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A DICTIONARY OF CHEMICAL SOLUBILITIES
INORGANIC



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TORONTO

A
DICTIONARY
OF
HEMICAL SOLUBILITIES
INORGANIC

FIRST EDITION

BY

ARTHUR MESSINGER COMEY, PH.D.

SECOND EDITION

ENLARGED AND REVISED

BY

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PREFACE TO FIRST EDITION

FOR many years a need has been felt by chemists for a book which shall collect into convenient form for ready reference the various data concerning the solubility of chemical substances that have been published from time to time in chemical periodicals and elsewhere.

The first mention that can be found of such a plan was made in 1731, when Peter Shaw delivered Chemical Lectures in London, as may be seen from the following:—

EXTRACTS from PETER SHAW's Chemical Lectures, publicly read at London in 1731 and 1732. London. Second Edition, London 1755. 8vo.

Page 97. Experiment I.—That Water as a Menstruum dissolves more of one body and less of another.

[He shows that two ounces of water dissolve two ounces of Epsom salt, five drachms of common salt, and eight grains of cream of tartar. Only in the latter case much remained undissolved until boiled.]

“It might be proper for the further Improvement of Chemistry and Natural Philosophy to form a Table of the Time and Quantity wherein all the known Salts are dissolvable in Water. . . . Such a Table regularly formed might ease the Trouble of refining Salts, by shewing at once without future Trial or Loss of Time how much Water each Salt required to dissolve it for Clarification, Filtration, or Crystallization. It would likewise supply us with a ready and commodious Way of separating any Mixture of Salts, by shewing which would first shoot out of the Mixture upon Crystallization. . . . The same Table might also direct us to a ready and commodious Method of separating two Salts without waiting for Crystallization. . . .”

It was many years, however, before the scheme suggested by Peter Shaw was put into execution. Professor F. H. Storer published the first work that undertook to carry out the idea in its entirety, in 1864, in a book, which he entitled “First Outlines of a Dictionary of Solubilities of Chemical Substances,” and which contained a compilation of nearly all the data on the subject published before 1860. It was at once recognized as a most valuable contribution to chemical literature; but for many years it has been difficult to obtain this work, as the limited edition which was published was soon wholly exhausted. Since then nothing has appeared on the subject except the brief tabulations found in various reference books, and no attempt has been made to cover the whole subject.

It is needless to state that the growth of chemical science since the publication of Professor Storer's book has been so enormous that that work has lost, at least to a great extent, the practical value it possessed thirty years ago. This growth has been indeed so great, and the data which have accumulated since 1860 so far surpass the earlier in volume, that a simple revision of Professor Storer's book was impracticable, and it therefore seemed best to start afresh.

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With the facilities offered by the various scientific libraries at Harvard University, the Massachusetts Institute of Technology, and other libraries in Boston, it has been possible to collect nearly all the data relating to the subject. For the work before 1860 Professor Storer's work has been found invaluable.

The method pursued has been to form a preliminary list of compounds with more or less data by consulting the two most complete works on inorganic chemistry—Gmelin-Kraut's "Handbuch der anorganischen Chemie" and Graham-Otto-Michaelis's "Lehrbuch." These statements have been verified and elaborated by consulting the original memoirs in all the periodicals devoted to chemical literature which were obtainable. The "Jahresbericht der Chemie" also has been used extensively in tracing references, but the original memoirs have always been consulted and references given to them when possible.

It has been found impracticable to draw any distinction as to reliability between the various data given by different observers. It was manifestly impossible to attempt to verify experimentally the statements of those who have carried on the researches, for the most assiduous labor of many could only cover a small portion of the attested facts. Therefore, even when two statements are directly contradictory, both have been given with the authority for each. The only exception to this has been made when more recent discoveries have shown beyond any reasonable doubt the falsity of previous work. In this way some of the older manifestly inaccurate work has been omitted. In a majority of cases the more recent work may be considered to be the more accurate, but this is not the invariable rule. A Synchronistic Table of the more common periodicals is given in the Appendix, whereby it is easy to determine the date of the publication of a research to which reference is made.

It may be objected by the practical chemist that most of the work previous to 1850 might well have been omitted, but a great deal of this work possesses at least a historical value, and often furnishes facts which have not since been verified. Much of the earlier work, when obviously of less importance, has been printed in smaller type.

The aim has been to include in this volume all analyzed inorganic substances, that is, all substances which do not contain carbon, but exception has been made in the case of CO_2 , CO , CS_2 , the carbonates, cyanides, ferro-cyanides, etc., which are here included.

The work has been brought up to March, 1894, when this volume went to press, and the results of researches published since that time are not included in the present edition.

It is hoped that this book will fill to some extent the want that has been felt by chemists for a compilation of this nature. While it has been attempted to make the book as free from errors as possible, nevertheless it is naturally impossible to avoid many mistakes, and the compiler will be very grateful to those who may call his attention to any errors or omissions.

A. M. C.

¹*CAMBRIDGE, MASS., Aug., 1895.*

PREFACE TO SECOND EDITION

DURING the twenty-five years which have elapsed since the publication of the first edition of this dictionary, a very large amount of work has been published in chemical periodicals, containing data concerning the solubility of inorganic chemical compounds. As it was impossible for the compiler of the first edition to devote the time necessary for the collecting of the published data, it was necessary to employ assistance, and Dr. Dorothy A. Hahn, Professor of Chemistry, Mt. Holyoke College, was engaged for this work. Dr. Hahn has collected the larger part of the material in this book, which work in spite of its arduous and tedious nature, she has performed in a most painstaking manner.

The compilation and arrangement of the data collected by Dr. Hahn, which devolved upon the original author, took much time. This, together with difficulties in printing, caused by the general conditions after the war, has delayed the publication until the present year, although the work was begun in 1916, and it has only been brought up to January 1st of that year.

Since the publication of the first edition of this work, Dr. Atherton Seidell has brought out two editions of his book, entitled "Solubilities of Inorganic and Organic Substances," which covers quite a different field, as he considers only quantitative data and those only for the commoner substances. Dr. Seidell has followed the plan in most cases where there are several available solubility determinations of a substance, of selecting and averaging the more reliable results, and embodying them in tables. Although this undoubtedly facilitates ready reference, it has seemed better to adhere to the original plan of the first edition, and to publish all the data in the form of the original authorities with references and dates, so that the user may be at liberty to use his own judgment in selection. Some few of the tables arranged by Dr. Seidell, however, have seemed to possess decided advantages over any other published data and they have been incorporated in the present volume. It is desired also to acknowledge indebtedness to Dr. Seidell's work for certain other tables where the original sources were not available to the present compilers.

The same plan and arrangement used in the first edition has been followed with certain elaboration, however, of the arrangement of data on the solubility of two or more salts in a solvent, which is explained in the Explanatory Preface.

Data published since the first edition on the cobalt and chromium ammonia compounds and those of the platinum group have been omitted, as it seemed that solubility data on those compounds possessed very little general interest.

As stated in the preface of the first edition, while every possible attempt has been made to avoid errors, it is manifestly impossible to avoid many mistakes in a work of this nature, and the compiler will be glad to have his attention called to any errors or omissions.

WILMINGTON, Del., Jan., 1921.

A. M. C.

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EXPLANATORY PREFACE

IN order to reduce this volume to a convenient size the subject-matter has been abbreviated and condensed as far as seemed compatible with clearness; but it has been the aim not to use any abbreviations which are not at once intelligible without consulting the explanatory table. The more common chemical formulæ have been universally used, thereby saving a large amount of space without detracting from ready intelligibility to chemists.

The solubility of the substance in water is first given, the data being arranged chronologically in the longer articles. Then follow the specific gravities of the aqueous solutions, and also any data obtainable regarding their boiling-points; other physical data concerning solutions are not included. Following this is the solubility of the substance in other solvents—first the inorganic acids, then alkali and salt solutions, and finally organic substances.

Owing to the great increase of data, published during the last twenty years, on the simultaneous solubility of two or more salts in a given solvent, it has been found necessary to plan some arrangement, whereby such data can easily be found, and the plan adopted is as follows: The data for the solubility of two salts in a solvent is placed under the salt which comes first according to the alphabetical arrangement in this dictionary, and the order of the data on various salts under the same heading follows the alphabetical order of the salts considered. Thus the data on the solubility of $\text{NH}_4\text{Cl} + \text{BaCl}_2$, $\text{NH}_4\text{Cl} + \text{CuCl}_2$, and $\text{NH}_4\text{Cl} + \text{PbCl}_2$, and $\text{NH}_4\text{Cl} + (\text{NH}_4)_2\text{SO}_4$ are placed under Ammonium Chloride and arranged in the given order. Certain exceptions have been made to this rule, where the data directly concerns the solubility of a salt in a solution of another salt, in which case, it is placed under the former. Numerous cross references, however, are given, which it is hoped will avoid confusion.

In many cases no definite distinction can be drawn between the phenomena of solution and decomposition. At present the theory of solution is in a confused state, and until what really takes place when a substance dissolves is thoroughly understood no distinct line can be drawn. The whole subject is unsettled at the present time; for while many chemists believe in the so-called "dissociation" theory, yet the "hydrate" theory is not without its supporters. It is not my intention to discuss the theoretical side of the question, which has been so well treated in many recent works. It is, however, obvious that the phenomena are essentially different, when, for example, sodium carbonate is dissolved in water, in which case the original salt is deposited on evaporation, and when iron is dissolved in sulphuric acid, and the solution deposits a sulphate of iron. Yet it is still the custom to speak of iron as soluble in sulphuric acid, although it would be much more accurate to say that the sulphuric acid was decomposed by the iron. It has thus been found impracticable to draw a sharp line between solu-

tion and decomposition, and the term "soluble" has in general been used where a solution of some sort is formed by the action of the solvent.

The matter of alphabetical arrangement of chemical compounds, in the present somewhat confused state of chemical nomenclature, has been a difficult question to decide. The plan followed has been practically that of the standard Dictionaries of Chemistry, whereby the compounds of metals with one of the non-metallic elements have been classified under the metals, while the salts of the other acids (the oxygen acids so called and some few others) have been arranged alphabetically under the acids. Thus barium chloride is found under barium, while barium chlorate is found under chloric acid. No exception has been made in the case of the rare metals, as is usually the custom in Dictionaries of Chemistry. Double salts are to be found under the word which comes first alphabetically; thus, "common alum," potassium aluminum sulphate, is found under aluminum sulphate as aluminum potassium sulphate (under sulphuric acid), but ammonia chrome alum is found under ammonium sulphate as ammonium chromium sulphate. In the same way the double sulphate and chromate of potassium is found under potassium chromate (chromic acid), and not under potassium sulphate (sulphuric acid). The double chloride of ammonium and magnesium is found under ammonium chloride (ammonium), while the double chloride of potassium and magnesium is found under magnesium chloride (magnesium). An exception is made, however, in the case of double compounds of salts of oxygen acids with salts containing a single non-metallic element, in which case they are always found under the oxygen acid. Thus the double sulphate and chloride of lead, PbSO_4 , PbCl_2 , is found under lead sulphate (sulphuric acid), and not under lead chloride (lead).

The above method in some cases widely separates analogous compounds, but it was found to be the only practical way to a strictly alphabetical arrangement, which is no necessary in a book containing so many very short articles.

The ammonia addition-products furnished another difficulty. While their nature is more or less definitely understood in the cobalt, platinum, etc., compounds, and a definite nomenclature is in general use, there is an absolute lack of anything of the kind in the less definite compounds. It is good usage to speak of cuprammonium compounds, but how shall we designate the analogous cadmium compounds? "Cadmammonium" has not yet received the sanction of chemists, and AlCl_3 , NH_3 is a still worse case for naming. I have, therefore, not attempted to name these compounds, but classified them all under the salts to which the ammonia is added, affixing the word ammonia, thus: aluminum chloride ammonia, cadmium chloride ammonia, and also cupric chloride ammonia for the salt now almost universally known as cuprammonium chloride.

The ammonia compounds of cobalt, chromium, mercury, and the platinum metals are arranged alphabetically according to their universally accepted names, a list of which is given under each of those elements.

It has further been necessary to settle arbitrarily the question whether a substance should be considered as a double salt or a salt of a compound acid con-

taining one of the metals. For example, "fluosilicates" (or silicofluorides, as some may prefer) is the general name for the double fluorides of SiF_4 and a metal, but this unanimity in usage gradually disappears as the basic elements become more nearly alike, so that it is impossible to draw a line between such compounds and a compound such as the double chloride of magnesium and potassium, for which indeed the name "potassium chloromagnesate" has been proposed. The aim has been in all these cases to follow the best usage rather than make an absolutely homogeneous system of nomenclature out of the existing confusion.

In the matter of formulæ no attempt at uniformity has been made. Thus in the case above some chemists write the formula of the double chloride of magnesium and potassium as KMgCl_3 , others as KCl, MgCl_3 . The form here used has been in most cases that of the author from whom the data are taken.

The prefixes mono, di, tri, ortho, pyro, etc., have in general been disregarded in the alphabetical arrangement, and have been printed in italics. Exceptions to this have been made, however, in the cobalt, chromium, etc., ammonium compounds, and in a few others, as dithionic, perchloric, etc., acids. Cross references have been used, so as to prevent any confusion arising from lack of uniformity in this respect.

In the Appendix will be found formulæ and tables for the conversion of the degrees of various hydrometer scales into specific gravity, and a Synchronistic Table of the Periodicals to which references are most frequently made.

Herz (Z. anorg. 25, 155) found that aluminum hydroxide which has been dried in a vacuum dessicator requires for solution in $\text{NaOH} + \text{Aq.}$ 3 atoms Na to 1 atom Al. Slade (Z. Elektrochem. 1911, 17, 261) was unable to obtain this result. Herz says Slade's error is due to insufficient shaking of the solution. (Herz, Z. Elektrochem. 1911, 17, 403.)

New solubility determinations verify the statement of Herz (Z. anorg. 25, 155) that the solubility of Al(OH)_3 in $\text{NaOH} + \text{Aq.}$ is proportional to the concentration of NaOH . They do not, however, verify his statement that the ratio Na : Al in the solutions is always 3 : 1, for the author finds that the ratio Na : Al varies from 2 : 1 to 10 : 1 depending on the conditions of precipitation and the method and duration of drying of the Al(OH)_3 . (Slade, Z. Elektrochem. 1912, 18, 1.)

Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ when freshly pptd., but presence of NH_4 salts diminish its solubility, and it separates out completely after long standing. (Fresenius.)

Somewhat sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ the more readily the larger the vol. of H_2O . Somewhat sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ but less than in $\text{NH}_4\text{OH} + \text{Aq.}$ Sl. sol. in dil. $\text{NH}_4\text{Cl} + \text{Aq.}$ unless that salt be in large excess. It is finally wholly pptd. if allowed to stand several days.

18752 pts. $\text{NH}_4\text{OH} + \text{Aq.}$ (4 % NH_4OH) dissolve an amt. of Al(OH)_3 corresponding to one pt. Al_2O_3 ; NH_4Cl prevents this solubility almost completely. (Hanamann, Pharm. Viertelj. 12, 527.)

Al(OH)_3 , prepared by ppt. of a solution of $\text{Al(NO}_3)_3$ with NH_4OH , filtered and washed, is insol. in $\text{NH}_4\text{OH} + \text{Aq.}$

Al(OH)_3 prepared by pptn. of a solution of potassium aluminate with NH_4Cl , is sol. in a large excess of NH_4OH if this is added to the ppt. at once. This modification which is sol. in NH_4OH is unstable and easily goes over into the modification which is insol. in NH_4OH . (Renz, B. 1903, 36, 2751.)

Conc. $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ does not dissolve Al(OH)_3 , and not a trace is dissolved by boiling conc. $\text{NH}_4\text{Cl} + \text{Aq.}$ (Weeren, Pogg. 92, 97.)

With $\text{NH}_4\text{F} + \text{Aq.}$ it forms a double salt, $\text{AlF}_3 \cdot 3\text{NH}_4\text{F}$, which is sol. in H_2O , but not in $\text{NH}_4\text{F} + \text{Aq.}$ (Helmholtz, Z. anorg. 3, 127.)

Insol. in $(\text{NH}_4)_2\text{S} + \text{Aq.}$ (Malaguti and Durocher, A. ch. (3) 17, 421.) Fuchs found, on the contrary, that it is not wholly insol. in $(\text{NH}_4)_2\text{S} + \text{Aq.}$ (Fresenius, Quant.)

Insol. in $\text{FeCl}_3 + \text{Aq.}$ (Béchamp.)

Determinations of the solubility of aluminum hydroxide in $\text{AlCl}_3 + \text{Aq.}$ show that part goes into solution to form a compound, while the greater part is in the colloidal form. (Fischer, Z. anorg. 1904, 40, 46.)

Only sl. sol. in conc. $\text{Al}_2(\text{SO}_4)_3 + \text{Aq.}$ but solubility increases with decrease in concentration of $\text{Al}_2(\text{SO}_4)_3$ until it reaches a maximum at a concentration of 32 % $\text{Al}_2(\text{SO}_4)_3$, at 20°, 28 % at 40°, and 38 % at 60°. With further decrease in concentration of $\text{Al}_2(\text{SO}_4)_3$

the solubility of Al(OH)_3 in $\text{Al}_2(\text{SO}_4)_3$ diminishes. (Kremann, C. A. 1909, 2422.)

Sol. in $\text{Ba(OH)}_2 + \text{Aq.}$ (Rose.)

Sol. in boiling $\text{Fe(NO}_3)_3$, $\text{Cr(NO}_3)_3$, $\text{Bi(NO}_3)_3$, $\text{Hg(NO}_3)_2$, HgNO_3 , SnCl_2 , and $\text{SbCl}_3 + \text{Aq.}$ (Persoz.)

Insol. in HCN or cold $\text{KCN} + \text{Aq.}$; but sl. sol. in hot $\text{KCN} + \text{Aq.}$ (Rose.)

Insol. in $\text{KC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ (Osann, 1821.)

When moist, sol. in $\text{H}_2\text{SO}_4 + \text{Aq.}$ from which it is repptd. on boiling. (Berthier, A. ch. (3) 7, 76.)

Somewhat sol. in $\text{NaC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ (Mercer.)

Not pptd. by $\text{NH}_4\text{OH} + \text{Aq.}$ in presence of Na citrate. (Spiller.)

Sol. in ethyl amine, amyl amine, sinkaline, ethyl picoline hydroxide, stibethylum hydroxide, triethyltoluenyl ammonium hydroxide + Aq. (Friedländer.)

Sol. in alkyl amines. (Renz, B. 1903, 36, 2751.)

Insol. in acetone. (Naumann, B. 1904, 37, 4328.)

Sol. to a considerable extent in $\text{K}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq.}$

Very sl. sol. in cane sugar + Aq. (Ramsey.)

Solubility in glycerine + Aq. containing about 60 % by vol. of glycerine. 100 cc. of the solution contain 0.25 g. Al_2O_3 . (Müller, Z. anorg. 1905, 43, 322.)

$\text{Al}_2\text{O}_3 \cdot 5\text{H}_2\text{O}$. Insol. in H_2O , $\text{NH}_4\text{OH} + \text{Aq.}$ and alcohol. Sol. in HCl and $\text{HNO}_3 + \text{Aq.}$ (Zunino, Gazz. ch. it. 1900, 30 (1), 194.)

$\text{Al}_6\text{O}_{14}\text{H}_{10}$, "Trialuminum hydroxide."

Not sol. in conc. acids in the cold; not sol. in KOH (cold) and only sl. sol. in hot KOH . Characterized by its solubility in exactly one mol. dil. HCl . Dil. solutions do not gelatinize even on long standing. Conc. solution of NH_4Cl and other salts cause ppt. which re-dissolves on addition of H_2O .

Alkalies and alkali carbonates decomp. the salt with HCl and ppt. trialuminium hydroxide. H_2SO_4 and sol. sulphates give insol. compds. with the hydrate. HNO_3 like HCl gives soluble compds. with the hydrate. (Structural formula given.) (Schlumberger, Bull. Soc. 1895, (5) 13, 41-65; C. C. 1895, I, 421.)

Aluminum iodide, AlI_3 .

Anhydrous. Fumes on air and deliquesces. Sol. in H_2O with evolution of much heat. Sol. in CS_2 and crystallizes from the hot sat. solution on cooling. (Weber.) Sol. in alcohol (Weber); ether and tetrachlormethane. (Gustavson.)

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913, 84, 26.)

+ $6\text{H}_2\text{O}$. Very sol. in H_2O .

Aluminum mercuric iodide, $\text{AlI}_3 \cdot \text{HgI}_2 + 8\text{H}_2\text{O}$.

Very deliquescent; sol. in H_2O without decomp. (Duboin, C. R. 1908, 146, 1028.)

ABBREVIATIONS OF REFERENCES

- A. —Annalen der Pharmacie, edited by Liebig and others, 1832–39; continued as Annalen der Chemie und Pharmacie, 1840–73; continued as Justus Liebig's Annalen der Chemie, 1874–1915+. 406 vols.
- A. ch.—Annales de Chimie et de Physique. Paris. 1st series, 1789–1816, 96 vols.; 2nd series, 1817–40, 78 vols.; 3rd series, 1841–63, 69 vols.; 4th series, 1864–73, 30 vols.; 5th series, 1874–83, 30 vols.; 6th series, 1884–93, 30 vols.; 7th series, 1893–1903, 30 vols.; 8th series, 1904–13, 30 vols.; 9th series, 1914+, 3 vols.
- Acta Lund.—Acta Universitatis Lundensis, or Lunds Universitets Års-skrift. Lund, 1864+.
- Am. Chemist.—The American Chemist. New York, 1870–77. 7 vols.
- Am. Ch. J.—The American Chemical Journal, edited by Remsen. Baltimore, 1879–1913. 50 vols.
- Am. J. Sci.—American Journal of Science and Arts, edited by Silliman, Dana, and others. New Haven. 1st series, 1818–45, 50 vols.; 2nd series, 1846–70, 50 vols.; 3rd series, 1871–95, 50 vols.; 4th series, 1896–1915+, 40 vols. Also numbered consecutively, 190 vols.
- Analyst.—The Analyst. London, 1876–1915+. 45 vols.
- Ann. chim. farm.—Annali di chimica e di farmacologia. Milan, 1886–90. 5 vols.
- Ann. des Mines.—See Ann. Min.
- Ann. Min.—Annales des Mines. Paris.
- Ann. Phil.—Annals of Philosophy. London. 1st series, 1813–20, 16 vols.; new series, 1821–26, 12 vols.
- Ann. Phys.—See Pogg. and W. Ann.
- Apoth. Z.—Apotheker-Zeitung. Berlin.
- Arb. Kais. Gesundheitsamt.—Arbeiten aus dem Kaiserlichen Gesundheitsamte.
- Arch. Néer. Sc.—Archives Néerlandaises des Sciences exactes et naturelles.
- Arch. Pharm.—Archiv der Pharmacie, continued from Archiv des Apothekervereins in Norddeutschland, which forms the 1st series. 1st series, 1822–34, 50 vols.; 2nd series, 1835–72, 150 vols.; 3rd series, 1873–94+, 32 vols. Also numbered consecutively, which system is exclusively used after 3rd series, vol. 253 (1915).
- Arch. sc. Phys. nat.—Archives des sciences physiques et naturelles de la Bibliothèque universelle de Genève.
- A. Suppl.—Annalen der Chemie und Pharmacie. Supplement-Bande. Vol. i. 1861; vol. ii, 1862–63; vol. iii. 1864–65; vol. iv. 1865–66; vol. v. 1867; vol. vi. 1868; vol. vii. 1870; vol. viii. 1872.
- B.—Berichte der deutschen chemischen Gesellschaft. Berlin, 1868–1915+. 48 vols.
- Att. Acc. Linc.—Atti della reale accademia dei Lincei, rendconditi, etc.
- B. A. B.—Sitzungsberichte der königlichen preussischen Akademie der Wissenschaften zu Berlin.
- Belg. Acad. Bull.—Bulletin de l'Académie Royale des Sciences, des Lettres, et des Beaux-Arts de Belgique.
- Berz. J. B.—Jahresbericht über die Fortschritte der physischen Wissenschaften, edited by Berzelius. 1822–47. 30 vols.
- Br. Arch.—Archiv des Apothekervereins im nördlichen Teutschland, etc., edited by Brandes. 1st series, 1822–31, 39 vols., corresponds to 1st series of Arch. Pharm.
- Bull. Acad. Crac.—Bulletin international de l'Académie des Sciences de Cracovie.
- Bull. Ac. St. Pétersb.—Bulletin de l'Académie Impériale des Sciences de St. Pétersbourg.
- Bull. Soc.—Bulletin des Séances de la Société chimique de Paris. 2nd series, 1864–88, 50 vols.; 3rd series, 1889–1906, 36 vols.; 4th series, 1907–15+, 18 vols.
- Bull. Soc. chim. Belg.—Bulletin de la Société chimique Belgique.
- Bull. Soc. ind. Mulhouse.—Bulletin de la Société industrielle de Mulhouse. 1828–49. 22 vols.
- Bull. Soc. Min.—Bulletin de la société française de Minéralogie. 1878–1915+. 37 vols.
- C. A.—Chemical Abstracts. American Chemical Society. New York.
- C. C.—Chemisches Centralblatt, continued from Pharmaceutisches Centralblatt.
- C. B. Miner.—Centralblatt für mineralogie, Geologie und Palæontologie. Berlin.
- Chem. Ind.—Die Chemische Industrie, edited by Jacobsen. Berlin.
- Chem. Soc.—Journal of the Chemical Society of London. 1st series, 1849–62, 15 vols.; 2nd series, 1863–78, 17 vols.; new series, 1878–1915+. The vols. are numbered consecutively from 1849. 1878 = vol. 32. Total, 108 vols.

amidophosphate and in $\text{NH}_4\text{OH} + \text{Aq.}$ Insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$

Acid. As the neutral salt.

Hydroxylamine amidophosphate, $(\text{NH}_2\text{O})\text{HPO}_2(\text{NH}_2).$

Sl. sol. in H_2O . (Stokes.)

Lithium amidophosphate, $\text{LiHPO}_2(\text{NH}_2).$

Sl. sol. in H_2O . (Stokes.)

Magnesium amidophosphate, $\text{MgPO}_2(\text{NH}_2) + 7\text{H}_2\text{O}$.

Very sl. sol. in H_2O ; quite easily sol. in dil. $\text{NH}_4\text{Cl} + \text{Aq.}$ Sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ (Stokes.)

$\text{MgH}_2(\text{PO}_2\text{NH}_2)_2 + 3\frac{1}{4}\text{H}_2\text{O}$. Insol. in $\text{NH}_4\text{Cl} + \text{Aq.}$ (Stokes.)

Manganese amidophosphate.

Neutral. Ppt.

Acid. Sl. sol. in H_2O .

Nickel amidophosphate.

Neutral. Ppt. Sol. in $\text{HC}_2\text{H}_3\text{O}_2$ or $\text{NH}_4\text{OH} + \text{Aq.}$

Acid. Sl. sol. in H_2O .

Potassium amidophosphate, $\text{K}_2\text{PO}_2(\text{NH}_2).$

Very sol. in H_2O and not decomp. by boiling. (Stokes.)

$\text{KHPO}_2(\text{NH}_2)$. Easily sol. in cold H_2O ; insol. in alcohol. (Stokes.)

Silver amidophosphate, $\text{Ag}_2\text{PO}_2(\text{NH}_2).$

Almost insol. in H_2O . Sol. in HNO_3 or $\text{NH}_4\text{OH} + \text{Aq.}$

$\text{AgHPO}_2(\text{NH}_2)$. Sl. sol. in H_2O ; easily sol. in dil. HNO_3 or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ also in $\text{NH}_4\text{OH} + \text{Aq.}$

Sodium amidophosphate, $\text{Na}_2\text{PO}_2(\text{NH}_2).$

Not deliquescent; very sol. in H_2O ; pptd. from aqueous solution by alcohol. (Stokes.)

$\text{NaHPO}_2(\text{NH}_2) + \frac{1}{4}(?)\text{H}_2\text{O}$. Nearly insol. in cold, and decomp. by hot H_2O . Insol. in alcohol.

Zinc amidophosphate.

Neutral. Perceptibly sol. in H_2O .

Acid. Sl. sol. in H_2O ; sol. in NH_4OH or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$

Diamidophosphoric acid, $\text{PO}(\text{NH}_2)_2\text{OH}$.

Sol. in cold H_2O ; almost insol. in alcohol; stable in the air but decomp. when heated and by boiling in aq. solution. (Stokes, Am. Ch. J. 1894, 16. 130.)

Barium diamidophosphate, $[\text{PO}(\text{NH}_2)_2\text{O}]_2\text{Ba}$.

Very sol. in H_2O ; insol. in alcohol; aq. solution decomp. slowly. (Stokes, Am. Ch. J. 1894, 16. 134.)

Magnesium diamidophosphate, $[\text{PO}(\text{NH}_2)_2\text{O}]_2\text{Mg}$.

Sol. in H_2O ; insol. in alcohol. (Stokes.)

Potassium diamidophosphate, $\text{PO}(\text{NH}_2)_2\text{OK}$.

Sol. in H_2O ; not deliquescent; insol. in alcohol. (Stokes.)

Silver diamidophosphate, $\text{PO}(\text{NH}_2)_2\text{OAg}$.

Very stable; insol. in H_2O . Very sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Stokes.)

Sodium diamidophosphate, $\text{PO}(\text{NH}_2)_2\text{ONa}$.

Sol. in H_2O ; not deliquescent; insol. in alcohol. (Stokes.)

Diamidotrihydroxyphosphoric acid.

Silver diamidotrihydroxyphosphate, $(\text{AgO})_3\text{P}(\text{NHAg})_3$.

(Stokes, Am. Ch. J. 1894, 16. 147.)

$(\text{AgO})_3\text{P}(\text{NH}_2)(\text{NHAg})$. Insol. in cold H_2O . (Stokes.)

$(\text{AgO})_3\text{P}(\text{NH}_2)_3$. Decomp. by cold H_2O . (Stokes.)

$+ 2\text{H}_2\text{O}$. Decomp. by boiling H_2O . (Stokes.)

Amidoimidophosphoric acid.

Amidoheximidoheptaphosphoric acid, $\text{OH} \cdot \text{PO} \cdot (\text{NH}_2)[\text{NH} \cdot \text{PO}(\text{OH})]_5 \cdot \text{NH} \cdot \text{PO}(\text{OH})_2$
 $= \text{P}_7\text{N}_7\text{O}_{18}\text{H}_{16}$.

Known only in solution in H_2O . (Stokes, Am. Ch. J. 1898, 20. 758.)

Silver diamidopyrimidophosphate, $\text{NH}(\text{PO} \cdot \text{NH}_2 \cdot \text{OAg})_2$.

Almost insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Stokes, Am. Ch. J. 1894, 16. 136.)

Silver amidoletrimido pentaphosphate, $\text{P}_5\text{N}_3\text{O}_{11}\text{H}_3\text{Ag}_7$.

Ppt. (Stokes, Am. Ch. J. 1898, 20. 752.)

Silver amidoheximidoheptaphosphate, $\text{P}_7\text{N}_7\text{O}_{18}\text{H}_3\text{Ag}_7$.

Ppt.; decomp. by acetic acid. (Stokes, Am. Ch. J. 1898, 20. 759.)

Sodium amidodiiimidotriphosphate, $\text{PO} \cdot \text{ONa} < \text{NH} \cdot \text{PO}(\text{ONa})_2$.

$= \text{P}_3\text{N}_3\text{O}_7\text{H}_4\text{Na}_4 + \text{H}_2\text{O}$.

Unstable; sol. in H_2O ; insol. in alcohol. (Stokes, Am. Ch. J. 1896, 18. 643.)

Sodium amidoheximidoheptaphosphate, $\text{P}_7\text{N}_7\text{O}_{18}\text{H}_3\text{Na}_7$.

Sol. in H_2O ; pptd. by alcohol. (Stokes, Am. Ch. J. 1898, 20. 758.)

Amidophosphimic acid.**Silver amidophosphimate, $P(NH)NH_2(OAg)_2$.**

Decomp. by heat; decomp. in contact with H_2O . (Stokes, Am. Ch. J. 1894, 16. 139.)

$(AgO)_2P(NAg)(NHA g)$.(?) Sl. sol. in $NH_4OH + Aq$. (Stokes, Am. Ch. J. 1894, 16. 149.)

Amidosulphonic acid, $HOSO_2NH_2$.

Easily sol. in H_2O , less easily in alcohol. (Berglund, B. 9. 252 and 1896.)

Very stable; less easily sol. in H_2O than its K salt. (Raschig, A. 241. 177.)

Stable in air. Non-deliquescent when cold. Sol. in 5 pts. H_2O at 0° and in $2\frac{1}{2}$ pts. H_2O at 70° . Solution in H_2O can be boiled several minutes without decomp. Solubility is decreased by addition of H_2SO_4 , so that if $1\frac{1}{5}$ – $1\frac{1}{4}$ pt. H_2SO_4 is added to H_2O , 100 pts. of the liquid dissolve only 3 pts. $HOSO_2NH_2$ in the cold. Pptd. from solution by HNO_3 or glacial acetic acid, but not by HCl . Solubility is decreased by presence of $NaHSO_4$. (Divers and Haga, Chem. Soc. 1896, 69. 1641.)

Amidosulphonates.

Easily sol. in H_2O ; sl. sol. in alcohol.

Aluminum amidosulphonate.

Very sol. in H_2O . (Berglund, Bull. Soc. (2) 29. 422.)

Ammonium amidosulphonate, $(NH_4)NH_2SO_3$.

Deliquescent. Sol. in H_2O ; insol. in alcohol.

Ammonium silver amidosulphonate, $NH_4SO_3(NH_2), AgSO_3(NH_2)$.

(Ephraim & Gurewitsch, B. 1910, 43. 148.)

Barium amidosulphonate, $Ba(NH_2SO_3)_2$.

Sol. in 3 pts. H_2O . (Berglund, l.c.)

Cadmium amidosulphonate, $Cd(NH_2SO_3)_2 + 5H_2O$.

Very sol. in H_2O . (B.)

Calcium amidosulphonate, $Ca(NH_2SO_3)_2 + 4H_2O$.

Very sol. in H_2O . (B.)

Cobalt amidosulphonate, $Co(NH_2SO_3)_2 + 3H_2O$.

Sol. in H_2O . (B.)

Copper amidosulphonate, $Cu(NH_2SO_3)_2 + 2H_2O$.

Sol. in H_2O . (B.)

Gold (auric) potassium amidosulphonate, $K_2Au_2(NSO_3)_2$.

Very sl. sol. in cold, more easily sol. in hot

H_2O . Sol. in dil. $HCl + Aq$. (Hofmann, B. 1912, 45. 1735.)

Lead amidosulphonate, $Pb(NH_2SO_3)_2 + H_2O$.

The most sol. of all amidosulphonates. (B.)

Lithium amidosulphonate, $LiNH_2SO_3$.

Deliquescent. (B.)

Magnesium amidosulphonate.

Very sol. in H_2O .

Manganese amidosulphonate, $Mn(NH_2SO_3)_2 + 3H_2O$.

Very sol. in H_2O . (B.)

Mercuric amidosulphonate, basic, $Hg(HgOSO_2NH_2)_2$.

Insol. in 3.5 % $HNO_3 + Aq$. Very sol. in 3 % $HCl + Aq$. (Hofmann, B. 1912, 45. 1733. $+ 2H_2O$. Insol. in hot H_2O . Sol. in $KOH + Aq$. (Divers and Haga, Chem. Soc. 1896, 69. 1649.)

Mercuric potassium amidosulphonate, $KHgNSO_3$.

Very sl. sol. in cold H_2O and cold dil. $KOH + Aq$. Sol. in 3 % $HCl + Aq$. (Hofmann, B. 1912, 45. 1732.)

Mercuric sodium amidosulphonate, $NaHgNSO_3$.

Nearly completely sol. in hot H_2O . (Hoffmann, B. 1912, 45. 1734.)

Nickel amidosulphonate, $Ni(NH_2SO_3)_2 + 3H_2O$.

Sol. in H_2O . (B.)

Potassium amidosulphonate, KNH_2SO_3 .

Sol. in H_2O . (Berglund.)

Potassium silver amidosulphonate, $NHAgSO_3K + H_2O$.

Decomp. by H_2O ; sol. in $NH_4OH + Aq$. (Hoffmann, B. 1912, 45. 1734.)

Silver amidosulphonate, $AgNH_2SO_3$.

Sol. in 15 pts. H_2O at 19° (B.)

Sodium amidosulphonate, $NaNH_2SO_3$.

Sol. in H_2O .

Strontium amidosulphonate, $Sr(NH_2SO_3)_2 + 4H_2O$.

Sol. in H_2O .

Thallium amidosulphonate, $TlNH_2SO_3$.

Sol. in H_2O .

Uranyl amidosulphonate.

Sol. in H_2O .

Zinc amidosulphonate, $Zn(NH_2SO_3)_2 + 4H_2O$.

Sol. in H_2O .

Amidosulphurous acid.**Ammonium amidosulphite, $\text{NH}_4\text{SO}_2\text{NH}_4$.**

Very deliquescent. Decomp. in the air with loss of NH_3 . Sol. in H_2O with decomp. Sol. in anhydrous alcohol. Sl. sol. in dry ether. (Divers, Chem. Soc. 1900, 77. 330.)

Ammonia, NH_3 .

Very sol. in H_2O , with evolution of much heat.

1 vol. H_2O absorbs 670 vols. ($\frac{1}{4}$ pt. by weight) NH_3 at $+10^\circ$ and 29.8 in. pressure; sp. gr. of solution = 0.875. (Davy.)

At low temperatures H_2O absorbs more than $\frac{1}{4}$ its weight of NH_3 , and sp. gr. of solution = 0.850. (Dalton.)

100 pts. H_2O absorb 8.41 pts. NH_3 at 24° , 5.96 pts. at 55° . (Osann.)

1 vol. H_2O absorbs 780 vols. NH_3 , 6 vols. H_2O increasing to 40 vols. sat. $\text{NH}_4\text{OH} + \text{Aq}$; 1 vol. sat. $\text{NH}_4\text{OH} + \text{Aq}$ contains 468 vols. NH_3 . (Thomson.)

1 vol. H_2O absorbs 450 vols. NH_3 at 15° . (Dumas.)

1 vol. H_2O absorbs 700 vols. NH_3 at ordinary temperature. (Otto.)

100 pts. H_2O absorb in NH_3 gas 47.7 pts. NH_3 by weight. (Berzelius.)

1 vol. H_2O absorbs 305 vols. NH_3 and vol. is increased to 1.5 vol., and sp. gr. becomes 0.900. (Ure.)

1 vol. H_2O at 0° and 760 mm. absorbs 1177.3 vols. NH_3 . (Sims.)

1 vol. H_2O at 0° and 760 mm. absorbs 1146 vols. NH_3 . (Roscoe and Dittmar.)

1 vol. H_2O at 0° and 760 mm. absorbs 1049.6 vols. NH_3 . (Carius.)

1 vol. H_2O at 0° and 760 mm. absorbs 1270 vols. NH_3 . (Berthelot.)

1 vol. H_2O at 0° and 760 mm. absorbs 1050 vols. NH_3 . (Bunsen.)

100 cc. H_2O absorb 64.50 g. NH_3 . (Raoult.)

Solubility of NH_3 in H_2O at 760 mm. and t° :
1 g. H_2O absorbs g. NH_3 , according to
Roscoe and Dittmar (A. 122. 347) (RD);
and according to Sims (A. 118. 345) (S).

t°	g. NH_3 RD	g. NH_3 S	t°	g. NH_3 RD	g. NH_3 S
0	0.875	0.899	36	0.343	0.363
2	0.833	0.853	38	0.324	0.350
4	0.792	0.809	40	0.307	0.338
6	0.751	0.765	42	0.290	0.326
8	0.713	0.724	44	0.275	0.315
10	0.679	0.684	46	0.259	0.304
12	0.645	0.646	48	0.244	0.294
14	0.612	0.611	50	0.229	0.284
16	0.582	0.578	52	0.214	0.274
18	0.554	0.546	54	0.200	0.265
20	0.526	0.518	56	0.186	0.256
22	0.499	0.490	58	0.247
24	0.474	0.467	60	0.238
26	0.449	0.446	70	0.194
28	0.426	0.426	80	0.154
30	0.403	0.408	90	0.114
32	0.382	0.393	98	0.082
34	0.362	0.378	100	0.074

Solubility of NH_3 by vol. in H_2O at 760 mm. and t° : 1 vol. H_2O at 760 mm. and t° dissolves V vols. NH_3 gas, vols. reduced to 0° and 760 mm.

t°	V	t°	V'
0	1049.60	13	759.55
1	1020.78	14	743.11
2	993.26	15	727.22
3	966.98	16	711.82
4	941.88	17	696.85
5	917.90	18	682.26
6	894.99	19	667.99
7	873.09	20	653.99
8	852.14	21	640.19
9	831.98	22	626.54
10	812.76	23	612.98
11	794.32	24	599.46
12	776.60	25	585.94

(Carius, A. 99. 144.)

Solubility of NH_3 in H_2O at P mm. pressure and 0° : 1 pt. H_2O absorbs pts. NH_3 at P mm. pressure and 0° .

P	Pts. NH_3	P	Pts. NH_3
10	0.044	900	0.968
20	0.084	950	1.101
30	0.120	1000	1.037
40	0.149	1050	1.075
50	0.175	1100	1.117
75	0.228	1150	1.161
100	0.275	1200	1.208
125	0.315	1250	1.258
150	0.351	1300	1.310
175	0.382	1350	1.361
200	0.411	1400	1.415
250	0.465	1450	1.469
300	0.515	1500	1.526
350	0.561	1550	1.584
400	0.607	1600	1.645
450	0.646	1650	1.707
500	0.690	1700	1.770
550	0.731	1750	1.835
600	0.768	1800	1.906
650	0.804	1850	1.976
700	0.840	1900	2.046
750	0.872	1950	2.120
800	0.906	2000	2.195
850	0.937		

(Roscoe and Dittmar, A. 112. 349.)

In proportion as the temperature is higher, so much the more nearly does the solubility of NH_3 in H_2O conform to the law of Henry and Dalton, but only obeys it completely when the temperature is 100° , as is seen in the following table.

Solubility of NH_3 in H_2O at various pressures and temperatures: P=partial pressure, i. e. total pressure minus the tension of aqueous vapour at the given temperature; G=grams NH_3 dissolved in 1 g. H_2O at the given pressure; G at 760=grams NH_3 that would be contained in 1 g. H_2O if the solubility was proportional to the pressure.

P	0°		20°		40°		100°	
	G at P	G at 760	G at P	G at 760	G at P	G at 760	G at P	G at 760
20	0.082	3.113
30	0.117	2.960
40	0.148	2.820
60	0.169	2.522	0.119	1.513
80	0.240	2.280	0.141	1.337	0.052	0.497
100	0.280	2.127	0.158	1.200	0.064	0.490
120	0.316	2.000	0.173	1.095	0.076	0.483
140	0.346	1.880	0.187	1.017	0.088	0.476
160	0.375	1.780	0.202	0.962	0.099	0.470
180	0.398	1.684	0.207	0.918	0.109	0.462
200	0.421	1.598	0.232	0.881	0.120	0.454
250	0.472	1.434	0.266	0.810	0.145	0.440
300	0.519	1.315	0.296	0.750	0.168	0.426
350	0.563	1.223	0.325	0.705	0.191	0.414
400	0.606	1.152	0.353	0.670	0.211	0.402
450	0.650	1.100	0.378	0.638	0.232	0.399
500	0.692	1.052	0.403	0.612	0.251	0.382
550	0.732	1.012	0.425	0.587	0.269	0.372
600	0.770	0.975	0.447	0.566	0.287	0.363
650	0.809	0.946	0.470	0.550	0.304	0.355
700	0.850	0.923	0.492	0.534	0.320	0.347	0.068	0.074
750	0.891	0.903	0.514	0.521	0.335	0.339	0.073	0.074
760	0.899	0.899	0.518	0.518	0.338	0.338	0.074	0.074
800	0.937	0.888	0.535	0.504	0.349	0.332	0.078	0.074
850	0.980	0.876	0.556	0.497	0.363	0.325	0.083	0.074
900	1.029	0.869	0.574	0.485	0.378	0.319	0.088	0.074
950	1.077	0.862	0.594	0.475	0.391	0.313	0.092	0.073
1000	1.126	0.855	0.613	0.466	0.404	0.307	0.096	0.073
1050	1.177	0.852	0.632	0.457	0.414	0.300	0.101	0.073
1100	1.230	0.850	0.651	0.450	0.425	0.294	0.106	0.073
1150	1.283	0.848	0.669	0.442	0.434	0.287	0.110	0.073
1200	1.336	0.846	0.685	0.433	0.445	0.282	0.115	0.073
1250	1.338	0.844	0.704	0.428	0.454	0.276	0.120	0.073
1300	1.442	0.843	0.722	0.422	0.463	0.271	0.125	0.073
1350	1.496	0.842	0.741	0.417	0.472	0.266	0.130	0.073
1400	1.549	0.841	0.761	0.413	0.479	0.260	0.135	0.073
1450	1.603	0.840	0.780	0.409	0.486	0.255
1500	1.656	0.839	0.801	0.406	0.493	0.250
1600	1.758	0.835	0.842	0.400	0.511	0.242
1700	1.861	0.832	0.881	0.394	0.530	0.237
1800	1.966	0.830	0.919	0.388	0.547	0.231
1900	2.070	0.828	0.955	0.382	0.565	0.226
2000	0.992	0.377	0.579	0.220
2100	0.594	0.215

(Sims, A. 118. 346.)

Solubility of NH_3 in H_2O at temps. below 0° . One gram H_2O dissolves

grams NH_3	Temp.
0.947	-3.9°
1.115	-10°
1.768	-20°
2.781	-30°
2.946	-40°

(Mallet, *Am. Ch. J.* 1897, 19, 807.)

The solubility of NH_3 in H_2O does not follow Dalton's law at ord. temp., but does at temp. near 100° . (Konowaloff, *J. Russ. Phys. Chem. Soc.* 1894, 26, 48; *Chem. Soc.* 1896, 70 (2) 351.)

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq.}$

% NH_3	Sp. gr.	% NH_3	Sp. gr.
32.3*	0.8750	14.53	0.9435
29.25	0.8857	13.46	0.9476
26	0.9000	12.40	0.9513
25.37*	0.9054	11.56	0.9545
22.07	0.9166	10.82	0.9573
19.54	0.9255	10.17	0.9597
17.52	0.9326	9.6	0.9616
15.88	0.9385	9.5*	0.9632

(H. Davy, *Elements*, 1, 241.)

* By direct experiment. The other numbers were obtained by calculation, making no allowance for compensation.

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 16° , according to Otto in his *Lehrbuch*.

% NH_3	Sp. gr.	% NH_3	Sp. gr.
12.000	0.9517	8.500	0.9650
11.875	0.9521	8.375	0.9654
11.750	0.9526	8.250	0.9659
11.625	0.9531	8.125	0.9664
11.500	0.9536	8.000	0.9669
11.375	0.9540	7.875	0.9673
11.250	0.9545	7.750	0.9678
11.125	0.9550	7.625	0.9683
11.000	0.9555	7.500	0.9688
10.950	0.9556	7.375	0.9692
10.875	0.9559	7.250	0.9697
10.750	0.9564	7.125	0.9702
10.625	0.9569	7.000	0.9707
10.500	0.9574	6.875	0.9711
10.375	0.9578	6.750	0.9716
10.250	0.9583	6.625	0.9721
10.125	0.9588	6.500	0.9726
10.000	0.9593	6.375	0.9730
9.875	0.9597	6.250	0.9735
9.750	0.9602	6.125	0.9740
9.625	0.9607	6.000	0.9745
9.500	0.9612	5.875	0.9749
9.375	0.9616	5.750	0.9754
9.250	0.9621	5.625	0.9759
9.125	0.9626	5.500	0.9764
9.000	0.9631	5.375	0.9768
8.875	0.9636	5.250	0.9773
8.750	0.9641	5.125	0.9778
8.625	0.9645	5.000	0.9783

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$, according to Ure in *Dict. of Arts*.

% NH_3	Sp. gr.	% NH_3	Sp. gr.
27.940	0.8914	15.900	0.9363
27.633	0.8937	14.575	0.9410
27.088	0.8967	13.250	0.9455
26.751	0.8983	11.925	0.9510
26.500	0.9000	10.600	0.9564
25.175	0.9045	9.275	0.9614
23.850	0.9090	7.950	0.9662
22.525	0.9133	6.625	0.9716
21.200	0.9177	5.300	0.9768
19.875	0.9227	3.975	0.9819
18.550	0.9275	2.650	0.9887
17.225	0.9320	1.325	0.9955

Sp. gr., b.-pt., and vols. gas in $\text{NH}_4\text{OH} + \text{Aq}$.

% NH_3	Sp. gr.	B.-pt.	Vols. gas in 1 vol. liquid
35.3	0.85	-3.3°	494
32.6	0.86	$+3.3^\circ$	456
29.9	0.87	10°	419
27.3	0.88	16.6°	382
24.7	0.89	23.3°	346
22.2	0.90	30°	311
19.8	0.91	36.6°	277
17.4	0.92	43.3°	244
15.1	0.93	50°	211
12.8	0.94	56.6°	180
10.5	0.95	63.3°	147
8.3	0.96	70°	116
6.2	0.97	78.3°	87
4.1	0.98	86.1°	57
2.0	0.99	91.1°	28

(Dalton, in *New System*, 2, 422.)

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ sat. at t° .

t°	Sp. gr.	t°	Sp. gr.	t°	Sp. gr.
0	0.8535	9	0.8746	18	0.8903
1	0.8561	10	0.8766	19	0.8916
2	0.8587	11	0.8785	20	0.8928
3	0.8611	12	0.8804	21	0.8940
4	0.8635	13	0.8823	22	0.8952
5	0.8658	14	0.8841	23	0.8963
6	0.8681	15	0.8858	24	0.8974
7	0.8703	16	0.8874	25	0.8984
8	0.8725	17	0.8889	

(Carius, *A.* 99, 141.)

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 14° , according to Carius (*A.* 99, 148).

% NH_3	Sp. gr.	% NH_3	Sp. gr.
36.0	0.8844	35.2	0.8860
35.8	0.8848	35.0	0.8864
35.6	0.8852	34.8	0.8868
35.4	0.8856	34.6	0.8872

of $\text{NH}_4\text{OH} + \text{Aq}$ at 14° , etc.—*Cont.*

Sp. gr.	% NH_3	Sp. gr.
0 8877	22.2	0 9185
0 8881	22.0	0 9191
0 8885	21.8	0 9197
0 8889	21.6	0 9203
0 8894	21.4	0 9209
0 8898	21.2	0 9215
0 8903	21.0	0 9221
0 8907	20.8	0 9227
0 8911	20.6	0 9233
0 8916	20.4	0 9239
0 8920	20.2	0 9245
0 8925	20.0	0 9251
0 8929	19.8	0 9257
0 8934	19.6	0 9264
0 8938	19.4	0 9271
0 8944	19.2	0 9277
0 8948	19.0	0 9283
0 8953	18.8	0 9289
0 8957	18.6	0 9296
0 8962	18.4	0 9302
0 8967	18.2	0 9308
0 8971	18.0	0 9314
0 8976	17.8	0 9321
0 8981	17.6	0 9327
0 8986	17.4	0 9333
0 8991	17.2	0 9340
0 8996	17.0	0 9347
0 9001	16.8	0 9353
0 9006	16.6	0 9360
0 9011	16.4	0 9366
0 9016	16.2	0 9373
0 9021	16.0	0 9380
0 9026	15.8	0 9386
0 9031	15.6	0 9393
0 9036	15.4	0 9400
0 9041	15.2	0 9407
0 9047	15.0	0 9414
0 9051	14.8	0 9420
0 9057	14.6	0 9427
0 9063	14.4	0 9434
0 9068	14.2	0 9441
0 9073	14.0	0 9449
0 9078	13.8	0 9456
0 9083	13.6	0 9463
0 9089	13.4	0 9470
0 9094	13.2	0 9477
0 9100	13.0	0 9484
0 9106	12.8	0 9491
0 9111	12.6	0 9498
0 9116	12.4	0 9505
0 9122	12.2	0 9512
0 9127	12.0	0 9520
0 9133	11.8	0 9527
0 9139	11.6	0 9534
0 9145	11.4	0 9542
0 9150	11.2	0 9549
0 9156	11.0	0 9556
0 9162	10.8	0 9563
0 9168	10.6	0 9571
0 9174	10.4	0 9578
0 9180	10.2	0 9586

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 14° , etc.—*Cont.*

% NH_3	Sp. gr.	% NH_3	Sp. gr.
10.0	0.9593	5.0	0.9790
9.8	0.9601	4.8	0.9799
9.6	0.9608	4.6	0.9807
9.4	0.9616	4.4	0.9815
9.2	0.9623	4.2	0.9823
9.0	0.9631	4.0	0.9831
8.8	0.9639	3.8	0.9839
8.6	0.9647	3.6	0.9847
8.4	0.9654	3.4	0.9855
8.2	0.9662	3.2	0.9863
8.0	0.9670	3.0	0.9873
7.8	0.9677	2.8	0.9882
7.6	0.9685	2.6	0.9890
7.4	0.9693	2.4	0.9899
7.2	0.9701	2.2	0.9907
7.0	0.9709	2.0	0.9915
6.8	0.9717	1.8	0.9924
6.6	0.9725	1.6	0.9932
6.4	0.9733	1.4	0.9941
6.2	0.9741	1.2	0.9950
6.0	0.9749	1.0	0.9959
5.8	0.9757	0.8	0.9967
5.6	0.9765	0.6	0.9975
5.4	0.9773	0.4	0.9983
5.2	0.9781	0.2	0.9991

Hager also gives a table in his *Commentar zur Pharmacopoea*, which is practically identical with those here given.

Strength of $\text{NH}_4\text{OH} + \text{Aq}$ of certain sp. gr. at 12°

Sp. gr.	1 kg. solution contains g. NH_3	1 l. solution contains g. NH_3	1 litre consists of	
			H_2O in cc.	liquid NH_3 in cc.
0 870	384.4	334.5	535.5	464.5
0 880	347.2	305.5	574.5	425.5
0 890	311.6	277.3	612.7	387.3
0 900	277.3	249.5	650.5	349.5
0 910	244.9	222.8	687.2	312.8
0 920	213.4	196.3	723.7	276.3
0 930	182.9	170.1	759.9	240.1
0 940	152.9	143.7	796.3	203.7
0 950	124.2	118.0	832.0	168.0
0 960	97.0	93.1	866.9	133.1
0 970	70.2	68.0	902.0	98.0
0 980	45.3	44.3	935.7	64.3
0 990	21.0	20.7	969.3	30.7

(Wachsmuth, *Arch. Pharm.* (3) 8. 510.)

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 15°
(Most careful experiments.)

Sp. gr.	% NH_3	Sp. gr.	% NH_3
0 990	2.15	0 926	19.50
0 974	6.10	0 916	22.50
0 950	12.54	0 910	24.40

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 15° —Continued

Sp. gr.	% NH_3	Sp. gr.	% NH_3
0.900	27.70	0.882	34.8
0.890	31.40	0.880	35.5
0.885	33.5

(Grüneberg, Chem. Ind. 12. 97.)

The following table is calculated from the above by interpolation:—

Sp. gr.	% NH_3	Sp. gr.	% NH_3
0.995	1.05	0.935	16.90
0.990	2.15	0.930	18.35
0.985	3.30	0.925	19.80
0.980	4.50	0.920	21.30
0.975	5.75	0.915	22.85
0.970	7.05	0.910	24.40
0.965	8.40	0.905	26.00
0.960	9.80	0.900	27.70
0.955	11.20	0.895	29.50
0.950	12.60	0.890	31.40
0.945	14.00	0.885	33.40
0.940	15.45	0.880	35.50

(Grüneberg.)

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 14° .

% HN_3	Sp. gr.	% NH_3	Sp. gr.
31	0.8933	15.6	0.9400
23.8	0.9116	11.7	0.9536
20.4	0.9246	5.1	0.9780

(Lunge and Smith, B. 17. 777.)

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 15° , according to Lunge and Wiernik (Zeit. f. angew. Ch. 1889. 183).

(Most carefully worked out and calculated.)

Sp. gr.	% NH_3	1 l. contains g. NH_3	Correction for $\pm 1^\circ$
1.000	0.00	0.0	0.00018
0.998	0.45	4.5	0.00018
0.996	0.91	9.1	0.00019
0.994	1.37	13.6	0.00019
0.992	1.84	18.2	0.00020
0.990	2.31	22.9	0.00020
0.988	2.80	27.7	0.00021
0.986	3.30	32.5	0.00021
0.984	3.80	37.4	0.00022
0.982	4.30	42.2	0.00022
0.980	4.80	47.0	0.00023
0.978	5.30	51.8	0.00023
0.976	5.80	56.6	0.00024
0.974	6.30	61.4	0.00024
0.972	6.80	66.1	0.00025
0.970	7.31	70.9	0.00025
0.968	7.82	75.7	0.00026

Sp. gr. of $\text{NH}_4\text{OH} + \text{Aq}$ at 15° , etc.—Continued

Sp. gr.	% NH_3	1 l. contains g. NH_3	Correction for $\pm 1^\circ$
0.966	8.33	80.5	0.00026
0.964	8.84	85.2	0.00027
0.962	9.35	89.9	0.00028
0.960	9.91	95.1	0.00029
0.958	10.47	100.3	0.00030
0.956	11.03	105.4	0.00031
0.954	11.60	110.7	0.00032
0.952	12.17	115.9	0.00033
0.950	12.74	121.0	0.00034
0.948	13.31	126.2	0.00035
0.946	13.88	131.3	0.00036
0.944	14.46	136.5	0.00037
0.942	15.04	141.7	0.00038
0.940	15.63	146.9	0.00039
0.938	16.22	152.1	0.00040
0.936	16.82	157.4	0.00041
0.934	17.42	162.7	0.00041
0.932	18.03	168.1	0.00042
0.930	18.64	173.4	0.00042
0.928	19.25	178.6	0.00043
0.926	19.87	184.2	0.00044
0.924	20.49	189.3	0.00045
0.922	21.12	194.7	0.00046
0.920	21.75	200.1	0.00047
0.918	22.39	205.6	0.00048
0.916	23.03	210.9	0.00049
0.914	23.68	216.3	0.00050
0.912	24.33	221.9	0.00051
0.910	24.99	227.4	0.00052
0.908	25.65	232.9	0.00053
0.906	26.31	238.3	0.00054
0.904	26.98	243.9	0.00055
0.902	27.65	249.4	0.00056
0.900	28.33	255.0	0.00057
0.898	29.01	260.5	0.00058
0.896	29.69	266.0	0.00059
0.894	30.37	271.5	0.00060
0.892	31.05	277.0	0.00061
0.890	31.75	282.6	0.00061
0.888	32.50	288.6	0.00062
0.886	33.25	294.6	0.00063
0.884	34.10	301.4	0.00064
0.882	34.95	308.3	0.00065

NH_3 is much less sol. in KOH , or $\text{NaOH} + \text{Aq}$ than in H_2O .

Solubility of NH_3 in H_2O , and $\text{KOH} + \text{Aq}$ of various strengths: 100 pts. solvent absorbs g. NH_3 at t° .

t°	H_2O	$\text{KOH} + \text{Aq}$ 11.25% K_2O	$\text{KOH} + \text{Aq}$ 25.25% K_2O
0	90.00	72.00	49.50
8	72.75	57.00	37.50
16	59.75	46.00	28.50
24	49.50	37.25	21.75

(Reault, A. ch. (5) 1. 262.)

100 pts. sat. KOH + Aq dissolve only 1 pt. NH₃.

Solubility in NaOH + Aq is the same as in KOH + Aq of the same strength.

NH₄Cl + Aq absorbs slightly less NH₃ than the same vol. H₂O. NaNO₃, and NH₄NO₃ + Aq absorb almost the same amount NH₃ as the same vol. H₂O. (Raoult, l.c.)

Solubility of NH₃ in 100 pts. Ca(NO₃)₂ + Aq.

t°	H ₂ O	Ca(NO ₃) ₂ + Aq 23.38% Ca(NO ₃) ₂	Ca(NO ₃) ₂ + Aq 59.03% Ca(NO ₃) ₂
0	90.00	96.25	104.50
8	72.75	78.50	84.75
16	59.75	65.00	70.50

(Raoult, l.c.)

Solubility in salt solutions at 25°C.

Salt	Mols. NH ₃ soluble in 1 liter of		
	.5-normal solution	1-normal solution	1.5-nor- mal solution
KCl	0.930	0.866	0.809
KBr	0.950	0.904	0.857
KI	0.970	0.942	0.900
KOH	0.852	0.716	0.607
NaCl	0.938	0.889	0.843
NaBr	0.965	0.916	0.890
NaI	0.995	0.992	0.985
NaOH	0.876	0.789	0.716
LiCl	0.980	1.008	1.045
LiBr	1.001	1.040	1.090
LiI	1.030	1.094	1.190
LiOH	0.865	0.808	0.768
KF	0.839	0.722	0.626
KNO ₃	0.923	0.862	0.804
KNO ₂	0.920	0.855	0.798
KCN	0.926	0.858	0.802
KCNS	0.932	0.868	0.814
1/2K ₂ SO ₄	0.875	0.772	0.678
1/2K ₂ SO ₃	0.865	0.768	0.675
1/2K ₂ CO ₃	0.788	0.650	0.554
1/2K ₂ C ₂ O ₄	0.866	0.771	0.675
1/2K ₂ CrO ₄	0.866	0.771	0.675
CH ₃ COOK	0.866	0.765	0.685
HCOOK	0.868	0.760	0.678
KBO ₂	0.814	0.677	0.560
1/2K ₂ HPO ₄	0.860	0.749	0.664
1/2Na ₂ S	0.887	0.795	0.726
KClO ₃ 0.25-norm.	0.927
KBrO ₃ 0.25-norm.	0.940
KIO ₃ 0.25-norm.	0.951

(Abegg & Riesenfeld, Z. phys. Ch. 1902, 40. 100.)

Solubility in salts + Aq at 35° C.

Salt	Concentration of the aq. solution	Mols. NH ₃ soluble in 1 liter of solution
KCl	0.5 normal	0.923
NaCl	"	0.966
CH ₃ COOK	"	0.902
1/2(COOK) ₂	"	0.902
KOH	"	0.870
NaOH	"	0.896
1/2K ₂ CO ₃	0.426 normal	0.914
1/2Na ₂ CO ₃	"	0.932

(Riesenfeld, Z. phys. Ch. 1903, 45. 462.)

The solubility of NH₃ in NaNO₃, NH₄NO₃, and in AgNO₃.2NH₃ + Aq is nearly the same as in pure H₂O. (Konowaloff, C. C. 1898, II. 659).

Distribution-coefficient of NH₃ between water and CHCl₃ = 26.3 at 20°; 24.9 at 25°; 23.2 at 30°.

The distribution-coefficient of NH₃ between CHCl₃ and a number of salt solutions has been determined for the purpose of studying the nature of metal-ammonia compounds in aqueous solution. (Dawson, Chem. Soc. 1900, 77. 1242.)

Distribution of NH₃ between H₂O and CHCl₃ at 18°.

NH ₃ concentration in aqueous solution. mols./litre	NH ₃ concentration in CHCl ₃ solution. mols./litre
0.9280	0.03506
1.921	0.07703
2.064	0.08350
2.274	0.09317
2.590	0.1083
3.700	0.1639
4.333	0.1996

(Dawson, Z. phys. Ch. 1909, 69. 120.)

Distribution of NH₃ between hydroxides + Aq and CHCl₃ at 18°.

Aqueous solution	NH ₃ concen- tration in the aqueous solution. mols./litre	NH ₃ concen- tration in the CHCl ₃ solution. mols./litre.
0.2-N. KOH	1.949	0.0841
0.5-N. KOH	1.978	0.0951
0.2-N. NaOH	2.016	0.0869
0.5-N. NaOH	1.944	0.0907
0.2-N. 1/2Ba(OH) ₂	2.076	0.08905
0.5-N. 1/2Ba(OH) ₂	3.397	0.1560

(Dawson, l.c.)

Distribution of NH₃ between Cu(OH)₂+Aq and CHCl₃ at 18°.

Conc. of Cu(OH) ₂ equivalents/litre	NH ₃ concentra- tion in aqueous solution. mols./litre	NH ₃ concentra- tion in CHCl ₃ solution. mols./litre
0.041	2.014	0.07968
0.0705	2.653	0.1087
0.081	3.011	0.1247

Dawson, l.c.)

Sol. in alcohol and ether.

Sol. in 3 pts. alcohol of 38°. (Boullay.)
1 vol. alcohol of 0.829 sp. gr. absorbs about 50 vols. NH₃. (Davy.)

Much less sol. in ethyl, propyl, or amyl alcohol than in H₂O. (Pagliano and Emo, Gazz. ch. it. 13. 278.)

Solubility of NH₃ in ethyl alcohol (absolute) at t°.

t°	% NH ₃	Pts. NH ₃ per 100 pts. alcohol
0	19.7	24.5
6	17.1	20.6
11.7	14.1	16.4
14.7	13.2	15.2
17	12.6	14.7
22	10.9	12.2
28.4	9.2	10.1

(de Bruyn, R. t. c. 11. 112.)

1 vol. abs. alcohol at 20° and 760 mm. pressure absorbs 340 vols. NH₃ gas. (Müller, W. Ann. 1891, 43. 567.)

1 l. methyl alcohol sat. with NH₃ contains 218 g. NH₃ at 0°; sp. gr. of solution = 0.770; coefficient of solubility = 425.0. (Delépine).

Solubility of NH₃ in alcohol at t°: weight NH₃ = weight NH₃ contained in a litre of solution sat. at 760 mm. and t°; sp. gr. = sp. gr. of solution; C = coefficient of solubility.

Temp.	Degree of Alcohol	100°	90°	80°	70°	60°	50°
0°	Weight NH ₃ .	130.5	146.0	206.5	246.0	304.5
	Sp. gr.	0.782	0.783	0.808	0.830	0.835
	C	209.5	245.0	390.0	504.5	697.7
10°	Weight NH ₃ .	108.5	120.0	167.0	198.25	227.0
	Sp. gr.	0.787	0.803	0.800	0.831	0.850
	C	164.3	186.0	288.0	373.0	438.6
20°	Weight NH ₃ .	75.0	97.5	119.75	137.5	152.5	182.7
	Sp. gr.	0.791	0.788	0.821	0.829	0.842	0.869
	C	106.6	147.8	190.5	223.0	260.8	338.2
30°	Weight NH ₃ .	51.5	74.0	81.75	100.3	129.5	152.0
	Sp. gr.	0.798	0.791	0.826	0.846	0.883
	C	97.0	186.7	121.6	211.6	252.0

(Delépine, J. Pharm. (5) 25. 496.)

Solubility of NH₃ in methyl alcohol (absolute) at t°.

t°	% NH ₃	Pts. NH ₃ per 100 pts. alcohol
0	29.3	41.5
6	26.0	35.2
11.7	23.5	30.7
14.7	21.8	27.9
17	20.8	26.3
22	18.3	22.4
28.4	14.8	17.4

(de Bruyn, l.c.)

Readily sol. in ether.
Sol. in 0.4 vol. petroleum from Amiano. (Saussure.)
1 vol. oil of turpentine absorbs 7.5 vols. NH₃ at 16°.
1 vol. oil of lemon absorbs 8.5 vols. NH₃ at 16°.

1 vol. oil of rosemary absorbs 9.75 vols. NH₃ at 29°.
1 vol. oil of lavender absorbs 47 vols. NH₃ at 20°. (Saussure.)
1 vol. caoutchine absorbs 3 vols. NH₃. (Himly.)
Valerol absorbs much NH₃. (Gerhardt, A. ch. (3) 7. 278.)
1 vol. ether at 760 mm. pressure absorbs 17.13 vols. NH₃ at 0°; 12.35 vols. at 10° and 10.27 vols. at 15°. (Christoff, Z. phys. Ch. 1912, 79. 459.)
+H₂O. Colorless crystals.
+½H₂O. Large transparent crystals. (Rupert, J. Am. Chem. Soc. 1909, 31. 868.)

Ammonia, with metal salts.
For the ammonia addition-products of metal salts, see under the respective metal salts, except in the case of Co, Cr, Hg, and the Pt metals, for which see cobalt ammonium, chromium ammonium, etc., compounds, for

further reference. New data on Co and Cr ammonium compounds and those of the Pt metals, published since the first edition, has not been included in the present edition.

ammonium amalgam, $\text{NH}_4, x\text{Hg}$.

Decomp. by H_2O , but more easily in presence of naphtha, alcohol, or ether.

ammonium azoimide, $\text{N}_4\text{H}_4 = \text{NH}_4\text{N}_3$.

Easily sol. in H_2O ; sl. sol. in absolute alcohol, easily in 80% alcohol. Insol. in ether or benzene. (Curtius, B. 24. 3344.)

ammonium cobalt azoimide, $\text{NH}_4\text{N}_3, \text{CoN}_6$.

Rather sol. in H_2O . (Curtius and Rissom, pr. 1898, (2) 58. 302.)

ammonium bromide, NH_4Br .

Easily sol. in H_2O with absorption of much heat.

1 pt. NH_4Br dissolves in pts. H_2O at t° .

t°	Pts. H_2O	t°	Pts. H_2O	t°	Pts. H_2O
10	1.51	30	1.23	100	0.78
16	1.39	50	1.06

(Eder, W. A. B. 82. (2) 1284.)

$\text{NH}_4\text{Br} + \text{Aq}$ containing 41.09% NH_4Br is sat. at 15° . (Gerlach.)

Sp. gr. of $\text{NH}_4\text{Br} + \text{Aq}$ at 15° .

% NH_4Br	Sp. gr.	% NH_4Br	Sp. gr.
5	1.0326	20	1.1285
10	1.0652	30	1.1921
15	1.0960	41.09	1.2920

(Eder.)

Sp. gr. of $\text{NH}_4\text{Br} + \text{Aq}$ at 16° .

% NH_4Br	Sp. gr.	% NH_4Br	Sp. gr.
2	1.0119	22	1.1375
3	1.0181	23	1.1440
4	1.0242	24	1.1506
5	1.0303	25	1.1573
6	1.0364	26	1.1642
7	1.0425	27	1.1713
8	1.0486	28	1.1787
9	1.0547	29	1.1862
10	1.0609	30	1.1938
11	1.0672	31	1.2018
12	1.0735	32	1.2098
13	1.0798	33	1.2180
14	1.0862	34	1.2260
15	1.0926	35	1.2342
16	1.0988	36	1.2425
17	1.1051	37	1.2509
18	1.1115	38	1.2594
19	1.1181	39	1.2679
20	1.1246	40	1.2765
21	1.1310	41	1.2850

(Hager, Comm. 1883.)

25 g. $\text{NH}_4\text{Br} + 50$ g. H_2O lower the temp. from 15.1° to -1.1° . (Rüdorff.)

Sol. in liquid NH_3 at -50° . (Moissan C. R. 1901, 133. 713.)

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.)

Sl. sol. in alcohol.

1 pt. NH_4Br dissolves in 32.3 pts. alcohol (0.806 sp. gr.) at 15° ; 9.5 pts. at 78° . (Eder, l.c.)

100 pts. absolute methyl alcohol dissolve 12.5 pts. at 19° ; 100 pts. absolute ethyl alcohol dissolve 3.22 pts. at 19° . (de Bruyn, Z. phys. Ch. 10. 783.)

Solubility in mixtures of methyl and ethyl alcohol at 25° .

P = % methyl alcohol in the solvent.

G = g. NH_4Br in 10 cc. of the solution.

S = sp. gr. of the sat. solution at $25^\circ/4^\circ$.

P	G	S
0.00	0.255	0.8065
4.37	0.299	0.8083
10.40	0.321	0.8117
41.02	0.506	0.8252
80.69	0.813	0.8501
84.77	0.847	0.8508
91.25	0.934	0.8551
100.00	0.983	0.8605

(Herz, Z. anorg. 1908, 60. 156.)

Solubility in mixtures of methyl and propyl alcohol at 25° .

P = % propyl alcohol in the solvent.

G = g. NH_4Br in 10 cc. of the solution.

S = Sp. gr. of the sat. solution at $25^\circ/4^\circ$.

P	G	S
0	0.983	0.8605
11.11	0.851	0.8524
23.8	0.690	0.8426
65.2	0.308	0.8184
91.8	0.128	0.8097
93.75	0.125	0.8089
100.	0.095	0.8059

(Herz, l.c.)

Solubility in mixtures of propyl and ethyl alcohol at 25° .

P = % propyl alcohol in the solvent.

G = g. NH_4Br in 10 cc. of the solution.

S = Sp. gr. of the sat. solution at $25^\circ/4^\circ$.

P	G	S
0	0.255	0.8065
8.1	0.251	0.8062
17.85	0.237	0.8052
56.6	0.163	0.8048
88.6	0.111	0.8042
91.2	0.105	0.8049
95.2	0.104	0.8059
100	0.095	0.8059

(Herz, l.c.)

Sol. in 809 pts. ether (0.729 sp. gr.). (Eder, l.c.)

Sol. in acetone. (Eidmann, C. C. 1899. II, 1014); (Naumann, B. 1904, 37. 4328.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethylacetate. (Naumann, B. 1910, 43. 314.)

Ammonium tribromide, NH_4Br_3 .

Gives off Br in air. Sol. in H_2O . (Rooseboom, B. 14. 2398.)

Decomp. in the air. Very sol. in H_2O . (Chattaway, Chem. Soc. 1915, 107. 106.)

Ammonium antimony bromide, $3\text{NH}_4\text{Br}$, 2SbBr_3 .

Easily sol. in abs. alcohol. (Caven, C. C. 1905. II, 293.)

$7\text{NH}_4\text{Br}$, 3SbBr_3 . Easily sol. in abs. alcohol. (Caven, C. C. 1905. II, 293.)

See also Bromantimonate, ammonium.

Ammonium bismuth bromide, NH_4Br , BiBr_3 , $+\text{H}_2\text{O}$.

Deliquescent. Decomp. by H_2O . Sol. in alcohol. (Nicklès, C. R. 51. 1097.)

Ammonium cadmium bromide, NH_4Br , CdBr_2 , $+\frac{1}{2}\text{H}_2\text{O}$.

Sol. in 0.73 pt. H_2O , 5.3 pts. abs. alcohol, 280 pts. ether (sp. gr. 0.729), and 24 pts. alcohol ether (1 : 1). (Eder, Dingl. 221. 89.)

Sol. in H_2O without decomp. between 1° and 110.1° .

100 pts. of the solution contain at:

t°	14.8°	52.2°	110.1°
53.82	58.01	65.32	75.83

pts. of the salt. (Rimbach, B. 1905, 38. 1555.)

$4\text{NH}_4\text{Br}$, CdBr_2 . Sol. in 0.96 pt. H_2O , from which it is pptd. by alcohol or ether. (Eder.)

Solubility in H_2O at t° .

Below 160° the salt is decomp. by H_2O ; at 160° it is sol. in H_2O without decomp.

t°	100 pts. of the solution contain			Solid phase
	Pts. Cd	Pts. Br	Pts. NH_4	
0.8	14.72	50.46	6.67	Double salt + NH_4Br
13.0	14.94	51.48	6.85	"
44.5	15.01	53.85	7.35	"
76.4	14.60	55.28	7.80	"
123.5	15.50	59.50	8.45	"
160.0	14.70	62.67	9.43	Double salt

(Rimbach, B. 1905, 38. 1558.)

Not sol. in $\text{HBr} + \text{Aq}$ without decomp. (Rimbach.)

Not sol. without decomp. in $\text{LiBr} + \text{Aq}$, $\text{CaBr}_2 + \text{Aq}$, $\text{MgBr}_2 + \text{Aq}$, $\text{NiBr}_2 + \text{Aq}$, or

$\text{CoBr}_2 + \text{Aq}$, even though very conc. solutions are used. Sol. without decomp. in $\text{ZnBr}_2 + \text{Aq}$. (Rimbach, B. 1905, 38. 1571.)

Ammonium chloromolybdenum bromide, $2\text{NH}_4\text{Br}$, $\text{Cl}_4\text{Mo}_3\text{Br}_3$.

Decomp. by pure H_2O . Can be crystallized from $\text{HBr} + \text{Aq}$. Apparently sol. without decomp. in alcohol. (Blomstrand.)

Ammonium cuprous bromide.

$4\text{NH}_4\text{Br}$, Cu_2Br_2 . Fairly stable in air.

$2\text{NH}_4\text{Br}$, $\text{Cu}_2\text{Br}_2 + \text{H}_2\text{O}$. Fairly stable in air. (Wells, Z. anorg. 1895, 10. 159.)

Ammonium cuprous bromide ammonia, NH_4Br , Cu_2Br_2 , 3NH_3 .

(Fleurent, C. R. 1891, 113. 1047.)

Ammonium cupric bromide, $2\text{NH}_4\text{Br}$, CuBr_2 , $+2\text{H}_2\text{O}$.

Very sol. in H_2O . (de Koninck, B. 21. 777 R.)

Ammonium iridium bromide.

See Bromiridate, ammonium.

Ammonium iron (ferric) bromide, $(\text{NH}_4)\text{FeBr}_4$, $+2\text{H}_2\text{O}$.

Very deliquescent; sol. in H_2O . (Walden, Z. anorg. 1894, 7. 332.)

Ammonium lead bromide, $12\text{NH}_4\text{Br}$, 7PbBr_2 , $+7\text{H}_2\text{O}$.

Decomp. on air, or with cold H_2O . (André, C. R. 96. 1502.)

$6\text{NH}_4\text{Br}$, $\text{PbBr}_2 + \text{H}_2\text{O}$. Decomp. by cold H_2O . (A.)

$7\text{NH}_4\text{Br}$, $\text{PbBr}_2 + 1\frac{1}{2}\text{H}_2\text{O}$. Stable on air; decomp. by cold H_2O . (A.)

None of the above compounds exist. (Wells, Sill. Am. J. 146. 25.)

$2\text{NH}_4\text{Br}$, PbBr_2 . Decomp. by H_2O . Sol. in conc. $\text{KOH} + \text{Aq}$ and in strong acids. (Fonze-Diacon, Bull. Soc. 1897, (3) 17.351.)

NH_4Br , 3PbBr_2 . (Wells.)

Ammonium magnesium bromide, NH_4Br , MgBr_2 , $+6\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . (Lerch, J. pr. (2) 28. 338.)

Ammonium mercuric bromide, 2HgBr_2 , NH_4Br .

Decomp. by H_2O into its constituent salts. (Ray, Chem. Soc. 1902, 81. 648.)

Ammonium molybdenum bromide, $2\text{NH}_4\text{Br}$, MoBr_3 , $+ \text{H}_2\text{O}$.

Easily sol. in H_2O . (Rosenheim, Z. anorg. 1905, 46. 322.)

Ammonium molybdenum bromide chloride.

See Ammonium chloromolybdenum bromide.

Ammonium osmium bromide.

See Bromosmate, ammonium.

Ammonium osmyl bromide, $(\text{NH}_4)_2\text{OsO}_2\text{Br}_4$.

Sol. in H_2O . (Wintrebert, A. ch. 1903, (7) 28. 95.)

Ammonium osmyl oxybromide,

(Wintrebert, A. ch. 1903 (7) 28. 117.)

Ammonium palladium bromide.

See Bromopalladate, ammonium, and Bromopalladite, ammonium.

Ammonium platinum bromide.

See Bromoplatinate, ammonium.

Ammonium rhodium bromide.

See Bromorhodite, ammonium.

Ammonium selenium bromide.

See Bromoselenate, ammonium.

Ammonium tellurium bromide.

See Bromotellurate, ammonium.

Ammonium thallic bromide, NH_4Br , TlBr_3 , + $2\text{H}_2\text{O}$.

Sol. in H_2O . (Willm.)

+ $4\text{H}_2\text{O}$. Efflorescent. Sol. in H_2O . (Nicklès.)

+ $5\text{H}_2\text{O}$. Sol. in H_2O . (Nicklès.)

Ammonium stannous bromide (ammonium bromostannite), NH_4Br , SnBr_2 , + H_2O .

Sol. in H_2O . (Benas, C. C. 1884. 958.)

$2\text{NH}_4\text{Br}$, SnBr_2 . Sol. in H_2O . (Raymann and Preis, A. 223. 323.)

+ H_2O . Sol. in H_2O . (Benas, l.c.)

+ $2\text{H}_2\text{O}$. (Richardson, Am. Ch. J. 14. 96.)

NH_4Br , 2SnBr_2 (?). (Benas.)

Ammonium stannic bromide, $2\text{NH}_4\text{Br}$, SnBr_4 .

See Bromostannate, ammonium.

Ammonium uranyl bromide, $2\text{NH}_4\text{Br}$, UO_2Br_2 , + $2\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O . (Sendtner.)

Ammonium zinc bromide, $2\text{NH}_4\text{Br}$, ZnBr_2 .

Deliquescent, and sol. in H_2O . (Bödeker, J. B. 1860. 17.)

+ H_2O . Very deliquescent, and sol. in H_2O . (André, A. ch. (6) 3. 104.)

+ $x\text{H}_2\text{O}$. (Ephraim, Z. anorg. 1908, 59. 66.)

$3\text{NH}_4\text{Br}$, ZnBr_2 . Sol. in H_2O . Decomp. only by great dilution. (Jones & Knight, Am. Ch. J. 1899, 22. 136.)

+ H_2O . Not hygroscopic. (Ephraim, Z. anorg. 1908, 59. 66.)

Ammonium bromide arsenic trioxide.

See Arsenite bromide, ammonium.

Ammonium bromide mercuric chloride, NH_4Br , 2HgCl_2 .

Ppt. (Ray, Chem. Soc. 1902, 81. 649.)

Ammonium bromide mercuric iodide, $2\text{NH}_4\text{Br}$, HgI_2 .

Decomp. by H_2O . Sol. in alcohol without decomp. (Grossmann, B. 1903, 36. 1602.)

$3\text{NH}_4\text{Br}$, 2HgI_2 . Decomp. by H_2O . Sol. in alcohol without decomp. (Grossmann, B. 1903, 36. 1602.)

Ammonium lead bromochloride, $\text{NH}_4\text{Pb}_2\text{Br}_4\text{Cl}$.

Decomp. by H_2O . (Fonzes-Diacon, Bull. Soc. 1897, (3) 17. 350.)

$\text{NH}_4\text{Pb}_2\text{Cl}_4\text{Br}$. Decomp. by H_2O . (Fonzes-Diacon, Bull. Soc. 1897, (3) 17. 349.)

Ammonium bromochloriodide, NH_4ClBrI .

Very stable; sol. in H_2O . (Chattaway, Chem. Soc. 1915, 107. 108.)

Ammonium lead bromiodide, NH_4PbBrI , + $2\text{H}_2\text{O}$ and $\text{NH}_4\text{Pb}_2\text{BrI}_4$.

Decomp. by H_2O . Sol. in conc. KOH + Aq and in strong acids. (Fonzes-Diacon, Bull. Soc. 1897, (3) 17. 352.)

Ammonium bromiodobromide, NH_4BrIBr .

Decomp. in the air. Sol. in ether. (Jackson, Am. Ch. J. 1900, 24. 28.)

Ammonium chloride, NH_4Cl .

(Sal-ammoniac.) Not deliquescent. Sol. in H_2O with reduction of temp.

Sol. in 2.24 pts. H_2O . (Wenzel.)

NH_4Cl + Aq sat. at 10° has sp. gr. = 1.072. (T.)

Sol. in 2.72 pts. cold, and 1 pt. boiling H_2O . (M. R., and P.)

Sol. in 3 pts. H_2O at 18.75° . (Abl.)

Sol. in 6 pts. cold, and 1 pt. boiling H_2O . (Fourcroy.)

100 pts. H_2O at 18.75° dissolve 36.75 pts. NH_4Cl . NH_4Cl + Aq sat. at its b.-pt. (114.2°) contains 88.9 pts. NH_4Cl in 100 pts. of the solution. (Berselius.)

100 pts. H_2O at 15° dissolve 33-36 pts.; and at 100° , 100 pts. NH_4Cl . (Ure's Dict.)

NH_4Cl + Aq sat. at 15° has sp. gr. = 1.075209, and contains at least 31.88 pts. NH_4Cl dissolved in every 100 pts. H_2O . (Michel and Krafft, A. ch. (3) 41. 478.)

NH_4Cl + Aq sat. at 10° contains 23.8% NH_4Cl . (Eller.)

NH_4Cl + Aq sat. in the cold contains 14.3% NH_4Cl . (Fourcroy.)

Sol. in 1 pt. H_2O at 113.5° , b.-pt. of sat. solution. (Griffiths.)

Sol. in 2.7 pts. H_2O at 18.75° , forming a liquid of 1.08 sp. gr. (Karsten, 1840.)

Sol. in 2.727 pts. H_2O at 10° . (Gren's Handbuch.)

100 pts. H_2O at 718 mm. pressure and t° dissolve pts. NH_4Cl .

t°	Pts. NH_4Cl	t°	Pts. NH_4Cl	t°	Pts. NH_4Cl	t°	Pts. NH_4Cl
0	28.40	30	41.72	60	55.04	90	68.36
10	32.84	40	46.16	70	59.48	100	72.80
20	37.28	50	50.60	80	63.92	110	77.24

(Alluard, C. R. 69. 500)

Solubility in 100 pts. H ₂ O at t°.							
t°	Pts. NH ₄ Cl	t°	Pts. NH ₄ Cl	t°	Pts. NH ₄ Cl	t°	Pts. NH ₄ Cl
0	29.7	30	41.4	60	55.2	90	71.3
1	30.0	31	41.8	61	55.7	91	71.9
2	30.3	32	42.2	62	56.2	92	72.5
3	30.6	33	42.7	63	56.7	93	73.1
4	31.0	34	43.1	64	57.2	94	73.7
5	31.4	35	43.6	65	57.7	95	74.3
6	31.8	36	44.0	66	58.2	96	74.9
7	32.2	37	44.4	67	58.7	97	75.5
8	32.6	38	44.9	68	59.2	98	76.1
9	33.0	39	45.3	69	59.7	99	76.7
10	33.3	40	45.8	70	60.2	100	77.3
11	33.7	41	46.2	71	60.7	101	78.0
12	34.1	42	46.7	72	61.2	102	78.6
13	34.5	43	47.1	73	61.7	103	79.2
14	34.8	44	47.6	74	62.3	104	79.9
15	35.2	45	48.0	75	62.8	105	80.5
16	35.6	46	48.5	76	63.4	106	81.2
17	36.0	47	49.0	77	63.9	107	81.8
18	36.4	48	49.5	78	64.5	108	82.5
19	36.8	49	49.9	79	65.1	109	83.1
20	37.2	50	50.4	80	65.6	110	83.8
21	37.6	51	50.9	81	66.2	111	84.4
22	38.0	52	51.3	82	66.7	112	85.1
23	38.4	53	51.8	83	67.3	113	85.7
24	38.8	54	52.3	84	67.8	114	86.4
25	39.3	55	52.8	85	68.4	115	87.1
26	39.7	56	53.2	86	69.0	115.65	87.3
27	40.1	57	53.7	87	69.6
28	40.5	58	54.2	88	70.2
29	40.9	59	54.7	89	70.7

(Mulder, calculated from his own and other observations. Scheik. Verhandel. 1864. 57.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. NH ₄ Cl	t°	Pts. NH ₄ Cl	t°	Pts. NH ₄ Cl
0	29.7	10.8	33.9	64.9	57.9
6.2	32.2	31.6	42.2	90.6	67.2

(Lindström, Pogg. 136. 315.)

NH₄Cl + Aq sat. at 13–16° contains 26.16% NH₄Cl. (v. Hauer, J. pr. 103. 114.)

Sol. in 2.72 pts. H₂O at 19°. (Schiff, A. 100. 326.)

Sol. in 2.803 pts. H₂O at 15°. (Gerlach.)

Sat. NH₄Cl + Aq at 75° contains 38.23% NH₄Cl. (Tschugaeff, Z. anorg. 1914. 86. 161.)

NH₄Cl + Aq sat. at 30° contains 29.5% NH₄Cl. (Meerburg, C. C. 1904. II, 1362.)

Solubility in H₂O at t°.

t°	1000 mols. H ₂ O dissolve mols. NH ₄ Cl	100 g. H ₂ O dissolve g. NH ₄ Cl
3.5	105.2	31.25
25.0	129.7	38.5
50.0	167.0	49.6

(Biltz and Marcus, Z. anorg. 1911, 71. 169.)

Solubility of NH ₄ Cl in H ₂ O at t°.		
t°	g. NH ₄ Cl in 100 g. of the solution	Solid phase
— 0.45	0.7 ^s	Ice
— 1.25	1.9 ^s	"
— 1.70	2.7 ^s	"
— 3.05	4.6	"
— 4.45	6.6 ^s	"
— 6.4	9.2 ^s	"
— 8.25	11.4	"
— 9.7	13.1	"
— 11.9	15.3	"
— 13.25	16.7	"
— 14.70	18.1 ^s	"
— 15.4	18.9	"
± 16.0	± 19.5	Ice + NH ₄ Cl
— 15.0	19.7	NH ₄ Cl
— 12.2	20.0	"
— 10.9	20.3	"
— 7.4	21.1	"
— 5.7	21.7	"
— 2.3	22.3	"
± 1.1	22.6	"
0	22.7	"

(Meerburg, Z. anorg. 1903, 37. 203.)

100 g. H₂O dissolve 29.5 g. NH₄Cl at 30°. (Schreinemakers, Arch. neer. Sc. (2) 15. 17.)

Spec. gravity of NH₄Cl + Aq. G = according to Gerlach at 15° (Z. anal. 8. 281); S = according to Schiff at 19° (A. 110. 74).

% NH ₄ Cl	Sp. gr.		% NH ₄ Cl	Sp. gr.	
	G	S		G	S
1	1.00316	1.0029	17	1.05086	1.0495
2	1.00632	1.0058	18	1.05367	1.0523
3	1.00948	1.0087	19	1.05648	1.0551
4	1.01264	1.0116	20	1.05929	1.0579
5	1.01580	1.0145	21	1.06204	1.0606
6	1.01880	1.0174	22	1.06479	1.0633
7	1.02180	1.0203	23	1.06754	1.0660
8	1.02481	1.0233	24	1.07029	1.0687
9	1.02781	1.0263	25	1.07304	1.0714
10	1.03081	1.0293	26	1.07375	1.0741
11	1.03370	1.0322	26.297	1.07658
12	1.03658	1.0351	27		1.0768
13	1.03947	1.0380	28		1.0794
14	1.04325	1.0409	29		1.0802
15	1.04524	1.0438	30		1.0846
16	1.04805	1.0467			...

For older determinations, see Storer's Dict.

Sp. gr. of NH₄Cl + Aq at 18°.

% NH ₄ Cl	Sp. gr.	% NH ₄ Cl	Sp. gr.
5	1.0142	20	1.0571
10	1.0289	25	1.0710
15	1.0430

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. at 20°/4° of a normal solution of NH_4Cl = 1.01454. (Haigh, J. Am. Chem. Soc. 1912, **34**. 1151.)

NH_4Cl + Aq containing 6.52% NH_4Cl has sp. gr. 20°/20° = 1.0195. (Le Blanc & Rohland, Z. phys. Ch. 1896, **19**. 272.)

Temp. of maximum density of NH_4Cl + Aq	g. mol. NH_4Cl in 1000 g. H_2O
2.640°	0.1899
0.055°	0.5407

(de Coppet, C. R. 1900, **131**. 178.)

Sp. gr. of dil. NH_4Cl + Aq at 20.004° and 731 mm. (corr.)

Conc. = g. equiv. NH_4Cl per l. at 20.004°.

Conc.	Sp. gr.
0.0000	1.000,000,0
0.0001	1.000,001,8
0.0002	1.000,003,7
0.0005	1.000,009,3
0.0010	1.000,018,5
0.0020	1.000,036,9
0.0050	1.000,091,3
0.0100	1.000,180,3

(Lamb and Lee, J. Am. Chem. Soc. 1913, **35**. 1688.)

Sp. gr. of dil. NH_4Cl + Aq.

NH_4Cl g. in 1000 g. of solution	Sp. gr. 16°/16°
0	1.000000
0.4431	1.000150
0.9061	1.000304
1.8085	1.000606
3.5947	1.001196
7.7845	1.002562
15.3425	1.004994
31.2364	1.010018

(Dijken, Z. phys. Ch. 1897, **24**. 107.)

B.-pt. of NH_4Cl + Aq, containing pts. NH_4Cl to 100 pts. H_2O . G = according to Gerlach (Z. anal. **26**. 439); L = according to Legrand (A. ch. (2) **59**. 436).

B.-pt.	G	L	B.-pt.	G	L
101°	6.5	7.8	109°	50.6	53.5
102	12.8	13.9	110	56.2	59.9
103	19.0	19.7	111	61.9	66.4
104	24.7	25.2	112	67.8	73.3
105	29.7	30.5	113	74.2	80.5
106	34.6	35.7	114	81.3	88.1
107	39.6	41.3	114.2	...	88.9
108	45.0	47.3	114.8	87.1	...

Sat. NH_4Cl + Aq boils at 115.8° at 718 mm. pressure. (Alluard, C. R. **59**. 500.)

NH_4Cl + Aq containing 74.2 pts. NH_4Cl to 100 pts. H_2O forms a crust at 113°; highest temperature observed, 114.8°. (Gerlach, Z. anal. **26**. 426.)

NH_4Cl + Aq containing 10% NH_4Cl boils at 101.7°; 20% NH_4Cl , at 104.4°. (Gerlach.)

NH_4Cl + Aq containing 10.6% NH_4Cl gives off NH_3 at 37°. (Leeds, Am. J. Sci. (3) **7**. 197.)

When NH_4Cl + Aq is boiled, or even evap. on water bath, a little NH_3 is expelled. (Fresenius.)

30 pts. NH_4Cl mixed with 100 pts. H_2O lower the temp. from 13.3° to -5.1°, that is 18.4°. (Rüdorff, B. **2**. 68.)

Freezing-point of sat. solution is -15.4°, the same temp. which is caused by mixing 25 pts. NH_4Cl with 100 pts. snow. (Rüdorff, Pogg. **122**. 337.)

Conc. HCl + Aq precipitates part of NH_4Cl from sat. NH_4Cl + Aq. (Vogel, J. pr. **2**. 199.)

Solubility of NH_4Cl in HCl + Aq at 0°. NH_4Cl = mols. NH_4Cl (in milligrammes) dissolved in 10 cc. of the liquid; HCl = mols. HCl (in milligrammes) dissolved in 10 cc. of the liquid.

NH_4Cl	HCl	Sum of mols.	Sp. gr.
46.125	0.0	46.125	1.076
43.6	2.9	46.5	1.0695
41.0	5.5	46.5	1.0705
39.15	7.85	47.0	1.0715
36.45	10.85	47.30	1.073
27.37	21.4	48.77	1.078
10.875	53.0	63.875	1.106
8.8	61.0	69.8	1.114

(Engel, Bull. Soc. (2) **45**. 655.)

Solubility of NH_4Cl in HCl + Aq.

t°	HCl concentration. g. mol. per 100 g. H_2O	Weight NH_4Cl dissolved in 1000 g. H_2O	Molecular solubility
0°	0	298.40	5.59
"	1/4	286.43	5.36
"	1/2	271.23	5.08
"	1	245.35	4.60
25°	0	395.10	7.40
"	1/4	380.85	7.13
"	1/2	366.00	6.85
"	1	339.05	6.35

(Armstrong & Eyre, Proc. R. Soc. (A.) **84**. 127.)

Solubility in $\text{NH}_4\text{OH} + \text{Aq.}$ $\text{NH}_4\text{Cl} = \text{mols.}$
 NH_4Cl (in mgs.) in 10 cc. solution;
 $\text{NH}_3 = \text{mols.}$ NH_3 (in mgs.) in 10 cc.
 solution.

NH_4Cl	NH_3	Sp. gr.
46.125	0	1.076
45.8	5.37	1.067
45.5	12.025	1.054
45.125	23.4	1.044
44.5	38.0	1.031
44.0	47	1.025
43.625	54.5	1.017
43.125	80.0	0.993
44.0	90.0	0.992
44.375	95.5	0.983
49.75	130	0.953
60.0	169.75	0.931

(Engel, Bull. Soc. (3) 6. 17.)

$\text{NH}_4\text{Cl} + \text{BaCl}_2$. 100 pts. H_2O dissolve 33.8
 pts. $\text{NH}_4\text{Cl} + 11.6$ pts. BaCl_2 at 20° . (Rüdorff,
 Pogg. 143. 467.)

Solubility of NH_4Cl and BaCl_2 in H_2O .

t°	Wt. per 100		Solid phase
	NH_4Cl	BaCl_2	
-16.2°	16.10	8.07	$\text{NH}_4\text{Cl} +$ $\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$
0	19.26	8.22	
30	24.89	8.19	
40	28.92	8.40	
50	30.63	8.55	

(Schreinemaker, Chem. Weekbl. 1910, 7. 333.)
 See also $\text{BaCl}_2 + \text{NH}_4\text{Cl}$ under BaCl_2 .

$\text{NH}_4\text{Cl} + \text{CdCl}_2$. Solubility of NH_4Cl and
 CdCl_2 .
 See Ammonium cadmium chloride.

$\text{NH}_4\text{Cl} + \text{CuCl}_2$. Solubility of NH_4Cl in
 H_2O at 30° in presence of varying amounts of
 CuCl_2 .

% by wt. CuCl_2	% by wt. NH_4Cl	Solid phase
0	29.5	NH_4Cl
1.9	28.6	$\text{NH}_4\text{Cl} + \text{CuCl}_2 \cdot 2\text{NH}_4\text{Cl} \cdot 2\text{H}_2\text{O}$
3.8	25.9	$\text{CuCl}_2 \cdot 2\text{NH}_4\text{Cl} \cdot 2\text{H}_2\text{O}$
7.7	19.8	"
10.5	16.5	"
12.3	14.9	"
15.6	12.1	"
19.9	9.4	"
24.0	7.1	"
29.4	4.9	"
35.1	3.4	"
41.4	2.1	"
43.2	2.0	$\text{CuCl}_2 \cdot 2\text{NH}_4\text{Cl} \cdot 2\text{H}_2\text{O} + \text{CuCl}_2 \cdot 2\text{H}_2\text{O}$
43.9	0.	$\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$

(Meerburg, Z. anorg. 1905, 45. 3)

$\text{NH}_4\text{Cl} + \text{PbCl}_2$. Solubility of NH_4Cl and
 PbCl_2 in H_2O at 22° .

g. equivalent in 1000 g. H_2O		Solid phase
NH_4Cl	PbCl_2	
0.0	0.0749	PbCl_2
0.1	0.0325	"
0.2	0.0194	"
0.3	0.0153	"
0.4	0.0138	"
0.5	0.0130	"
0.52	0.0127	$\text{PbCl}_2 + \text{NH}_4\text{Cl} \cdot 2\text{PbCl}_2$
0.55	0.0123	$\text{NH}_4\text{Cl} \cdot 2\text{PbCl}_2$
0.6	0.0113	"
0.65	0.0105	"
0.7	0.0099	"
0.8	0.0087	"
0.9	0.0083	"
1.0	0.0080	"
1.2	0.0075	"
1.5	0.0073	"
2.0	0.0077	"
2.5	0.0092	"
3.0	0.0112	"
4.0	0.0182	"
5.0	0.0296	"
6.0	0.0473	"
7.0	0.0774	"
7.29	0.0828	$\text{NH}_4\text{Cl} + \text{NH}_4\text{Cl} \cdot 2\text{PbCl}_2$
7.29	0.0000	NH_4Cl

(Brönstedt, Z. phys. Ch. 1911, 77. 132.)

Solubility of NH_4Cl and $2\text{PbCl}_2 \cdot \text{NH}_4\text{Cl}$ in
 H_2O at 100° .

NH_4Cl g. equivalent		PbCl_2 g. equivalent		Solid phase
in 1000 g. solution	in 1000 g. H_2O	in 1000 g. solution	in 1000 g. H_2O	
1.277	1.404	0.160	0.176	NH_4Cl $+ 2\text{PbCl}_2 \cdot \text{H}_2\text{O}$

(Brönstedt, l. c.)

$\text{NH}_4\text{Cl} + \text{MgCl}_2$. Solubility of NH_4Cl and
 $\text{NH}_4\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$.

t°	In 1000 g. mols. H_2O		Solid phase
	NH_4Cl	MgCl_2	
3.5°	27.5	55.7	$\text{NH}_4\text{Cl} + \text{NH}_4\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$
25	42.1	56.4	"
50	62.9	59.1	"

(Bilts and Marcus, Z. anorg. 1911, 71. 170.)

Solubility of $\text{NH}_4\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$ and $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$.

t°	In 1000 g. mol. H_2O		Solid phase
	g. mol. NH_4Cl	g. mol. MgCl_2	
3.5°	0.5	99.5	$\text{MgCl}_2 \cdot 6\text{H}_2\text{O} + \text{NH}_4\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$
25°	0.5	103.8	
50°	0.8	111.2	

(Biltz and Marcus, Z. anorg. 1911, 71. 170.)

$\text{NH}_4\text{Cl} + \text{NH}_4\text{NO}_3$. 100 pts. H_2O dissolve 29.1 pts. $\text{NH}_4\text{Cl} + 173.8$ pts. NH_4NO_3 at 19.5° . (Rüdorff, B. 6. 482.)

$\text{NH}_4\text{Cl} + \text{Ba}(\text{NO}_3)_2$. 100 pts. H_2O dissolve at 18.5° —

	1	2	3	4	5
NH_4Cl	36.7	38.6	38.06	39.18	...
$\text{Ba}(\text{NO}_3)_2$	8.6	16.73	17.02	8.9

2, sat. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ treated with NH_4Cl ; 3, sat. $\text{NH}_4\text{Cl} + \text{Aq}$ treated with $\text{Ba}(\text{NO}_3)_2$; 4, simultaneous treatment of both salts with H_2O . (Karsten.)

$\text{NH}_4\text{Cl} + \text{KNO}_3$. 100 pts. H_2O dissolve at 18.5° —

	1	2	3	4	5	6
KNO_3	29.9	30.56	37.68	38.62	34.2
NH_4Cl	44.33	37.98	39.84	36.7	38.8
		74.89	75.66	78.46		73.0

1 and 5, according to Mulder; 2, sat. $\text{KNO}_3 + \text{Aq}$ treated with NH_4Cl ; 3, sat. $\text{NH}_4\text{Cl} + \text{Aq}$ treated with KNO_3 ; 4, simultaneous treatment of NH_4Cl and KNO_3 (Karsten); 6, by warming solution with excess of both salts, and cooling to 14.8° . The amount of excess of one or the other salt has no influence. (Rüdorff.)

$\text{NH}_4\text{Cl} + \text{NaNO}_3$. Slowly sol. in sat. $\text{NaNO}_3 + \text{Aq}$, at first to a clear solution, but afterwards NaCl separates out. (Karsten.)

$\text{NH}_4\text{Cl} + \text{KCl}$. 100 pts. H_2O dissolve—

	(Rüdorff) 15°	(Karsten) 18.75°		
KCl	16.97	34.4	16.27
NH_4Cl	28.90	29.83	37.02

	(Rüdorff) 22°	(Mulder) At b.-pt.		
KCl	19.1	58.5	21.9
NH_4Cl	30.4	67.7	87.3

100 pts. sat. solution of $\text{NH}_4\text{Cl} + \text{KCl}$ contain 30.61 pts. of the two salts at $13-16^\circ$. (v. Hauer, J. pr. 103. 114.)

$\text{NH}_4\text{Cl} + \text{NaCl}$. 100 pts. H_2O dissolve—

	10-20°	(Mulder) 10°	10°	(v. Hauer) $13-16^\circ$
NH_4Cl	19.50	33.3	18.8-20.3
NaCl	35.8	30.00	24.6-26.1
		49.50		43.4-46.4

	(Karsten) 18.75°	(Rüdorff) 18.7°	(Mulder) At b.-pt.
NH_4Cl	22.06	37.02	22.9
NaCl	26.38	23.9
	48.44	46.8	100.8

Sp. gr. of sat. solution of $\text{NH}_4\text{Cl} + \text{NaCl}$ is 1.179. (Karsten.)

$\text{NH}_4\text{Cl} + (\text{NH}_4)_2\text{SO}_4$. 100 pts. H_2O dissolve 26.8 pts. $\text{NH}_4\text{Cl} + 46.5$ pts. $(\text{NH}_4)_2\text{SO}_4$ at 21.5° . (Rüdorff, B. 6. 484.)

Solubility in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ at 30° .

Composition of the solution		Solid phase
% by wt. NH_4Cl	% by wt. $(\text{NH}_4)_2\text{SO}_4$	
0	44	$(\text{NH}_4)_2\text{SO}_4$
6.86	36.15	"
14.62	28.6	"
17.60	25.69	$(\text{NH}_4)_2\text{SO}_4 + \text{NH}_4\text{Cl}$
17.93	25.81	"
19.07	23.22	NH_4Cl
19.97	21.3	"
22.3	16.33	"
24.06	12.72	"
29.5	0	"

(Schreinemakers, Z. phys. Ch. 1909, 69. 562.)

$\text{NH}_4\text{Cl} + \text{CuSO}_4$. Sol. in sat. $\text{CuSO}_4 + \text{Aq}$, at first to a clear solution, but a double sulphate of NH_4 and Cu soon separates. (Karsten.)

$\text{NH}_4\text{Cl} + \text{MgSO}_4$. Slowly and difficultly sol. in sat. $\text{MgSO}_4 + \text{Aq}$ with subsequent separation of double sulphate. (Karsten.)

$\text{NH}_4\text{Cl} + \text{K}_2\text{SO}_4$. 100 pts. H_2O dissolve, at 18.75° —

		a	b	c	
K_2SO_4	10.8	11.1	13.26	13.28
NH_4Cl	38.2	37.94	37.92	36.7
		49.3	51.20	51.20	

In (a) NH_4Cl was added to sat. $\text{K}_2\text{SO}_4 + \text{Aq}$. In (b) K_2SO_4 was added to sat. $\text{NH}_4\text{Cl} + \text{Aq}$. In (c) NH_4Cl and K_2SO_4 were treated together with H_2O . (Karsten.)

100 pts. H₂O at 14° dissolve 14.1 pts. K₂SO₄+36.8 pts. NH₄Cl=50.9 pts. K₂SO₄+NH₄Cl, under all conditions. (Rüdorff, Pogg. 148. 565.)

100 pts. H₂O dissolve at b.-pt.—

K ₂ SO ₄ . . .	26.75	33.3– 33.9	87.3
NH ₄ Cl	90.4–111.8	
		123.7–145.7	

(Mulder.)

NH₄Cl+Na₂SO₄. 100 pts. H₂O dissolve 28.9 pts. NH₄Cl+24.7 pts. Na₂SO₄, if NH₄Cl +Aq sat. at 10° is sat. with Na₂SO₄ at 11°. 100 pts. H₂O dissolve 31.8 pts. NH₄Cl+9.0 pts. Na₂SO₄, if Na₂SO₄+Aq sat. at 10° is sat. with NH₄Cl at 11°. (Mulder, J. B. 1866. 68.)

Sol. in sat. Na₂SO₄+Aq. (Karsten.)
Sol. in sat. ZnSO₄+Aq. (Karsten.)

Sl. sol. in liquid NH₃ at –50°. (Moissan, C. R. 1901, 133. 713.)
Very sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 826.)

Very sl. sol. in absolute alcohol.
100 pts. alcohol of 0.939 sp. gr. dissolve—
at 4° 8° 27° 38° 56°
11.2 12.6 19.4 23.6 30.1 pts. NH₄Cl.
(Gerardin, A. ch. (4) 5. 129.)
14 pts. boiling highest rectified spirit dissolve 1 pt. NH₄Cl. (Wenzel.)
100 pts. alcohol of—
0.900 sp. gr. dissolve 0.5 pts. NH₄Cl.
0.872 " " " 4.75 " "
0.834 " " " 1.5 " "
(Kirwan.)

Though somewhat sol. in pure absolute alcohol, NH₄Cl is absolutely insol. in alcohol in presence of methyl amine chlorides. (Winkles, A. 93. 324.)
100 pts. absolute methyl alcohol dissolve 3.35 pts. at 19°.
100 pts. absolute ethyl alcohol dissolve 0.62 pt. at 19°. (de Bruyn, Z. phys. Ch. 10. 783.)

Solubility of NH₄Cl in methyl alcohol.

t°	Alcohol concentration, mol. g. alcohol for 1000 g. H ₂ O	Solubility in 1000 g. H ₂ O	Molecular solubility
0°	0	298.40	5.59
"	1/4	297.35	5.57
"	1/2	296.55	5.55
"	1	292.65	5.47
"	3	283.15	5.30
25°	0	395.10	7.40
"	1/4	394.75	7.39
"	1/2	393.85	7.37
"	1	392.90	7.36
"	3	386.20	7.23

(Armstrong and Eyre, Proc. R. Soc. Lond. (A) 84. 127.)

Solubility of NH₄Cl in ethyl alcohol at 0°.

Alcohol concentration, mol. g. alcohol for 1000 g. H ₂ O	Solubility in 1000 g. H ₂ O	Molecular solubility
0	298.40	5.59
1/4	295.50	5.53
1/2	291.95	5.47
1	286.40	5.37
3	266.25	4.99

(Armstrong and Eyre, l.c.)

See also ammonium cupric chloride.

Solubility of NH₄Cl in propyl alcohol.

t°	Alcohol concentration, mol. g. alcohol for 1000 g. H ₂ O	Solubility in 1000 g. H ₂ O	Molecular solubility
0°	0	298.46	5.59
"	1/4	295.40	5.53
"	1/2	291.30	5.45
"	1	284.00	5.32
25°	0	395.10	7.40
"	1/4	393.50	7.37
"	1/2	390.80	7.32
"	1	384.80	7.21

(Armstrong and Eyre, l.c.)

Solubility in mixtures of methyl and ethyl alcohol at 25°.
P = % methyl alcohol in the solvent.
G = g. NH₄Cl in 10 cc. of the solution.
S = sp. gr. of the sat. solution at 25°/4°.

P	G	S
0.00	0.0533	0.7908
4.37	0.0583	0.7909
10.40	0.0658	0.7910
41.02	0.118	0.7957
80.69	0.217	0.8020
84.77	0.227	0.8026
91.25	0.247	0.8040
100.00	0.276	0.8062

(Herz, Z. anorg. 1908, 60. 155.)

Solubility in mixtures of methyl and propyl alcohol at 25°.
P = % propyl alcohol in the solvent.
G = g. NH₄Cl in 10 cc. of the solution.
S = Sp. gr. of the sat. solution at 25°/4°.

P	G	S
0	0.276	0.8062
11.11	0.231	0.8035
23.8	0.182	0.8008
65.2	0.071	0.8005
91.8	0.026	0.8002
93.75	0.023	0.8000
100.00	0.018	0.8009(?)

(Herz, Z. anorg. 1908, 60. 157.)

Solubility in mixtures of propyl and ethyl alcohol at 25°.

P = % propyl alcohol in the solvent.

G = g. NH_4Cl in 10 cc. of the solution.

S = Sp. gr. of the sat. solution at 25°/4°.

P	G	S
0	0.0533	0.7908
8.1	0.0505	0.7910
17.85	0.0455	0.7916
56.6	0.0312	0.7963
88.6	0.0210	0.7996
91.2	0.0203	0.8001
95.2	0.0190	0.8003
100	0.0177	0.8009

(Herz, Z. anorg. 1908, 60. 160.)

Insol. in ether and CS_2 . (Fordos and Gélis, A. ch. (3) 32. 393.)

Very sl. sol. in acetone. (Krug and M'Elroy, J. anal. appl. Ch. 6. 184.)

Solubility of NH_4Cl in acetone + Aq at 25°.

A = cc. acetone in 100 cc. acetone + Aq.

NH_4Cl = millimols. NH_4Cl in 100 cc. of the solution.

A	NH_4Cl	Sp. gr.
0	585.1	1.0793
10	534.1	1.0618
20	464.6	1.0451
30	396.7	1.0263
40	328.5	0.99984
46.5	283.7	0.97998
to	18.9	0.8390
85.7		
90	9.4	0.8274

(Herz, Z. anorg. 1905, 45. 263.)

Solubility of NH_4Cl in glycerine + Aq at 25°.

G = g. glycerine in 100 g. glycerine + Aq.

NH_4Cl = millimols. NH_4Cl in 100 cc. of the solution.

G	NH_4Cl	Sp. gr.
0	585.1	1.0793
13.28	544.6	1.0947
25.98	502.9	1.1127
45.36	434.4	1.1452
54.23	403.5	1.1606
83.84	291.4	1.2225
100	228.4	1.2617

(Herz, l.c.)

Insol. in acetone. (Naumann, B. 1904, 37. 4328.); (Eidmann, C. C. 1899. II, 1014.)

Insol. in anhydrous pyridine. Sol. in 97% pyridine + Aq, 95% pyridine + Aq and in 93% pyridine + Aq. (Kahlenberg, J. Am. Chem. Soc. 1908, 30. 1107.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Very sol. in ethyl amine. (Shinn, J. phys. Chem. 1907, 11. 538.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Sol. in formic acid. (Zanninovich-Tessarini, Z. phys. Ch. 1896, 19. 251.)

Ammonium antimony chloride, $\text{SbCl}_4(\text{NH}_4)_2$, $\text{SbCl}_4(\text{NH}_4)_3$.

Ppt. Decomp. by H_2O . (Weinland, B. 1905, 38. 1085.)

$\text{SbCl}_4(\text{NH}_4)_2$, SbCl_5 , NH_4OH . Very deliquescent; sl. sol. in H_2O with decomp. (Weinland, B. 1901, 34. 2635.)

Ammonium antimonous chloride, NH_4Cl , SbCl_3 .

Deliquescent. (Dehérain, C. R. 52. 734.)

$2\text{NH}_4\text{Cl}$, $\text{SbCl}_3 + 2\text{H}_2\text{O}$. Permanent in dry air; decomp. by much H_2O . (Poggiale.)

$3\text{NH}_4\text{Cl}$, $\text{SbCl}_3 + 3\text{H}_2\text{O}$. As above.

Ammonium antimonic chloride, $3\text{NH}_4\text{Cl}$, SbCl_5 .

Decomp. by H_2O . (Dehérain, C. R. 52. 734.)

$4\text{NH}_4\text{Cl}$, SbCl_5 . Decomp. by H_2O . (D.)

See also Chlorantimonate, ammonium.

Ammonium antimony platinum chloride, $(\text{Sb}, \text{Pt})\text{Cl}_4(\text{NH}_4)_2$.

Ppt. (Weinland, B. 1905, 38. 1084.)

Ammonium antimony tin chloride, $(\text{Sb}, \text{Sn})\text{Cl}_4(\text{NH}_4)_2$.

Ppt. (Weinland, B. 1905, 38. 1085.)

Ammonium arsenyl chloride, $2\text{NH}_4\text{Cl}$, $\text{AsOCl}_2 + \frac{1}{2}\text{H}_2\text{O}$.

(Wallace, Phil. Mag. (4) 16. 358.)

Ammonium bismuth chloride, NH_4Cl , 2BiCl_3 .

Deliquescent. (Dehérain, C. R. 54. 724.)

$2\text{NH}_4\text{Cl}$, BiCl_3 . Decomp. by H_2O . (Arppe.) Pogg. 64. 237.)

$+ 2\frac{1}{2}\text{H}_2\text{O}$. (Rammelsberg.)

$3\text{NH}_4\text{Cl}$, BiCl_3 . Decomp. by H_2O . (Arppe.)

$5\text{NH}_4\text{Cl}$, 2BiCl_3 . (Rammelsberg.)

Ammonium bismuth potassium chloride, $2\text{NH}_4\text{Cl}$, BiCl_3 , KCl .

(Dehérain, C. R. 54. 724.)

Ammonium cadmium chloride, NH_4Cl , CdCl_2 .

Solubility of NH_4Cl , CdCl_2 in H_2O at t°.

t°	Pts. by weight in 100 pts. of solution			g. in 100 g. solution	Grams in 100 H_2O	Mols. H_2O free salt dissolved by 100 mols. H_2O
	Cl	Cd	NH_4			
2.4°	13.44	14.26	2.24	29.94	42.74	3.25
16.0	15.07	15.82	2.56	33.45	50.26	3.83
41.2	17.46	18.61	2.89	38.96	63.83	4.86
63.8	19.73	20.92	3.34	43.99	78.54	5.98
105.9	23.52	24.70	4.01	52.58	109.33	8.30

(Rimbach, B. 1897, 30. 3076.)

+ $\frac{1}{2}$ H₂O. Sl. sol. in H₂O, alcohol, and wood spirit. (v. Hauer, W. A. B. 13. 449.)

4NH₄Cl, CdCl₂. Sol. in H₂O. (v. Hauer.)

Decomp. by H₂O to NH₄Cl, CdCl₂. Decomp. increases with decrease of temp. At 3.9° approximately wholly decomp. to NH₄Cl, CdCl₂. At 113.9° very nearly all is 4NH₄Cl, CdCl₂. (Rimbach, B. 1897, 30. 3077.)

Solubility of 4NH₄Cl, CdCl₂ in H₂O at t°.

t°	Pta. dissolved in 100 pta. by weight of solution.		
	Cd	Cl	NH ₄
3.9	5.75	18.17	7.37
16.1	6.93	20.26	7.97
40.2	9.91	23.84	8.92
58.5	12.50	26.53	9.35
112.9	16.66	31.79	10.78
113.9	16.51	32.71	11.30

(Rimbach, B. 1897, 30. 3071.)

Sol. without decomp. in 37.3% HCl(d=1.19) and 24.8% HCl(d=1.125). (Rimbach, B. 1905, 38. 1569.)

Solubility of 4NH₄Cl, CdCl₂+NH₄Cl in H₂O at t°.

t°	In 100 pta. by wt. of the solution			Composition of the solid phase	
	Pts. by wt. Cd	Pts. by wt. Cl	Pts. by wt. NH ₄	Mol. % NH ₄ Cl	Mol. % Tetra-salt
1.0	2.82	17.11	7.82	59.0	41.0
13.2	2.76	18.84	8.71	74.0	26.0
40.1	3.16	22.56	10.49	71.0	29.0
58.2	3.51	25.21	11.72	69.0	31.0

(Rimbach, B. 1902, 35. 1300.)

Solubility of 4NH₄Cl, CdCl₂+NH₄Cl, CdCl₂ in H₂O at t°.

t°	In 100 pta. by wt. of the solution			Composition of the solid phase	
	Pts. by wt. Cd	Pts. by wt. Cl	Pts. by wt. NH ₄	Mol. % Mono-salt	Mol. % Tetra-salt
1.1	5.34	17.62	7.27	49.6	50.4
14.0	7.12	19.86	7.84	47.0	53.0
40.7	10.24	23.82	8.85	77.0	23.0
58.5	12.50	26.53	9.35

(Rimbach, B. 1902, 35. 1300.)

Sol. without decomp. in 50% LiCl+Aq, 33.3% CaCl₂+Aq and 50% MgCl₂+Aq. (Rimbach, B. 1905, 38. 1569.)

Ammonium chloromolybdenum chloride, 2NH₄Cl, Cl₄Mo₃Cl₂+2H₂O.

Decomp. by pure H₂O; can be crystallized from HCl+Aq. (Blomstrand.)

Ammonium chromium chloride, 2NH₄Cl, CrCl₃+H₂O.

Sol. in H₂O with decomp. (Neumann, A. 244. 229.)

+6H₂O=2NH₄Cl, [CrCl₃.4H₂O]Cl+2H₂O.

Hygroscopic. Decomp. by H₂O and by alcohol. (Weinland, B. 1907, 40. 3770.)

Ammonium cobaltous chloride, NH₄Cl, CoCl₂+6H₂O.

Deliquescent in moist air. Very easily sol. in H₂O. (Hautz, A. 66. 284.)

Ammonium cobaltous chloride ammonia, NH₄Cl, CoCl₂, NH₃. (F. Rose.)

Ammonium cuprous chloride, 4NH₄Cl.Cu₂Cl₂.

Decomp. in the air.

4NH₄Cl, 3Cu₂Cl₂. Decomp. by H₂O, not by alcohol. (Ritthausen, J. pr. 59. 369.)

Fairly stable in air. (Wells, Z. anorg. 1895, 10. 158.)

Ammonium cupric chloride, NH₄Cl, CuCl₂.

Solubility of NH₄Cl, CuCl₂ in absolute alcohol at 25°.

% CuCl ₂	Solid phase	% CuCl ₂	Solid phase
4.65	NH ₄ Cl+NH ₄ Cl, CuCl ₂	12.90	NH ₄ Cl, CuCl ₂
4.74	NH ₄ Cl+NH ₄ Cl, CuCl ₂	34.92	NH ₄ Cl, CuCl ₂ +CuCl ₂ .C ₂ H ₅ OH
6.45	NH ₄ Cl, CuCl ₂	34.50	"

(Foote and Walden, J. Am. Ch. Soc. 1911, 33. 1032.)

+2H₂O. Sol. in 2 pta. H₂O. (Hautz, A. 66. 280.)

Does not exist, (Meerburg, C. C. 1904. II. 1362.)

2NH₄Cl, CuCl₂+2H₂O. Easily sol. in H₂O, also in alcohol, even when absolute. (Cap and Henry, J. pr. 13. 184.)

Solubility of 2NH₄Cl, CuCl₂ in H₂O at t°.

g. 2NH ₄ Cl, CuCl ₂ in 100 g. of the solution	t°	Solid phase
3.87	-1.5°	ice
5.88	-2.48	"
8.78	-3.95	"
9.97	-4.60	"
13.12	-6.40	"
15.84	-8.04	"
17.64	-9.24	"
20.12	-10.80	"
20.3	-11.0	ice+2NH ₄ Cl, CuCl ₂ .2H ₂ O
20.46	-10	2NH ₄ Cl, CuCl ₂ .2H ₂ O
21.16	-5	"
22.02	0	"
24.26	+12	"
25.95	20	"
27.70	30	"
30.47	40	"
33.24	50	"
36.13	60	"
39.25	70	"
43.36	80	"

(Meerburg, Z. anorg. 1905, 45. 8.)

Somewhat sol. in liquid NH_3 . (Franklin and Kraus, Am. Ch. J. 1898, 20. 827.)

Is the only hydrate of $2\text{NH}_4\text{Cl}$, CuCl_2 , existing between -11° and $+80^\circ$. (Meerburg, C. C. 1904. II, 1362.)

$+3\text{H}_2\text{O}$. (Bourgeois, Bull. Soc. 1898, (3) 19. 786.)

Ammonium cupric chloride ammonia,
 $2\text{NH}_4\text{Cl}$, CuCl_2 , 2NH_3 .

Decomp. by H_2O , less easily by alcohol. Decomp. by acids. (Ritthausen.)

Ammonium indium chloride, $2\text{NH}_4\text{Cl}$, InCl_3 , $+ \text{H}_2\text{O}$.

Easily sol. in H_2O . (Meyer.)

Ammonium iodine chloride, NH_4Cl , ICl_3 .

More sol. in H_2O than KCl , ICl_3 . (Filhol, J. Pharm. 25. 441; Berz. J. B. 20. (2) 110.)

Ammonium iridium trichloride.

See Chloriridite, ammonium.

Ammonium iridium tetrachloride.

See Chloriridate, ammonium.

Ammonium iron (ferrous) chloride, NH_4Cl , FeCl_2 .

Easily sol. in H_2O ; insol. in alcohol. (Winkler.)

Ammonium iron (ferric) chloride, $2\text{NH}_4\text{Cl}$, $\text{FeCl}_3 + \text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O without decomp. (Fritzsche); sol. in 3 pts. H_2O at 18.75° . (Abl.)

Sol. in H_2O . (Walden, Z. anorg. 1894, 1. 32.)

Ammonium iron (ferric) potassium chloride,
 NH_4Cl , FeCl_3 , $\text{KCl} + 1\frac{1}{2}\text{H}_2\text{O}$.

Min. *Kremersite*. Deliquescent.

Ammonium lead chloride, NH_4Cl , $2\text{PbCl}_2 + 3\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (?). (André, C. R. 96. 1502.)

$6\text{NH}_4\text{Cl}$, $\text{PbCl}_2 + \text{H}_2\text{O}$.

$9\text{NH}_4\text{Cl}$, $\text{PbCl}_2 + 1\frac{1}{2}\text{H}_2\text{O}$.

$9\text{NH}_4\text{Cl}$, $2\text{PbCl}_2 + 2\frac{1}{2}\text{H}_2\text{O}$.

$10\text{NH}_4\text{Cl}$, $\text{PbCl}_2 + \text{H}_2\text{O}$.

$11\text{NH}_4\text{Cl}$, $2\text{PbCl}_2 + 3\frac{1}{2}\text{H}_2\text{O}$.

$18\text{NH}_4\text{Cl}$, $\text{PbCl}_2 + 4\text{H}_2\text{O}$.

All these salts are decomp. by H_2O . (André A. ch. (6) 3. 104.)

Of the salts prepared by André, only one NH_4Cl , 2PbCl_2 exists. (Wells, Sill. Am. J. 146. 25.)

Solubility determinations show that NH_4Cl , 2PbCl_2 is the only double salt formed at 25° . (Foote, Am. Ch. J. 1907, 37. 121.)

NH_4Cl , $\text{PbCl}_2 + \frac{1}{2}\text{H}_2\text{O}$. (Wells, l.c.)

Ammonium lead tetrachloride.

See Chloroplumbate, ammonium.

Ammonium magnesium chloride, $\text{NH}_4\text{MgCl}_2 + 6\text{H}_2\text{O} = \text{NH}_4\text{Cl}$, $\text{MgCl}_2 + 6\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O .

Sol. in 6 pts. cold H_2O . (Fourcroy.)

Solubility in $\text{NH}_4\text{Cl} + \text{Aq}$ at t° .

t°	Per 1000 Mol. H_2O	
	Mol. NH_4Cl	Mol. MgCl_2
3.5°	27.5	55.7
25 0	42.1	56.4
50.0	62.9	59.1

(Biltz, Z. anorg. 1911, 71. 170.)

$4\text{NH}_4\text{Cl}$, $5\text{MgCl}_2 + 33\text{H}_2\text{O}$. Sol. in H_2O . (Berthelot and André, A. ch. (6) 11. 294.)

Ammonium manganous chloride, NH_4Cl , $\text{MnCl}_2 + \frac{1}{2}\text{H}_2\text{O}$.

Sol. in $1\frac{1}{2}$ pts. H_2O at ordinary temp. (Hautz, A. 66. 280); does not exist. (Saunders, Am. Ch. J. 14. 134.)

$2\text{NH}_4\text{Cl}$, $\text{MnCl}_2 + \text{H}_2\text{O}$. Sol. in H_2O (Rammelsberg); does not exist. (Saunders.)

$+2\text{H}_2\text{O}$. Easily sol. in H_2O , but with decomp. into NH_4Cl and MnCl_2 . (Saunders.)

Ammonium manganic chloride, $2\text{NH}_4\text{Cl}$, MnCl_3 .

Sol. in H_2O ; less sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. Unstable. (Neuman, M. 1894, 15. 490.)

$+ \text{H}_2\text{O}$. Decomp. by H_2O . Sol. in HCl apparently without decomp. (Rice, Chem. Soc. 1898, 73. 260.)

Ammonium mercuric chloride, $2\text{NH}_4\text{Cl}$, $\text{HgCl}_2 + \text{H}_2\text{O}$ (sal alembroth).

Sol. in 0.66 pt. H_2O at 10° , and in nearly every proportion of hot H_2O .

NH_4Cl , HgCl_2 . Easily sol. in H_2O .

$+ \frac{1}{2}\text{H}_2\text{O}$. Easily sol. in H_2O . (Kane.)

$2\text{NH}_4\text{Cl}$, $3\text{HgCl}_2 + 4\text{H}_2\text{O}$. Easily sol. in H_2O . (Holmes, C. N. 5. 351.)

NH_4Cl , 2HgCl_2 . Very sol. in H_2O . (Ray, Chem. Soc. 1902, 81. 648.)

NH_4Cl , 5HgCl_2 . (Strömholm, J. pr. 1902, (2) 66. 441.)

Ammonium mercuric sodium chloride, NH_4Cl , HgCl_2 , 4NaCl (?).

Sol. in H_2O . (Kossmann, A. ch. (3) 27. 243.)

Ammonium molybdenum chloride, $2\text{NH}_4\text{Cl}$, $\text{MoCl}_5 + \text{H}_2\text{O}$.

Very sol. in H_2O . Nearly insol. in alcohol and ether. (Chilesotti, C. C. 1903. II, 652.)

See also Ammonium chloromolybdenum chloride.

Ammonium molybdenum chloride iodide.

See Ammonium chloromolybdenum iodide.

Ammonium molybdenyl chloride, $2\text{NH}_4\text{Cl}$, $\text{MoO}_2\text{Cl}_2 + 2\text{H}_2\text{O}$.

(Weinland, Z. anorg. 1905, **44**. 98.)

$2\text{NH}_4\text{Cl}$, MoOCl_3 . Sol. in H_2O ; insol. in H_2O sat. with HCl . (Klason, B. 1901, **34**. 149.)

Ammonium nickel chloride, NH_4Cl , $\text{NiCl}_2 + 6\text{H}_2\text{O}$.

Deliquescent in moist air. Easily sol. in H_2O . (Hautz.)

$4\text{NH}_4\text{Cl}$, $\text{NiCl}_2 + 7\text{H}_2\text{O}$ (?).

Ammonium osmium tetrachloride.

See Chlorosmate, ammonium.

Ammonium osmium sesquichloride.

See Chlorosmite, ammonium.

Ammonium osmyl chloride, $(\text{NH}_4)_2\text{OsO}_2\text{Cl}_4$.

Sol. in H_2O . Decomp. by HCl . (Wintrebert, A. ch. 1903, (7) **28**. 92.)

Ammonium osmyl oxychloride, $(\text{NH}_4)_2\text{OsO}_2\text{Cl}_2$.

Very sl. sol. in H_2O . Sol. in $\text{KOH} + \text{Ag}$ with decomp. (Wintrebert, A. ch. 1903, (7) **28**. 116.)

Ammonium palladium chlorides.

See Chloropalladate, ammonium and chloropalladite, ammonium.

Ammonium rhodium dichloride, $4\text{NH}_4\text{Cl}$, $\text{RhCl}_2 + 3\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O , but decomp. slowly. (Willm. B. 16. 3033.)

Does not exist. (Leidié, A. ch. (6) **17**. 277.)

Ammonium rhodium trichloride.

See Chlororhodite, ammonium.

Ammonium rhodium chloride ammonium nitrate, Rh_2Cl_6 , $6\text{NH}_4\text{Cl}$, $2\text{NH}_4\text{NO}_3$.

See Chlororhodite nitrate, ammonium.

Ammonium ruthenium trichloride.

See Chlororuthenite, ammonium.

Ammonium ruthenium tetrachloride.

See Chlororuthenate, ammonium.

Ammonium tellurium chloride.

See Chlorotellurate, ammonium.

Ammonium thallic chloride, $3\text{NH}_4\text{Cl}$, TiCl_3 .

Easily sol. in H_2O . (Willm.)

$+2\text{H}_2\text{O}$. Easily sol. in H_2O and alcohol. (Nicklès, J. Pharm. (4) **1**. 28.)

Ammonium thorium chloride, $8\text{NH}_4\text{Cl}$, $\text{ThCl}_4 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Chydenius.)

Ammonium tin (stannous) chloride (ammonium chlorostannite), NH_4Cl , $\text{SnCl}_2 + \text{H}_2\text{O}$.

Decomp. by H_2O . Resembles K salt. (Richardson, Am. Ch. J. **14**. 93.)

$2\text{NH}_4\text{Cl}$, $\text{SnCl}_2 + \text{H}_2\text{O}$. Sol. in H_2O , but decomp. by boiling. (Rammelsberg.)

Contains $2\text{H}_2\text{O}$. (Richardson.)

$4\text{NH}_4\text{Cl}$, $\text{SnCl}_2 + 3\text{H}_2\text{O}$. Decomp. by H_2O . (Poggiale, C. R. **20**. 1182.)

Does not exist. (Richardson.)

Ammonium tin (stannic) chloride.

See Chlorostannate, ammonium.

Ammonium titanium chloride, $2\text{NH}_4\text{Cl}$, $\text{TiCl}_4 + 2\text{H}_2\text{O}$.

Ppt.; decomp. in moist air; sol. in fuming HCl ; insol. in ether. (Rosenheim, Z. anorg. 1901, **26**. 242.)

Ammonium titanium chloride, $3\text{NH}_4\text{Cl}$, TiCl_4 . Sol. in H_2O .

$6\text{NH}_4\text{Cl}$, TiCl_4 . Sol. in H_2O . (Rose.)

Ammonium tungsten chloride, $(\text{NH}_4)_2\text{W}_2\text{Cl}_9 = 3\text{NH}_4\text{Cl}$, 2WCl_6 .

Easily sol. in H_2O . Nearly insol. in most organic solvents. (Olsson, B. 1913, **46**. 577.)

Ammonium uranyl chloride.

Very deliquescent, and sol. in H_2O . (Peligot.)

$2\text{NH}_4\text{Cl}$, $(\text{UO}_2)\text{Cl}_2 + 2\text{H}_2\text{O}$. Solution at 15° contains in 100 g. 3.51 g. NH_4 , 40.67 g. UO_2 and 19.15 g. Cl , hence there is considerable decomp. (Rimbach, B. 1904, **37**. 466.)

Ammonium vanadium chloride, $2\text{NH}_4\text{Cl}$, $\text{VCl}_3 + \text{H}_2\text{O}$.

Difficulty sol. in H_2O and alcohol. (Stähler, B. 1904, **37**. 4412.)

Ammonium zinc chloride, NH_4Cl , $\text{ZnCl}_2 + 2\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O . (Hautz, A. **66**. 287.)

$2\text{NH}_4\text{Cl}$, ZnCl_2 . Sol. in H_2O . (Rammelsberg, Pogg. **94**. 507.)

$+ \text{H}_2\text{O}$. Deliquescent in moist air. Sol. in $\frac{2}{3}$ pt. cold H_2O with absorption of heat. Sol. in 0.28 pt. hot H_2O (Golfier-Bassayre, A. ch. **70**. 344); sol. in $\frac{1}{2}$ pt. cold H_2O . (Hautz, A. **66**. 287.)

$3\text{NH}_4\text{Cl}$, ZnCl_2 . Sol. in H_2O . (Marignac.) $+ \text{H}_2\text{O}$. (Berthelot, A. ch. (6) **11**. 294.)

$4\text{NH}_4\text{Cl}$, ZnCl_2 . (Dehéraïn.)

$6\text{NH}_4\text{Cl}$, $\text{ZnCl}_2 + \frac{2}{3}\text{H}_2\text{O}$. (Berthelot, l.c.)

Ammonium chloride zinc oxychloride, 2ZnCl_2 , $8\text{NH}_4\text{Cl}$, ZnO .

Sol. in a little H_2O , but decomp. by excess. (André.)

3ZnCl_2 , $10\text{NH}_4\text{Cl}$, ZnO . As above. (André, A. ch. (6) **3**. 88.)

Ammonium chloride antimony fluoride, NH_4Cl , SbF_3 .

Easily sol. in H_2O . (de Haen, B. **21**. 901 R.)

Ammonium chloride arsenic trioxide.

See Arsenite chloride, ammonium.

Ammonium chloride bismuth bromide,
 $\text{NH}_4\text{Cl}, \text{BiBr}_3 + \text{H}_2\text{O}$.

Deliquescent; decomp. by H_2O . (Muir, Soc. **31**. 148.)

$\text{NH}_4\text{Cl}, \text{BiBr}_3 + 3\text{H}_2\text{O}$. Decomp. by H_2O .

$\text{NH}_4\text{Cl}, 2\text{BiBr}_3 + \text{H}_2\text{O}$. Decomp. by H_2O .

Ammonium chloride chromic oxychloride,
 $\text{NH}_4\text{Cl}, \text{CrOCl}_2$.

Stable in the air. Sol. in conc. HCl but decomp. (Weinland, B. 1906, **39**.)

Ammonium chloride cuprocupric thiosulphate,
 $\text{NH}_4\text{Cl}, \text{Cu}_2\text{O}, \text{CuO}, 3\text{S}_2\text{O}_3$.

Thiosulphate ammonium chloride, cuprocupric.

Ammonium chloride lead iodide, $3\text{NH}_4\text{Cl}, \text{PbI}_2$.

Decomp. with H_2O . (Behrens, Pogg. **62**.)

$\text{NH}_4\text{Cl}, \text{PbI}_2 + 2\text{H}_2\text{O}$. Decomp. with H_2O . (Gale, C. R. **20**. 1180.)

Ammonium chloride mercuric bromide,
 $\text{NH}_4\text{Cl}, \text{HgBr}_2$.

Rem-Bey, Dissert. 1885.)

Ammonium chloride platinum sulphite.
 Chloroplatosulphite, ammonium.

Ammonium chloride tin (stannous) bromide,
 $\text{NH}_4\text{Cl}, \text{SnBr}_2 + \text{H}_2\text{O}$.

Stable in H_2O . (Raymann and Preis, A. **223**.)

Ammonium dichloriodide, $\text{NH}_4\text{Cl}_2\text{I}$.

Stably decomp. when exposed to dry air at room temp. Very sol. in H_2O . (Chattaway, Soc. 1915, **107**. 107.)

Ammonium tetrachloriodide, $\text{NH}_4\text{Cl}_4\text{I}$.

Stable in the air. (Chattaway, Chem. Soc. 1915, **107**. 107.)

Ammonium lead chloriodide, $\text{NH}_4\text{PbClI}_2 + \text{H}_2\text{O}$ and $(\text{NH}_4)_2\text{PbCl}_2\text{I}_2 + 2\text{H}_2\text{O}$.

Stable in $\text{KOH} + \text{Aq}$ and in strong acids; decomp. by H_2O . (Fonzes-Diacon, Bull. Soc. **3**) **17**. 348.)

Ammonium fluoride, NH_4F .

Readily sol. in H_2O ; sl. sol. in alcohol. (Marignac, Ann. Min. (5) **15**. 221.)

Sl. in liquid NH_3 . (Ruff and Geisel, B. **36**. 820.)

Most insol. in liquid NH_3 at 50° . (Moissan, R. 1901, **133**. 713.)

Sl. in methyl alcohol. (Carrara, Gazz. **1896**, **26**. 119.)

Ammonium hydrogen fluoride, $\text{NH}_4\text{F}, \text{HF}$.
 Deliquescent in moist air. Sol. in H_2O .

Ammonium antimony fluoride, $2\text{NH}_4\text{F}, \text{SbF}_3$.

Deliquescent; sol. in 0.9 pt. cold H_2O . Insol. in alcohol or ether. (Flückinger, A. **84**. 248.)

$\text{NH}_4\text{F}, 4\text{SbF}_3$. 3 pts. sol. in 2 pts. H_2O . (Raad and Hauser, B. 1890, **23**. R. 125.)

$\text{NH}_4\text{F}, \text{SbF}_3$. Easily sol. in H_2O . (Marignac, A. **145**. 239.)

Ammonium bismuth fluoride, $2\text{NH}_4\text{F}, \text{BiF}_3$.

Insol. in H_2O . Rather difficultly sol. in acids. (Helmholt, Z. anorg. **3**. 115.)

Ammonium cadmium fluoride, $\text{NH}_4\text{F}, \text{CdF}_2$.

Insol. in H_2O . Sol. in acids on boiling. (Helmholt, Z. anorg. **3**. 115.)

Ammonium chromium fluoride, $3\text{NH}_4\text{F}, \text{CrF}_3$.

Easily sol. in H_2O . Sl. sol. in $\text{NH}_4\text{F} + \text{Aq}$. (Petersen, J. pr. (2) **40**. 52.)

$2\text{NH}_4\text{F}, \text{CrF}_3 + \text{H}_2\text{O}$. (Wagner, B. **19**. 896.)

Ammonium cobaltous fluoride, $2\text{NH}_4\text{F}, \text{CoF}_2 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Wagner, B. **19**. 896.)

Easily sol. in H_2O . (Helmholt, Z. anorg. **3**. 132.)

Ammonium columbyl fluoride.

See Fluoxycolumbate, ammonium.

Ammonium columbium fluoride oxyfluoride,
 $3\text{NH}_4\text{F}, \text{CbF}_3, \text{CbOF}_3$.

See Fluoxycolumbate columbium fluoride, ammonium.

Ammonium copper fluoride, $2\text{NH}_4\text{F}, \text{CuF}_2 + 2\text{H}_2\text{O}$.

Insol. in H_2O . (Helmholt, Z. anorg. **3**. 115.)

Nearly insol. in H_2O but decomp. thereby. (Haas, Ch. Z. 1908, **32**. 8.)

Ammonium glucinum fluoride, $2\text{NH}_4\text{F}, \text{GlF}_2$.

Sol. in H_2O . (Marignac, A. ch. (4) **30**. 51.)

Very sol. in H_2O . (Helmholt, Z. anorg. **3**. 130.)

Ammonium iron (ferrous) fluoride, $2\text{NH}_4\text{F}, \text{FeF}_2$. (Wagner, B. **19**. 896.)

$\text{NH}_4\text{F}, \text{FeF}_2 + 2\text{H}_2\text{O}$. (W.)

Ammonium iron (ferric) fluoride, $2\text{NH}_4\text{F}, \text{FeF}_3$.

More sol. in H_2O than the corresponding K compound. Decomp. by boiling. (Nicklès, J. Pharm. (4) **7**. 15.)

$3\text{NH}_4\text{F}, \text{FeF}_3$. Sl. sol. in H_2O . (Marignac, A. ch. (3) **60**. 306.)

Easily sol. in acids. (Helmholt, Z. anorg. **3**. 124.)

Ammonium manganic fluoride, $2\text{NH}_4\text{F}, \text{MnF}_4$.

More sol. than the K salt. (Nicklès, C. R. **65**. 107.)

True composition is $4\text{NH}_4\text{F}$, Mn_2F_8 . (Christensen, J. pr. (2) 34. 41.)

See also Fluomanganate, ammonium.

Ammonium manganyl fluoride.

See Fluoxymanganate, ammonium.

Ammonium molybdenum fluoride.

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. (Berzelius.)

See also Fluomolybdate, ammonium.

Ammonium molybdenyl fluoride.

See Fluoxymolybdate, ammonium.

Ammonium nickel fluoride, $2\text{NH}_4\text{F}$, $\text{NiF}_2 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Wagner, B. 19. 896.)

Easily sol. in H_2O . (Helmholtz, Z. anorg. 3. 143.)

Ammonium scandium fluoride, $(\text{NH}_4)_2\text{ScF}_6$.

Easily sol. in H_2O . Aqueous solution is not decomp. by boiling. Decomp. by acids. (R. I. Meyer, Z. anorg. 1914, 86. 275.)

Ammonium silicon fluoride.

See Fluosilicate, ammonium.

Ammonium silver fluoride, $2\text{NH}_4\text{F}$, $\text{AgF} + \text{H}_2\text{O}$.

Not hygroscopic. Sol. in H_2O ; sol. in conc. $\text{NH}_4\text{F} + \text{Aq}$. Sol. in alcohol. (Grützner, Arch. Pharm. 1900, 238. 3.)

$15\text{NH}_4\text{F}$, $\text{AgF} + 4\text{H}_2\text{O}$. More deliquescent than NH_4F . (Böhm, Dissert. 1906.)

Ammonium tantalum fluoride.

See Fluotantalate, ammonium.

Ammonium tantaly fluoride.

See Fluoxytantalate, ammonium.

Ammonium tellurium fluoride, NH_4F , TeF_4 .

Decomp. by H_2O . (Högbom, Bull. Soc. (2) 35. 60.)

Ammonium tin (stannous) fluoride, $2\text{NH}_4\text{F}$, $\text{SnF}_2 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Wagner, B. 19. 896.)

Ammonium tin (stannic) fluoride, $2\text{NH}_4\text{F}$, SnF_4 .

See Fluostannate, ammonium.

Ammonium titanium sesquifluoride.

See Fluotitanate, ammonium.

Ammonium titanyl fluoride.

See Fluoxypertitanate, ammonium.

Ammonium tungstyl fluoride.

See Fluoxytungstate, ammonium.

Ammonium uranyl fluoride.

See Fluoxyuranate, ammonium.

Ammonium vanadium sesquifluoride.

See Fluovanadate, ammonium.

Ammonium vanadyl fluoride.

See Fluoxyvanadate, ammonium.

Ammonium zinc fluoride, $2\text{NH}_4\text{F}$, ZnF_2 .

Sol. in H_2O . (R. Wagner.)

+ $2\text{H}_2\text{O}$. Very sl. sol. in H_2O . Easily sol. in dil. acids. (Helmholtz.)

Ammonium zirconium fluoride.

See Fluozirconate, ammonium.

Ammonium fluoride manganic oxyfluoride, $2\text{NH}_4\text{F}$, MnOF_2 .

Precipitate. (Nicklès.)

See also Fluoxymanganate, ammonium.

Ammonium fluoride molybdenum trioxide, $2\text{NH}_4\text{F}$, MoO_3 .

Decomp. by H_2O . (Mauro, Gazz. ch. it. 18. 120.)

Ammonium fluoride tungsten oxyfluoride.

See Fluoxytungstate, ammonium.

Ammonium fluoride tungsten oxyfluoride ammonium tungstate, $4\text{NH}_4\text{F}$, WO_3F_2 , $(\text{NH}_4)_2\text{WO}_4$.

See Fluoxytungstate tungstate, ammonium.

Ammonium fluoride vanadium oxyfluoride.

See Fluoxyvanadate, and fluoxyhypovanadate, ammonium.

Ammonium hydroselenide, NH_4HSe .

Sol. in H_2O with decomp. (Bineau, A. ch. (2) 67. 229.)

Ammonium hydrosulphide, NH_4SH .

Sol. in H_2O and alcohol. Solutions decomp. on air.

Ammonium hydroxide, NH_4OH .

See Ammonia,

Ammonium imidosulphamide,

$(\text{S}_2\text{O}_4\text{N}_2\text{H}_4)\text{NH}_4$.

(Hantzsch, B. 1905, 38. 1033.)

Ammonium iodide, NH_4I .

Very deliquescent. Sol. in 0.60 pt. H_2O . (Eder, Dingl. 221. 89.)

Sp. gr. of aqueous solution of NH_4I at 18° containing—

10	20	30	40	50% NH_4I
1.0652	1.1397	1.2260	1.3260	1.4415

(Kohlrausch, W. Ann. 1879. 1.)

$\text{NH}_4\text{I} + \text{Aq}$ containing 12.51% NH_4I has sp. gr. $20^\circ/20^\circ = 1.0846$.

$\text{NH}_4\text{I} + \text{Aq}$ containing 19.19% NH_4I has sp. gr. $20^\circ/20^\circ = 1.1359$.

(Le Blanc and Robland, Z. phys. Ch. 1896, 19. 279.)

Very easily sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.)

Very sol. in liquid NH_3 at -50° . (Moissan, C. R. 1901, 133. 713.)

Sol. in SOCl_2 . (Walden, Z. anorg. 1900, 25. 216.)

Sol. in liquid SO_2 . (Walden, Z. anorg. 1902, 30. 160.)

Sol. in 4.0 pts. abs. alcohol. (Eder, l.c.)

" 210 " ether. (Eder, l.c.)

" 20 " alcohol-ether (1 : 1). (Eder, l.c.)

Sol. in acetone. (Eidmann, C.C. 1899, II. 1014.); (Naumann, B. 1904, 37. 4328.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Ammonium diiodide, NH_4I_2 .

Sol. in alcohol, ether, CS_2 , and $\text{KI} + \text{Aq}$; less sol. in chloroform. (Guthrie, Chem. Soc. (2) 1. 239.)

Ammonium triiodide, NH_4I_3 .

Sl. deliquescent. Sol. in little H_2O , but decomp. by much H_2O . (Johnson, Chem. Soc. 33. 397.)

Ammonium antimony iodide, NH_4I , $\text{SbI}_3 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Nicklès, C. R. 51. 1097.)

$3\text{NH}_4\text{I}$, $4\text{SbI}_3 + 9\text{H}_2\text{O}$. Decomp. by H_2O , with separation of SbOI . Sol. in $\text{HC}_2\text{H}_3\text{O}_2$, HCl , and $\text{H}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq}$. Decomp. by CS_2 . (Schäffer, Pogg. 109. 611.)

$3\text{NH}_4\text{I}$, $\text{SbI}_3 + 3\text{H}_2\text{O}$. As above.

$4\text{NH}_4\text{I}$, $\text{SbI}_3 + 3\text{H}_2\text{O}$. As above.

Ammonium bismuth iodide, NH_4I , $\text{BiI}_3 + \text{H}_2\text{O}$.

Deliquescent; decomp. by H_2O . (Nicklès, C. R. 51. 1097.)

$4\text{NH}_4\text{I}$, $\text{BiI}_3 + 3\text{H}_2\text{O}$. As above. (Linau, Pogg. 111. 240.)

$2\text{NH}_4\text{I}$, $\text{BiI}_3 + 2\frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O , or MCl , MBr , or $\text{MI} + \text{Aq}$. (Nicklès, J. pr. (2) 39. 116.)

Ammonium cadmium iodide, $2\text{NH}_4\text{I}$, $\text{CdI}_2 + 2\text{H}_2\text{O}$.

Deliquescent. (Croft.)

Sol. at 15° in 0.58 pt. H_2O , 0.70 pt. abs. alcohol., 8.9 pts. ether (sp. gr. 0.729), and 1.8 pts. alcohol-ether (1 : 1). (Eder, Dingl. 221. 89.)

100 pts. of the solution in H_2O contain 85.97 pts. of the salt at 14.5° . (Rimbach, B. 1905, 33. 1563.)

NH_4I , $\text{CdI}_2 + \frac{1}{2}\text{H}_2\text{O}$. Sol. at 15° in 0.90 pt. H_2O , 0.88 pt. abs. alcohol, and 2.4 pts. ether (sp. gr. 0.729). (Eder, l.c.)

$+ \text{H}_2\text{O}$. (Grossmann, Z. anorg. 1902, 33. 154.)

Ammonium chloromolybdenum iodide, $2\text{NH}_4\text{I}$, $\text{Cl}_4\text{Mo}_3\text{I}_2 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . Cryst. from $\text{HI} + \text{Aq}$. (Blomstrand.)

Ammonium cuprous iodide, $2\text{NH}_4\text{I}$, $\text{Cu}_2\text{I}_2 + \text{H}_2\text{O}$.

Decomp. on the air, or by H_2O , or alcohol. (Saglier, C. R. 104. 1440.)

$+ \frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O with separation of Cu_2I_2 . (Gossner, Zeit. Kryst. 1903, 38. 501.)

Ammonium cupric iodide ammonia, $2\text{NH}_4\text{I}$, CuI_2 , $2\text{NH}_3 + 2\text{H}_2\text{O}$.

Insol. in H_2O or alcohol; sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

$+ 6\text{H}_2\text{O}$. Unstable. (Saglier, C. R. 104. 1440.)

NH_4I , 2CuI_2 , 3NH_3 . (Fleurent, C. R. 1891, 113. 1047.)

Ammonium iridium diiodide, $2\text{NH}_4\text{I}$, IrI_2 .

Insol. in cold or hot H_2O , and in alcohol. Sol. in warm dil. acids. (Oppler.)

Ammonium iridium sesquiiodide.

See Iodiridite, ammonium.

Ammonium iridium tetraiodide.

See Iodiridate, ammonium.

Ammonium lead iodide, NH_4I , $\text{PbI}_2 + 2\text{H}_2\text{O}$.

Decomp. by much H_2O . (Wells, Sill. Am. J. 146. 25.)

$4\text{NH}_4\text{I}$, $3\text{PbI}_2 + 6\text{H}_2\text{O}$. Sl. sol. in H_2O . (Mosnier, C. R. 1895, 120. 444.)

Sol. in H_2O with decomp. Sol. in strong $\text{KOH} + \text{Aq}$ and in strong acids. (Fonzes-Diacon, Bull. Soc. 1897, (3) 17. 347.)

Ammonium magnesium iodide, NH_4I , $\text{MgI}_2 + 6\text{H}_2\text{O}$.

Very deliquescent. (Lerch, J. pr. (2) 28. 338.)

Ammonium mercuric iodide, NH_4I , $\text{HgI}_2 + \text{H}_2\text{O}$.

Decomp. into its constituents by H_2O . (Boullay, A. ch. (2) 34. 345.)

Sol. without decomp. in alcohol and ether. NH_4I , 2HgI_2 . Decomp. by H_2O . Sol. in $\text{KI} + \text{Aq}$. Very sol. in alcohol, ether and nitrobenzol. (Löw, Zeit. Kryst, 51. 138.)

Ammonium silver iodide, $2\text{NH}_4\text{I}$, AgI .

Deliquescent. Decomp. by H_2O . (Poggiale.)

Ammonium thallic iodide, NH_4I , TlI_3 .

Sol. in H_2O . (Nicklès, J. Pharm. (4) 1. 32.)

Ammonium tin (stannous) iodide, NH_4I , SnI_2 .

Decomp. by small amt. H_2O but completely sol. in a large amt. (Boullay, A. ch. (2) 34. 376.)

$+ 1\frac{1}{2}\text{H}_2\text{O}$. (Personne.)

Ammonium zinc iodide, $2\text{NH}_4\text{I}$, ZnI_2 .

Extremely deliquescent, and sol. in H_2O . (Rammelsberg, Pogg. 43. 665.)

NH_4I , $\text{ZnI}_2 + 4\frac{1}{2}\text{H}_2\text{O}$. Hygroscopic. (Ephraim, Z. anorg. 1910, **67**. 384.)

Ammonium iodide arsenic trioxide.

See **Arsenite iodide, ammonium.**

Ammonium cobalt nitride.

See **Ammonium cobalt azoimide.**

Ammonium ruthenium dihydronitrosobromide, $\text{NO}.\text{Ru}_2\text{H}_2(\text{NH}_3)_6\text{Br}_2.2\text{HBr}$.

Ppt. (Brizard, A. ch. 1900, (7) **21**. 363.)

Ammonium ruthenium nitrosochloride, $3\text{NH}_4\text{Cl}.2\text{HCl}.\text{NORu}_2\text{H}_2\text{Cl}_3$.

Ppt. (Brizard, A. ch. 1900, (7) **21**. 354.)

Ammonium ruthenium dihydronitrosochloride, $\text{NO}.\text{Ru}_2\text{H}_2(\text{NH}_3)_6\text{Cl}_2.2\text{HCl}$.

Ppt. (Brizard, A. ch. 1900, (7) **21**. 358.)

Ammonium peroxide, $(\text{NH}_4)_2\text{O}_2$.

M.-pt. -2° . Sl. sol. in ether without decomp. (D'Ans, B. 1913, **46**. 3076.)

Sol. in alcohol; insol. in ether; decomp. slowly in aq. solution. (Melikoff, B. 1897, **30**. 3145.)

Ammonium hydrogen peroxide, $(\text{NH}_4)_2\text{O}_2, \text{H}_2\text{O}_2$.

Decomp. at ordinary temp. (Melikoff, B. 1898, **31**. 447.)

+ H_2O . Unstable; deliquesces at ordinary temp.; sol. in alcohol; insol. in light petroleum. (Melikoff, B. 1898, **31**. 152.)

Ammonium selenide, $(\text{NH}_4)_2\text{Se}$.

Sol. in H_2O with decomp. (Bineau, A. ch. (2) **67**. 229.)

Stable in the air. Sol. in H_2O ; aq. solution decomp. slowly. (Lenher and Smith, J. Am. Chem. Soc. 1898, **20**. 277.)

Ammonium hydrogen selenide, NH_4HSe .

Sol. in H_2O . (Fabre, C. R. **103**. 269.)

Ammonium monosulphide, $(\text{NH}_4)_2\text{S}$.

Decomp. on air. Sol. in H_2O , but solution decomposes rapidly.

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 826.)

Ammonium disulphide, $(\text{NH}_4)_2\text{S}_2$.

Sol. in H_2O with decomp.

Does not exist. (Bloxam, Chem. Soc. 1895, **67**. 293.)

Ammonium tetrasulphide, $(\text{NH}_4)_2\text{S}_4$.

Easily sol. in H_2O . Conc. solution is stable, dil. solution decomp. on air. Easily sol. in alcohol without decomp., but solution decomp. on the air more rapidly than the aqueous solution. (Fritzsche, J. pr. **32**. 313.)

+ $\frac{1}{2}\text{H}_2\text{O}$. When dissolved in H_2O , it is at once dissociated with deposition of S. (Bloxam, Chem. Soc. 1895, **67**. 303.)

Ammonium pentasulphide, $(\text{NH}_4)_2\text{S}_5$.

Decomp. on air. Sol. in H_2O with separation of S. Sol. in alcohol without decomp., but solution decomposes quickly on standing. (Fritzsche, J. pr. **32**. 313.)

Rapidly decomp. by H_2O with separation of S. (Bloxam, Chem. Soc. 1895, **67**. 298.)

+ H_2O . Decomp. by H_2O with separation of S. (Bloxam, Chem. Soc. 1895, **67**. 298.)

Ammonium heptasulphide, $(\text{NH}_4)_2\text{S}_7$.

More stable on air, and less easily decomposed by H_2O than $(\text{NH}_4)_2\text{S}_5$.

+ $1\frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O with separation of S. Slowly attacked by dil. $\text{HCl} + \text{Aq}$. (Bloxam, Chem. Soc. 1895, **67**. 307.)

Tetrammonium heptasulphide, $(\text{NH}_4)_4\text{S}_7 + 4\text{H}_2\text{O}$.

Sol. in H_2O . Solution can be kept for a long time without depositing S. (Bloxam, Chem. Soc. 1895, **67**. 298.)

Diammonium enneasulphide, $(\text{NH}_4)_2\text{S}_9 + \frac{1}{2}\text{H}_2\text{O}$.

Decomposed by H_2O with separation of S. Not attacked by boiling dil. $\text{HCl} + \text{Aq}$ on account of formation of a hard crust of S on the crystals. (Bloxam, Chem. Soc. 1895, **67**. 306.)

Tetrammonium enneasulphide, $(\text{NH}_4)_4\text{S}_9$.

Solution in H_2O deposits crystals of $(\text{NH}_4)_2\text{S}_9$ on standing. (Bloxam, Chem. Soc. 1895, **67**. 302.)

+ $3\frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O with separation of S. (Bloxam, Chem. Soc. 1895, **67**. 299.)

Ammonium polysulphides.

Conc. $\text{NH}_3 + \text{Aq}$ dissolves H_2S to form $(\text{NH}_4)_2\text{S}.2\text{NH}_4\text{SH}$. On dilution more H_2S is absorbed to form $(\text{NH}_4)_2\text{S}.4\text{NH}_4\text{SH}$, then $(\text{NH}_4)_2\text{S}.8\text{NH}_4\text{SH}$, then $(\text{NH}_4)_2\text{S}.18\text{NH}_4\text{SH}$ and finally NH_4SH . (Bloxam, Chem. Soc. 1895, **67**. 284.)

Ammonium copper sulphide, $(\text{NH}_4)_2\text{S}.2\text{CuS}_2$ (?).

Sol. in warm H_2O , but decomp. on standing. Warm $\text{KOH} + \text{Aq}$ acts similarly; sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$, $\text{Na}_2\text{CO}_3 + \text{Aq}$, or absolute alcohol. Insol. in ether. Decomp. by dil. acids. (Priwoznik, B. **6**. 1291.)

Correct formula is NH_4CuS_4 . Sl. sol. in H_2O . Decomp. by conc. and dil. acids. Easily sol. in NaOH . Sl. sol. in alcohol. (Biltz, B. 1907, **40**. 976.)

Ammonium gold polysulphide, $\text{AuS}_3.\text{NH}_4$.

Ppt. (Hofmann, B. 1903, **36**. 3092; B. 1904, **37**. 245.)

Ammonium iridium pentadecasulphide, $\text{IrS}_{15}(\text{NH}_4)_3$.

Ppt. (Hofmann, B. 1904, **37**. 247.)

Ammonium palladium undecasulphide,
 $\text{PdS}_{11}(\text{NH}_4)_2 + \frac{1}{2}\text{H}_2\text{O}.$

Ppt. (Hofmann, B. 1904, 37. 248.)

Ammonium platinum pentadecasulphide,
 $\text{PtS}_{15}(\text{NH}_4)_2 + 2\text{H}_2\text{O}.$

Can be washed with CS_2 without decomp.
 Sol. in alcohol. Insol. in ether. (Hofmann, B. 1903, 36. 3091.)

Ammonium stannic sulphide.

See Sulphostannate, ammonium.

Ammonium telluride, $\text{NH}_4\text{HTe}.$

Easily sol. in H_2O . (Bineau, A. ch. (2) 67. 229.)

Ammonium sulphide ammonia, $(\text{NH}_4)_2\text{S}, 2\text{NH}_3.$

Very unstable. (Bloxam, Chem. Soc. 1895, 67. 294.)

Ammonium acisulphomelid, $(\text{NSO.ONH}_4)_2$ (?)

(Hantzsch and Stuer, B. 1905, 38. 1039.)

Ammonplatindiamine comps.

See Platintriamine comps.

Ammondisulphonic acid, $\text{NH}_2(\text{SO}_3\text{H})_2.$

Known only in its salts. (Claus, A. 158. 52 and 194.)

Contains 2 at. H less, and is identical with imidosulphonic acid $\text{NH}(\text{SO}_3\text{H})_2$, which see. (Raschig, A. 241. 161.)

Ammontrisulphonic acid, $\text{NH}_2(\text{SO}_3\text{H})_3.$

Known only in its salts. (Claus, A. 158. 52 and 194.)

Contains 2 at. H less, and is nitrilosulphonic acid $\text{N}(\text{SO}_3\text{H})_3$, which see. (Raschig, A. 241. 161.)

Ammontetrasulphonic acid, $\text{NH}(\text{SO}_3\text{H})_4.$

Known only in its salts. (Claus, A. 158. 52 and 194.)

Does not exist, but was impure nitrilosulphonic acid, which see. (Raschig, A. 241. 161.)

Anhydroarseniotungstic acid, $\text{H}_2\text{AsW}_3\text{O}_{22}.$

See under Arseniotungstic acid.

Anhydrooxycobaltamine chloride,



Easily sol. in H_2O , but decomposes after a few minutes; can be recrystallized from dil. $\text{HCl} + \text{Aq}$. Precipitated from sat. H_2O solution by conc. $\text{HCl} + \text{Aq}$, or alcohol. (Vortmann, M. Ch. 6. 404.)

$\text{Co}_2(\text{NH}_3)_{10} \left(\begin{array}{c} \text{Cl} \\ \text{OH} \end{array} \right) \text{Cl}_4.$ Sol. in H_2O . (Vortmann.)

Anhydrooxycobaltamine chloride mercuric chloride, $\text{Co}_2(\text{NH}_3)_{10}(\text{ClO}_2\text{H})\text{Cl}_4, 3\text{HgCl}_2.$

Can be recryst. from very dil. hot $\text{HCl} + \text{Aq}$.

— **chloroplatinate, $\text{Co}_2(\text{NH}_3)_{10}(\text{ClO}_2\text{H})\text{Cl}_4, 2\text{PtCl}_4.$**

Can be recrystallized from H_2O containing HCl .

— **chloronitrate, $\text{Co}_2(\text{NH}_3)_{10}\text{Cl}(\text{O.OH})(\text{NO}_2)_4 + \text{H}_2\text{O}.$**

Can be recrystallized from dil. $\text{HCl} + \text{Aq}$.

$\text{Co}_2(\text{NH}_3)_{10}\text{Cl}(\text{O.OH})\text{Cl}_2(\text{NO}_2)_2 + \text{H}_2\text{O}.$
 More easily sol. in H_2O than the preceding comp.

— **chlorosulphate, $\text{Co}_2(\text{NH}_3)_{10}\text{Cl}(\text{O.OH})(\text{SO}_4)_2.$**

— **dichromate, $[\text{Co}_2(\text{NH}_3)_{10}\text{O.OH}]_2(\text{Cr}_2\text{O}_7)_2 + 8\text{H}_2\text{O}.$**

Sl. sol. in H_2O .

— **nitrate, $\text{Co}_2(\text{NH}_3)_{10}(\text{NO}_2)(\text{O.OH})(\text{NO}_2)_4 + \text{H}_2\text{O}.$**

Sl. sol. in pure H_2O with immediate decomp.
 Can be recrystallized from H_2O containing HNO_3 .

— **sulphate, $[\text{Co}_2(\text{NH}_3)_{10}\text{O.OH}]_2(\text{SO}_4)_2, 2\text{H}_2\text{SO}_4 + 2\text{H}_2\text{O}.$**

Sl. sol. in cold H_2O . When crystallized from dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, is converted into—

$[\text{Co}_2(\text{NH}_3)_{10}\text{O.OH}]_2(\text{SO}_4)_2, \text{H}_2\text{SO}_4 + 3\text{H}_2\text{O},$
 which by further recrystallization from very dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ becomes—

$[\text{Co}_2(\text{NH}_3)_{10}\text{O.OH}]_2(\text{SO}_4)_2 + 8\text{H}_2\text{O}.$ Sl. sol. in cold H_2O . (Vortmann.)

Anhydrophospholuteotungstic acid, $\text{H}_2\text{PW}_3\text{O}_{22}.$

See under Phosphotungstic acid.

Antimonic Acid.

Metantimonic acid, $\text{HSbO}_3.$

Very sl. sol. in H_2O ; sol. in conc. $\text{HCl} + \text{Aq}$; sl. sol. in dil. $\text{HNO}_3 + \text{Aq}$; easily sol. in tartaric acid + Aq ; easily sol. in hot KOH , or $\text{NaOH} + \text{Aq}$; completely insol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Fremy, A. ch. (3) 23. 407.)

Sl. sol. in H_2O . Very sl. sol. in KOH and $\text{K}_2\text{CO}_3 + \text{Aq}$. Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in $\text{HNO}_3 + \text{H}_2\text{SO}_4$. Slowly sol. in cold, quickly in hot $\text{HCl} + \text{Aq}$. Sl. sol. in tartaric and oxalic acid and in $\text{KHC}_2\text{O}_4 + \text{Aq}$. (Senderens, Bull. Soc. 1899, (3) 21. 48.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Pyroantimonic acid, $\text{H}_4\text{Sb}_2\text{O}_7.$

More sol. in H_2O and acids than H_2SbO_4 . Sol. in cold NH_4OH , or $\text{KOH} + \text{Aq}$. (Fremy.)

Slowly sol. in cold H_2O .

5.88 g. Sb_2O_5 in 1 l. H_2O at 15°

8.55 " " " 1 l. " " 25°

21.30 " " " 1 l. " " 60°

(Delacroix, J. Pharm. 1897, 6. 337-41.)

Sl. sol. in H_2O . Very sl. sol. in KOH and $K_2CO_3 + Aq$. Insol. in $NH_4OH + Aq$, and in $HNO_3 + H_2SO_4$. Slowly sol. in cold, quickly in hot $HCl + Aq$. Sl. sol. in tartaric acid, oxalic acid and $KHC_2O_4 + Aq$. (Senderens, Bull. Soc. 1899, (3) 21. 48.)

Orthoantimonic acid, H_3SbO_4 .

Sl. sol. in H_2O . Insol. in $NH_4OH + Aq$. Easily sol. in $KOH + Aq$. (Freymy.)

Does not exist. (Raschig, B. 18. 2745.)

Has, however, been prepared by Daubrawa (A. 186. 110), Conrad (C. N. 40. 198), and Beilstein and Blaese (Bull. Ac. St. Petersb. 33. 97).

Very sol. in H_2O . (Delacroix, Bull. Soc. 1899 (3) 21. 1049.)

Very sl. sol. in H_2O , in KOH and $K_2CO_3 + Aq$. Slowly sol. in cold, quickly in hot $HCl + Aq$. Insol. in $NH_4OH + Aq$, and in $HNO_3 + H_2SO_4$. Sl. sol. in tartaric acid, oxalic acid and $KHC_2O_4 + Aq$. (Senderens, Bull. Soc. 1899, (3) 21. 52.)

+ $\frac{1}{2}H_2O$. (Beilstein and Blaese.)

According to Beilstein and Blaese only one antimonic acid, H_3SbO_4 , exists.

Tetrantimonic acid, $Sb_2O_5 + 4H_2O = H_3Sb_2O_9$.

Slowly sol. in cold H_2O .

Solution sat. at t° contains g. Sb_2O_5 per litre—

t°	15°	25°	60°	70°
g. Sb_2O_5	5.88	8.3–8.75	21.30	53.89

Decomp. in solution by heating to 100° or long standing in the cold to Sb_2O_5 , $3H_2O$. (Delacroix, Bull. Soc. 1899, (3) 21. 1049.)

Insol. in H_2O . Very sl. sol. in KOH and $K_2CO_3 + Aq$. Slowly sol. in cold, quickly in hot $HCl + Aq$. Insol. in $NH_4OH + Aq$. Insol. in $HNO_3 + H_2SO_4$. Sl. sol. in tartaric acid, oxalic acid and in $KHC_2O_4 + Aq$. (Senderens, Bull. Soc. 1899, (3) 21. 51.)

Hexantimonic acid, $Sb_2O_5 + 6H_2O = H_{11}Sb_2O_{11}$.

Sol. in H_2O to the extent of 22 g. Sb_2O_5 per l. but on standing becomes turbid and a white powder is pptd. until finally only 3 g. Sb_2O_5 are dissolved per l. (Senderens, Bull. Soc. 1899, (3) 21. 48–49.)

Antimonates.

a. Antimonates. From $HSbO_3$. Some of the K and NH_4 salts are sol. in H_2O , the others are slightly sol. or insol.

β . Pyroantimonates. From $H_4Sb_2O_7$. As a class, insol. in H_2O , but decomp. thereby except in presence of large excess of alkali. (Freymy, A. ch. (3) 12. 499.)

Probably do not exist. (Beilstein and Blaese.)

Aluminum antimonate, $Al_2O_3, 3Sb_2O_5$ (?).

Ppt. Somewhat sol. in excess of Al salts + Aq . Insol. in $K_4Sb_2O_7 + Aq$.

$Al(SbO_3)_3 + 15H_2O = AlH_6(SbO_4)_3 + 12H_2O$.

Ppt. (Beilstein and Blaese, Bull. Ac. St. Petersb. 33. 101.)

$Al(SbO_3)_3 + 7H_2O = AlH_6(SbO_4)_3 + 4H_2O$.

Ppt. (B. and B.)

$Al_2O_3, Sb_2O_5 + 9H_2O$. Ppt. (Ebel, B. 22. 3043.)

Ammonium antimonate, $NH_4SbO_3 + 2H_2O$.

Insol. in H_2O .

+ $2\frac{1}{2}H_2O$. Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 56.)

+ $6H_2O$. See $(NH_4)_2H_3Sb_2O_7 + 5H_2O$.

Ammonium pyroantimonate, $(NH_4)_2Sb_2O_7$.

Known only in solution.

$(NH_4)_2H_3Sb_2O_7 + 5H_2O$.

Sol. in H_2O , but decomp. by standing or boiling into insol. salt. Insol. in alcohol. (Freymy, J. pr. 45. 215). Composition is $NH_4SbO_3 + 6H_2O$, according to Raschig (B. 18. 2743).

Barium antimonate, $Ba(SbO_3)_2$.

Ppt. Scarcely sol. in H_2O . Slowly sol. in $BaCl_2 + Aq$.

+ $2H_2O$. Somewhat sol. in H_2O . Easily sol. in $HCl + Aq$. (Delacroix, Bull. Soc. 1899, (3) 21. 1051.)

+ 5, or $6H_2O$. Ppt.

$BaSb_4O_7 + 5H_2O$. Sol. in conc. HCl . (Delacroix, Bull. Soc. 1899, (3) 21. 1051.)

$BaO, 3Sb_2O_5 + 5H_2O$. Insol. in H_2O . Incompletely sol. in HCl . (Delacroix, *l.c.*)

$BaO, 4Sb_2O_5 + 15H_2O$. (Delacroix, *l.c.*)

$9BaO, 10Sb_2O_5 + 18H_2O$. Insol. in $HCl + Aq$. (Delacroix, *l.c.*)

Bismuth antimonate, $BiSbO_4 + H_2O$.

Ppt. Insol. in H_2O ; sol. in $HCl + Aq$. (Cavazzi, Gazz. ch. it. 15. 37.)

$3Bi_2O_3, Sb_2O_5 + H_2O$. Insol. in H_2O ; sol. in $HCl + Aq$. (Cavazzi.)

$2Bi_2O_3, Sb_2O_5$. As above. (Cavazzi.)

Cadmium antimonate, $Cd(SbO_3)_2 + 2H_2O$.

Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 56.)

+ $3\frac{1}{2}H_2O$. Very sol. in H_2O . Sol. in $HCl + Aq$. (Ebel, Dissert. 1890.)

+ $5H_2O$. Insol. in H_2O . (Senderens, *l.c.*)

+ $6H_2O$. Ppt. Insol. in H_2O . (Ebel, B. 22. 3043.)

Calcium antimonate, $Ca(SbO_3)_2$.

Ppt.

+ $5H_2O$. Ppt. (Heffter, Pogg. 86. 418.)

+ $6H_2O$. Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 56.)

$3CaO, 2Sb_2O_5 + 6H_2O$. Min. *Ullmanite*.

Chromic antimonate, $Cr(SbO_3)_3 + 14H_2O$.

Ppt. (Beilstein and Blaese.)

Cobaltous antimonate, $Co(SbO_3)_2 + 5H_2O$.

Insol. in H_2O . Loses $3H_2O$ in the presence of H_2SO_4 and passes into $Co(SbO_3)_3 + 2H_2O$.

also insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 55.)

+6 H_2O . Ppt. (Ebel, B. 22. 3043.)

+7 H_2O . Sl. sol. in H_2O . Sl. sol. in boiling solutions of cobalt salts.

+12 H_2O . Ppt. (Heffter, Pogg. 86. 448.)

Cobaltous hydrogen antimonate, $CoH_4(SbO_4)_2 + H_2O$.

(Gorgeul, Ann. Phys. Beibl. 1897, 21. 198.)

Cupric antimonate, $3CuO, 2Sb_2O_3$.

Ppt. (Beilstein and Blaese.)

$Cu(SbO_3)_2$. Insol. in H_2O , acids, or alkalies. (Berzelius.)

+2 H_2O . Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 55.)

+5 H_2O . Ppt. (Ebel, B. 22. 3043.)

Insol. in H_2O . (Senderens, l.c.)

$CuO, 2Sb_2O_3 + 9H_2O$. Insol. in H_2O . Sol. in $Sb_2O_3, 4H_2O + Aq$. (Delacroix, Bull. Soc. 1899, (3) 21. 1054.)

$2CuC_2, 3Sb_2O_3 + 10H_2O$. Insol. in H_2O . Sol. in NH_4OH and in triantimonic acid + Aq. (Delacroix, l. c.)

$CuO, 6Sb_2O_3 + 16H_2O$. (Delacroix, l.c.)

Cupric antimonate ammonia, $Cu(SbO_3)_2, 4NH_3 + 4H_2O$.

Insol. in H_2O and $NH_4OH + Aq$. (Schiff, A. 123. 39.)

$CuSb_2N_3H_{21}O_{12} = Cu(ONH_4)OH, 2(NH_4SbO_3 + 2H_2O)$. (Raschig, B. 18. 2743.)

$Cu(SbO_3)_2, 3NH_3 + 9H_2O$. (Delacroix, Bull. Soc. 1901, (3) 25. 289.)

Glucinum antimonate, $Gl(SbO_3)_2 + 6H_2O$.

Somewhat sol. in hot H_2O . Easily sol. in warm HCl . (Ebel, Dissert. 1890.)

Iron (ferrous) antimonate.

Sl. sol. in H_2O . (Berzelius.)

Iron (ferric) antimonate.

Insol. in H_2O . (B.)

$Fe_2O_3, Sb_2O_3 + 7H_2O$. Ppt. (Ebel, B. 22. 3043.)

$Fe_2O_3, 2Sb_2O_3 + 11H_2O$. Ppt. (Beilstein and Blaese.)

$Fe(SbO_3)_3 + 6\frac{1}{2}H_2O$. Ppt. (B. and B.)

Lead antimonate, basic, $Pb_3(SbO_3)_2(OH)_4 + 2H_2O = Pb_3(SbO_4)_2 + 4H_2O$.

Min. *Bleinerite, Bindheimite*.

$2Pb(SbO_3)_2, PbO + 11H_2O$. Ppt. (B. and B.)

Lead antimonate, $Pb(SbO_3)_2$.

Insol. in H_2O . Incompletely decomp. by acids. (Berzelius.)

Naples Yellow. Insol. in H_2O .

+2 H_2O . Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 57.)

+5 H_2O . Ppt. (Ebel, B. 22. 3043.)

+6 H_2O . Ppt. (Beilstein and Blaese.)

+9 H_2O . Insol. in H_2O . (Senderens, l.c.)

Lead antimonate chloride, $Pb(SbO_3)_2, PbCl_2$.

Min. *Nadorite*. Sol. in HCl, HNO_3 , and tartaric acid + Aq.

Lithium antimonate, $LiSbO_3$.

Sl. sol. in cold, sol. in hot H_2O , and crystallizes on cooling. Much more sol. than $NaSbO_3$.

+3 H_2O . Ppt. Sl. sol. in H_2O . (Beilstein and Blaese.)

Magnesium antimonate, $Mg(SbO_3)_2 + 12H_2O$.

Sol. in hot, less sol. in cold H_2O . (Heffter.)

Sol. in $MgSO_4 + Aq$; insol. in $KSbO_3 + Aq$. (Berzelius.)

Manganous antimonate, $Mn(SbO_3)_2$.

Difficultly sol. in H_2O .

When heated, is sol. only in strong acids.

+2 H_2O . Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 56.)

+5 H_2O . Ppt. (Ebel, B. 22. 3043.)

+6 H_2O . Insol. in H_2O . (Senderens, l.c.)

+7 H_2O . Ppt. (Beilstein and Blaese.)

Mercurous antimonate.

Insol. in H_2O . (Berzelius.)

Mercuric antimonate, $Hg(SbO_3)_2$.

Insol. in H_2O , alkalies, and most acids.

Sl. attacked by boiling H_2SO_4 , and $HCl + Aq$.

+2 H_2O . Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 55.)

+5 H_2O . Insol. in H_2O . (Senderens.)

+6 H_2O . Ppt. (Beilstein and Blaese.)

Nickel antimonate, $Ni(SbO_3)_2 + 2H_2O$.

Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 54.)

+5 H_2O . Insol. in H_2O . (Senderens.)

+6 H_2O . Ppt. Insol. in H_2O . (Heffter, Pogg. 86. 446.)

+12 H_2O . Sl. sol. in H_2O . (Heffter.)

Potassium antimonate, $KSbO_3$.

Insol. in H_2O . Sol. in warm $KOH + Aq$, but separates nearly completely on cooling. By boiling with H_2O , or by standing for a long time with cold H_2O , it gradually dissolves as $2KSbO_3 + 5H_2O$, or $K_2H_2Sb_2O_7 + 4H_2O$, or $2KH_2SbO_4 + 3H_2O$.

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

+ H_2O . Insol. in H_2O . (Senderens, Bull. Soc. 1899, (3) 21. 57.)

+1 $\frac{1}{2}$ H_2O (= $2KSbO_3 + 5H_2O$ of Fremy). Easily sol. in H_2O , especially if warm. Solution is pptd. by $NH_4Cl + Aq$. (Freymy, A. ch. (3) 12. 499.)

+2 $\frac{1}{2}$ H_2O . 100 pts. H_2O at 20° dissolve 2.81 pts. anhydrous salt; sp. gr. of solution sat. at 18° = 1.0263. Composition is given as $K_2H_2Sb_2O_7 + 4H_2O$. (Knorre and Olschewsky, B. 20. 3043.)

+3 $\frac{1}{2}$ H_2O . Insol. in H_2O . (Senderens, l.c.)

+4½H₂O. Sol. in H₂O. (Delacroix, J. Pharm. 1897, (6) 6. 533.)

2K₂O, 3Sb₂O₃+10H₂O. Sl. sol. in H₂O. (Delacroix, J. Pharm. 1897, 6. 337.)

+10H₂O. (Delacroix, *l.c.*)

Potassium pyroantimonate, K₄Sb₂O₇.

Deliquescent; decomp. by boiling with H₂O into K₂SbO₃+5H₂O, by cold H₂O into K₂H₂SbO₇+6H₂O. (Fremy.)

Does not exist. (Knorre and Olschewsky.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Potassium hydrogen pyroantimonate, K₂H₂SbO₇.

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

+2½H₂O. (Senderens, Bull. Soc. 1899, (3) 21. 57.)

+3½H₂O. Very difficultly sol. in hot or cold H₂O. (Knorre and Olschewsky, B. 18. 2358.)

+6H₂O. Quite difficultly sol. in cold H₂O. Not precipitated by NH₄Cl+Aq. Aqueous solution gradually decomposes. (Fremy.)

+4H₂O. See 2KSbO₃+5H₂O.

Potassium antimonate sulphantimonate, KSbO₃, K₂SbS₄+5H₂O.

Decomp. on air, and with cold H₂O. Sol. in hot H₂O. (Rammelsberg.)

Silver antimonate.

Insol. in H₂O. (Berzelius.)

AgSbO₃+3H₂O=AgH₂SbO₄+2H₂O. Easily sol. in NH₄OH+Aq, when freshly pptd. (Beilstein and Blaese.)

+1½H₂O. Ppt. (Ebel, B. 22. 3043.)

Silver antimonate ammonia, AgH₂SbO₄, 2NH₃+H₂O.

(Beilstein and Blaese.)

Sodium antimonate, NaSbO₃.

Sol. in much H₂O, but soon becomes decomposed into Na₂H₂SbO₇.

+3½H₂O, composition of Na₂H₂SbO₇+6H₂O, according to Beilstein and Blaese.

1000 pts. H₂O dissolve 0.31 pt. NaSbO₃+3½H₂O at 12.3°.

1000 pts. alcohol of 15.8% dissolve 0.13 pt. NaSbO₃+3½H₂O at 12.3°.

1000 pts. alcohol of 25.6% dissolve 0.07 pt. NaSbO₃+3½H₂O at 12.3°.

Somewhat more sol. when freshly precipitated.

Absolutely insol. in glacial HC₂H₃O₂.

Presence of NaOH or Na salts diminish solubility, while NH₄OH or K salts increase it

slightly. (Beilstein and Blaese, Bull. Ac. St. Petersb. 33. 201.)

+4½H₂O. Sol. in H₂O. (Delacroix, Bull. Soc. 1899, (3) 21. 1051.)

2Na₂O, 3Sb₂O₃+10H₂O. (Delacroix, *l.c.*)

Na₂O, 3Sb₂O₃+11H₂O. (Delacroix, *l.c.*)

Sodium pyroantimonate, Na₂H₂SbO₇+6H₂O.

Boiling H₂O dissolves ⅜ pt. of this salt. (Fremy.) 1000 pts. H₂O dissolve 2.5 pts. salt. (Ebel, B. 22. 3044.) See also NaSbO₃+3½H₂O.

+5H₂O. (Knorre and Olschewsky.)

Strontium antimonate, Sr(SbO₃)₂+6H₂O.

Ppt. Less sol. in H₂O than SrSO₄. (Heffter, Pogg. 86. 418.)

Thallos antimonate, TlSbO₃+2H₂O= TlH₂SbO₄+H₂O.

Somewhat sol. in H₂O, when freshly precipitated; insol. when dried. (Beilstein and Blaese.)

Tin (stannous) antimonate, 2SnO, Sb₂O₃.

Ppt. (Lenssen, A. 114. 113.)

Sn(SbO₃)₂+2H₂O. Attacked with difficulty by acids or alkalies, most easily by hot conc. H₂SO₄. (Schiff, A. 120. 55.)

2SnO, 3Sb₂O₃+4H₂O.

SnO, 2Sb₂O₃.

Tin (stannic) antimonate.

Insol. in H₂O. (Levol, A. ch. (3) 1. 504.)

Uranium antimonate, 5UO₂, 3Sb₂O₃+15H₂O.

Ppt. Sol. in hot conc. HCl+Aq, and in UCl₃+Aq. (Rammelsberg.)

Zinc antimonate, Zn(SbO₃)₂.

Very slightly sol. in H₂O (Berzelius); sol. in solutions of Zn salts.

+2H₂O. (Ebel, Dissert. 1890.)

Insol. in H₂O. (Senderens, Bull. Soc. 1899, (3) 21. 57.)

+5H₂O. Not wholly insol. in cold, moderately sol. in hot H₂O. (Ebel, Dissert. 1890.)

+6H₂O. Insol. in H₂O. (Senderens.)

Antimoniomolybdic acid.

Ammonium antimoniomolybdate, 5(NH₄)₂O, 4Sb₂O₃, 7MoO₃+12H₂O.

Readily sol. in hot H₂O. (Gibbs, Am. Ch. J. 7. 392.)

Antimoniotungstic acid, 3Sb₂O₃, 4WO₃+11H₂O.

Sol. in H₂O. (Hallopeau, C. R. 1896, 123. 1068.)

Potassium antimoniotungstate, 3K₂O, 3Sb₂O₃, 4WO₃+4H₂O.

Much more sol. in hot than in cold H₂O. Decomp. by HCl, H₂SO₄ and HNO₃. (Hallopeau, C. R. 1896, 123. 1066.)

+16H₂O. Much more easily sol. in hot than cold H₂O. Decomp. by HCl, H₂SO₄, and HNO₃. (Hallopeau, l.c.)

6K₂O, 4Sb₂O₃, 12WO₃+25H₂O.

Sl. sol. in H₂O. (Gibbs, Am. Ch. J. 7. 392.)

Antimoniuretted hydrogen.

See Antimony hydride.

Antimonosomolybdic acid.

Ammonium antimonosomolybdate, 6(NH₄)₂O, 3Sb₂O₃, 17MoO₃+21H₂O.

Insol. in cold H₂O. (Gibbs, Am. Ch. J. 7. 313.)

Antimonosophosphotungstic acid.

Potassium antimonosophosphotungstate, 12K₂O, 5Sb₂O₃, 6P₂O₅, 22WO₃+48H₂O.

Nearly insol. in cold or warm H₂O. (Gibbs, Am. Ch. J. 7. 392.)

Antimonosotungstic acid.

Ammonium antimonosotungstate.

Sol. in H₂O.

Barium antimonosotungstate, 4BaO, 6Sb₂O₃, 22WO₃+36H₂O.

Precipitate; very sl. sol. in hot H₂O. (Gibbs, Am. Ch. J. 7. 313.)

Antimonous acid, HSbO₂.

(Long, J. Am. Chem. Soc. 1895, 17. 87.)

+1½H₂O. Ppt. (Schaffner, A. 51. 182.)

H₂SbO₃. Ppt. (Clarke and Stallo, B. 13. 1793.)

Does not exist. (Guntz, C. R. 102. 1472.)

H₂Sb₂O₅. When freshly pptd., is sol. in dil. KOH, and NaOH+Aq. Scarcely sol. in NH₄OH+Aq, or in (NH₄)₂CO₃, or KHCO₃+Aq.

Completely sol. in K₂CO₃, and Na₂CO₃+Aq, especially if warm. When recently pptd. is sl. sol. in succinic acid+Aq.

Calcium antimonite, CaSb₂O₄ (?).

Min. *Romeite*. Insol. in acids.

Cobaltous antimonite (?).

Sl. sol. in H₂O. (Berzelius.)

Cuprous antimonite, Cu₂(SbO₃)₂.

Insol. in H₂O. Sol. in acids; most easily in conc. HCl+Aq. (Hausmann and Stromeyer, Schw. J. 19. 241.)

Cupric antimonite (?).

Insol. in H₂O. (Berzelius.)

CuSb₂O₆. Min. *Ammiolite*.

CuSb₂O₄. Sol. in HCl+Aq, tartaric and citric acids. (Harding, Z. anorg. 1899, 20. 238.)

Iron (ferrous) antimonite (?).

More sol. in H₂O than the antimonate. (Dumas.)

Potassium antimonite, K₂O, 3Sb₂O₃.

Easily decomp. by cold H₂O. Not decomp. by KOH+Aq containing over 20.9% K₂O. (Corimimbœuf, C. R. 115. 1305.)

+3H₂O. As above. (C.)

Potassium antimonite iodide, K₂O, 8Sb₂O₃, 2KI.

Insol. and not decomp. by cold or hot H₂O. Not decomp. by acids or alkalies. Aqua regia decomp. slowly. Tartaric acid dissolves gradually. (Grühl, Dissert. 1897.)

Sodium antimonite, NaSbO₃+3H₂O.

Difficultly sol. in H₂O. (Terreil, A. ch. (4) 7. 380.)

2Na₂O, 3Sb₂O₃+H₂O. Decomp. by H₂O, but not by NaOH+Aq containing 94.3 g. NaOH per l. (Corimimbœuf.)

Na₂O, 2Sb₂O₃. Decomp. by H₂O but not by NaOH+Aq containing 188.6 g. NaOH per l. (C.)

Na₂O, 3Sb₂O₃. Decomp. by H₂O, but not by NaOH+Aq containing 113.2 g. NaOH per l. (C.)

+2H₂O=NaH₂(SbO₂)₃. (Terreil.)

Antimony, Sb.

Does not decomp. H₂O. Not attacked by HCl+Aq (Berzelius); slowly sol. in conc. HCl+Aq (Debray); slowly sol. in conc. warm HCl+Aq (Troost). Attacked by very conc. HCl+Aq only when finely divided (Schützenberger, Willm); very sl. attacked by dil. or conc. acid (Guntz). Not attacked by boiling HCl+Aq (Gmelin). By careful experiments, pure Sb is absolutely insol. in dil. or conc., hot or cold HCl+Aq, except when in contact with oxygen. (Ditte and Metzner, A. ch. (6) 29. 889.)

Insol. in dil. or cold conc., but sol. in hot conc. H₂SO₄. Oxidized but not dissolved by HNO₃+Aq. Easily and completely sol. in aqua regia.

Very slowly attacked by pure HNO₃+Aq of 1.51–1.42 sp. gr.; weaker acid has no marked action whether it contains NO₂ or not. HCl+HNO₃ has no action if dil. or at low temp., but when even very dil. and KNO₃ is added, the action will begin. (Millon, A. ch. (3) 6. 101.)

Not attacked in 10 months by 2% HNO₃+Aq. Sb is not dissolved by HNO₃+Aq of any concentration, a white powder being always left, which is insol. in HNO₃+Aq or H₂O. (Montemartini, Gazz. ch. it. 22. 384.)

Insol. in alkalies+Aq.

Somewhat sol. in distilled H₂O. More or less sol. in solutions of acids, alkalies and salts and in alcohol and ether. Only sl. sol. in a mixture of alcohol and ether. (Ruff and Albert, B. 1905, 38. 54.)

Alkaline H₂O₂ converts Sb into antimonie acid, but neutral H₂O₂ is without action. (Clark, Chem. Soc. 1893, 63. 886.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 826.)

Easily attacked by pyrosulphuryl chloride. (Heumann and Köchlin, B. 16. 479.)

Sb is sol. in a mixture of HNO_3 and tartaric acid or other polybasic acids. (Czerwek, Z. anal. 1906, 45. 507.)

Not attacked by a mixture of alcohol and ether. (Cohen, Z. phys. Ch. 1904, 47. 12.)

$\frac{1}{2}$ cc. oleic acid dissolves 0.0007 g. Sb in 6 days. (Gates, J. phys. Ch. 1911, 15. 143.)

There are three modifications.

1. Ordinary gray metallic.

2. Black amorphous. Unstable at ord. temp. By boiling with H_2O is changed to metallic Sb.

3. Yellow. Very unstable. At -50° goes over rapidly into the ordinary black modification. Sol. in CS_2 at a little above -90° . (Stock, B. 1903, 37. 898.)

Unstable above -90° . (Stock, B. 1905, 38. 3837.)

Antimony arsenide, Sb_2As_3 .

(Descamps, C. R. 86. 1065.)

Antimony tribromide, SbBr_3 .

Deliquescent; decomp. by H_2O .

Very sol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 826.)

Very sol. in warm liquid AsBr_3 , forming a solution with sp. gr. = 3.685 at 47° . (Retgers, Z. phys. Ch. 1893, 11. 339.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, 25. 217.)

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913, 84. 27.)

Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, 25. 211.)

Sol. in alcohol and CS_2 .

Sol. in ether forming two layers. (Hayes, J. Chem. Soc. 1902, 24. 360.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Solubility of SbBr_3 in organic liquids. Data in parentheses indicate labile equilibrium.

Solvent	t°	Mols. per 100	t°	Mols. per 100	t°	Mols. per 100
Benzene	5.6°	0	65°	17.1	91.5°	73.7
	4.5	1.9	75	24.9	90	76.7
	15	3.0	80	30.7	85	84.9
	25	4.3	85	38.4	90	91.4
	35	6.0	90	48.2	92	94.8
	45	8.6	91.5	58.1	94	100
	55	12.1	92.5	66.6		
Chlor-benzene	-45.2°	0	0°	7.2	60°	37.6
	-47	1.7	10	9.2	70	50.0
	-40	2.2	20	11.8	80	66.6
	-30	3.2	30	15.4	90	89.6
	-20	4.3	40	20.8	94	100
	-10	5.6	50	28.1		
Brom-benzene	-31°	0	15°	17.4	75°	65.2
	-32	2.6	25	22.2	85	81.1
	-25	4.4	35	22.7	90	90.0
	-15	6.9	45	34.4	94	100
	-5	9.9	55	42.6		
	5	13.4	65	52.6		

Solubility of SbBr_3 in organic liquids.—Cont.

Solvent	t°	Mols. per 100	t°	Mols. per 100	t°	Mols. per 100
Iod-benzene	-28.6°	0	10°	26.3	70°	67.0
	-30.5	4.0	20	31.5	80	78.2
	-32	8.7	30	37.3	90	91.9
	-20	13.5	40	43.7	94	100
	-10	17.5	50	50.7		
	0	21.7	60	58.5		
Paradi-chlor-benzene	54.5°	0	65°	29.5	85°	68.9
	51.5	6.3	70	37.0	90	85.2
	48.5	12.8	75	45.6	94	100
	55	18.7	80	56.2		
Paradi-brom-benzene	88°	0	65°	52.0	90°	91.8
	85	6.8	70	59.1	92	95.4
	80	18.0	75	66.5	94	100
	75	29.5	80	74.4		
	70	41.5	85	83.0		
Nitro-benzene	6°	0	-5°	32.3	55°	59.1
	1	8.6	5	35.3	65	66.4
	-4	17.0	15	38.8	75	74.9
	-9	24.0	25	42.8	85	86.0
	-15	29.7	35	47.4	90	93.0
	(-17)	(31.9)	45	52.8	94	100
Metadi-nitro-benzene	90°	0	55°	49.1	70°	70.8
	85	8.1	50	53.0	75	76.0
	80	16.2	47.5	54.4	80	81.7
	75	24.2	50	56.1	85	87.8
	70	31.8	55	58.8	90	94.2
	65	38.5	60	62.2	94	100
	60	44.3	65	66.2		
Toluene	-93°	0	10°	28.8	70°	69.4
	-93.5	0.3	20	36.7	80	79.4
	-70	1.2	30	47.5	85	85.2
	-50	2.6	(34)	(54.0)	90	92.6
	-30	5.2	40	51.5	94	100
	-10	13.3	50	56.3		
	-1	22.4	60	62.3		
Ethyl-benzene	-93°	0.1	10°	9.8	60°	59.8
	-60	0.4	20	19.5	70	67.4
	-40	1.0	25	28.6	80	77.4
	-20	2.3	29	37.8	85	85
	-10	3.9	40	44.6	90	92.6
	0	6.4	50	51.6	94	100
Propyl-benzene	-80°	0.4	(-1.5°)	(33.8)	50°	44.3
	-60	1.2	(-20)	(23.3)	60	51.5
	-40	3.4	0	25.8	70	61.5
	-30	5.5	10	27.8	80	73.5
	-20	9.5	20	30.5	90	90
	-10	17.2	30	34.1	94	100
	-5	24.3	40	38.6		
Isoamyl-benzene	-70°	1.9	(-13°)	(24.9)	50°	35.8
	-50	3.6	-10	17	60	43.3
	-40	5.1	0	18.2	70	54.0
	-30	7.1	10	19.9	80	68.5
	-20	13.4	20	22.5	90	90
	-17	16.4	30	25.9	94	100
	(-15)	(19.4)	40	30.3		

(Menschutkin, Ann. Inst. Pol. P. le Gr. 13. 1.)

Antimony bromide with MBr.

See Bromantimonate, M.

Also below.

Antimony hydrogen bromide, SbBr_3 , $\text{HBr} + 3\text{H}_2\text{O}$.

Very hygroscopic. Decomp. by H_2O . (Weinland and Feige, B. 1903, 36. 256.)

See Metabromantimonic acid.

Antimony caesium bromide, $2\text{SbBr}_3, 3\text{CsBr} + 2\text{H}_2\text{O}$.

Loses Br_2 in the air. (Weinland, B. 1903, **36**. 257.)

Antimony calcium bromide, $\text{SbBr}_3, \text{CaBr}_2 + 8\text{H}_2\text{O}$.

Easily decomp. (Benedict, Proc. Am. Acad. 1895, **30**. 9.)

Antimony glucinum bromide, $3\text{SbBr}_3, 2\text{GlBr}_2 + 18\text{H}_2\text{O}$.

Hydroscopic. Easily decomp. (Weinland, B. 1903, **36**. 258.)

Antimony magnesium bromide, $\text{SbBr}_3, \text{MgBr}_2 + 8\text{H}_2\text{O}$.

As Ca salt. (Benedict, Proc. Am. Acad. 1895, **30**. 9.)

Antimony potassium bromide, $10\text{SbBr}_3, 23\text{KBr} + 27\text{H}_2\text{O}$.

(Herty, Am. Ch. J. 1894, **16**. 496.)

Antimony rubidium bromide, $2\text{SbBr}_3, 3\text{RbBr}$.

Decomp. by H_2O ; can be recryst. from dil. $\text{HBr} + \text{Aq}$. (Wheeler, Z. anorg. **5**. 258.)

SbRb_2Br_6 . Slowly loses Br_2 in the air. Decomp. by H_2O . (Weinland, B. 1903, **36**. 259.)

$10\text{SbBr}_3, 23\text{RbBr}$ (?). Cryst. from conc. $\text{HBr} + \text{Aq}$. (Wheeler.)

The composition assigned to this salt by Wheeler (Z. anorg. **5**. 253) is incorrect. (Ephraim, B. 1903, **36**. 1817.)

Antimony vanadium bromide, $\text{SbBr}_3, \text{VBr}_4 + 7\text{H}_2\text{O}$.

Hydroscopic. Decomp. by H_2O . Sol. in dil. HCl and in tartaric acid. (Weinland, B. 1903, **36**. 260.)

Antimony bromide potassium chloride, $\text{SbBr}_3, 3\text{KCl} + 1\frac{1}{2}\text{H}_2\text{O}$.

Slowly deliquescent. Very sol. in H_2O .

Sat. solution contains 120.5 g. to 100 cc. H_2O , and has sp. gr. = 1.9.

Decomp. by much H_2O . (Atkinson, Chem. Soc. **43**. 290.)

Does not exist. (Herty, Am. Ch. J. 1894, **16**. 497.)

See also Antimony chloride potassium bromide.

Antimony bromofluoride, SbF_4Br .

Decomp. by H_2O . (Ruff, B. 1906, **39**. 4319.)

Antimony trichloride, SbCl_3 .

Deliquescent. Decomp. by H_2O with precipitation of SbOCl . This precipitation is prevented by tartaric, citric, or hydrochloric acid, or by conc. solutions of chlorides of alkalies and alkaline earths.

Solubility in H_2O .
100 pts. SbCl_3 sol. in pts. H_2O at t° .

t°	Pts. H_2O
0°	16.6
15°	12.3
20°	10.9
25°	10.1
30°	9.4
35°	8.7
40°	7.3
50°	5.2
60°	2.2

(Meerburg, Z. anorg. 1903, **33**. 299.)

Solubility in $\text{HCl} + \text{Aq}$.
100 mol. H_2O dissolve mol. SbCl_3 in presence of mol. HCl at 20°.

Mol. HCl	Mol. SbCl_3
0	72.1–72.8
2.4	73.0
6.5	67.5
8.4	67.6
8.6	66.5
9.8	65.0
12.2	65.3
29.6	54.5

(Meerburg, Z. anorg. 1903, **33**. 304.)

Solubility in $\text{HCl} + \text{Aq}$.

Solid phase	100 mol. H_2O dissolve at 20°			
	1	2	3	4
	Mol. SbCl_3	Mol. HCl	Mol. SbOCl	Mol. HCl
SbOCl	8.7	7.2	9.8	6.9
	8.6	7.5	16.1	7.9
	19.6	8.0	21.7	7.4
	19.8	8.9	25.0	8.8
			28.0	8.6
$(\text{SbOCl})_x, (\text{SbCl}_3)_y$	37.5	8.7	32.0	7.9
	44.0	6.8	35.8	7.9
	63.7	6.2	59.5	6.4
	69.1	5.6	61.0	6.5
	66.1	4.6	62.7	4.4
	69.8	5.3		
SbCl_3 and $(\text{SbOCl})_x, (\text{SbCl}_3)_y$	69.3	4.3		
	68.3	3.6		

1 & 2. (Meerburg, Z. anorg. 1903, **33**. 302.)

3 & 4. (Noodt, Z. anorg. 1903, **33**. 302.)

Somewhat sol. in liquid $(\text{CN})_2$. (Centnerszwer, Bull. Soc. 1901, (3) **28**. 405.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 826.)

Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, **25**. 211.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, **25**. 217.)

Easily sol. in AsBr_3 . (Walden, Z. anorg. 1902, 29, 374.)

Sol. in alcohol without decomp. Very sol. in hot CS_2 , but solubility diminishes rapidly on cooling. (Cooke, Proc. Am. Acad. 18, 72.)

1 g. SbCl_3 is sol. in 0.186 g. acetone at 18° . Sp. gr. of sat. solution $18^\circ/4^\circ = 2.216$. (Naumann, B. 1904, 37, 4332.)

Sol. in ethyl acetate. (Naumann, B. 1904, 37, 3601.)

1 pt. sol. in 16.97 pts. of ethyl acetate at 18° . Sp. gr. of sat. solution $18^\circ/4^\circ = 1.7968$. (Naumann, B. 1910, 43, 320.)

Sol. in benzonitrile. (Naumann, B. 1914, 47, 1369.)

Sol. in methylal. (Eidmann, C. C. 1899, II, 1014.)

Solubility of SbCl_3 in organic liquids.
Data in parentheses indicate labile equilibrium

Solvent	t°	Mols. per 100	t°	Mols. per 100	t°	Mols. per 100
Benzene	5.6°	0	50°	27.2	75°	78.5
	4	2.6	60	34.7	70	83.3
	1	7.1	70	45.2	62	89.3
	10	10.1	75	53.1	67.5	94.2
	20	13.1	77.5	58.7	73	100
	30	16.8	79	66.6		
	40	21.4	77.5	73.4		
Chlorobenzene	-45.2°	0	-10°	14.4	30°	47.1
	-47	2.2	-5	19.4	40	56.2
	-40	3.6	0	28.1	50	66.6
	-30	6.0	(4)	(41.1)	60	78.7
	-20	9.0	10	32.5	70	94.3
	-15	11.6	20	38.7	73	100
Bromobenzene	-31°	0	-5°	21.7	40°	59.2
	-32.5	3.4	0	26.6	50	68.8
	(-35)	(6.4)	3	31.8	60	80.6
	-30	4.8	(0)	(41.9)	65	87.2
	-25	7.6	(7)	(50.0)	70	95.0
	-20	10.7	10	36.4	73	100
	-15	14.1	20	43.2		
	10	17.8	30	50.8		
Iodobenzene	-28.6°	0	(-5°)	(40.7)	25°	53.9
	-30	2.4	-34.5	10.7	35	60.4
	(-35)	(11.7)	-25	16.4	45	67.5
	(-40)	(20.8)	-15	24.7	55	76.2
	(-45)	(27.2)	-5	39.1	65	87.4
	(-35)	(30.9)	(-3)	(47.2)	70	95.0
	(-25)	(33.9)	5	44.5	73	100
	(-15)	(37.2)	15	48.7		
Para-chlorobenzene	54.5°	0	39.5°	29.5	60°	66.5
	50	6.3	45	37.5	65	74.1
	45	15.5	50	46.4	70	91.1
	40	28.0	55	56.0	73	100
Para-bromobenzene	88°	0	65°	45.4	95°	97.1
	85	5.9	60	53.8	70	95.2
	80	15.8	49.5	64.9	73	100
	75	25.7	55	72.5		
	70	35.7	60	79.8		
Nitrobenzene	6°	0	-13.5°	27.3	15°	59.2
	2	7.0	-10.5	29.8	25	63.0
	-2	12.1	-7.5	35.2	35	67.6
	-6	16.5	-4.5	40.7	45	72.8
	-10	20.3	-3	50.0	55	79.0
	-14	23.5	-0.5	52.9	65	87.2
	(-18)	(26.2)	-5	53.0	70	92.7
	-16.5	25.2	5	55.9	73	100

Solubility of SbCl_3 in organic liquids—Cont.

Solvent	t°	Mols. per 100	t°	Mols. per 100	t°	Mols. per 100
Metadi-nitrobenzene	90°	0	(-10°)	(57.7)	(0°)	(78.1)
	80	14.3	(10)	(62.4)	20	65.2
	70	25.3	(27.5)	(44.5)	30	68.8
	60	33.8	(28.5)	(50.0)	40	73.2
	40	45.6	27.5	55.0	50	78.5
	(20)	(53.6)	25	60.2	60	85.8
	(1)	(59.9)	(20)	(66.2)	70	95.2
	(-11)	(62.2)	(10)	(73.5)	73	100
Toluene	-93°	0	-10°	14.4	40°	59.3
	-94	0.5	0	22.1	42.5	66.6
	-70	1.4	6	28.6	40	71.1
	-50	3.3	11	35.7	50	77.1
	-40	5.1	(-8)	(27.0)	60	83.8
	-30	7.2	20	40.5	70	94.7
	-20	10	30	47.6	73	100
Ethylbenzene	-93°	0.1	35°	36.4	(35.8°)	(68.1)
	-50	0.6	39	50	(33)	(65.7)
	-30	1.1	37	57.7	40	70.3
	-10	3.6	35	61.8	50	77.3
	0	5.6	(33)	(65.7)	60	85.5
	10	9.4	(15)	(37.8)	65	90.3
	20	16.8	(25)	(47.5)	70	95.6
	30	27.2	37	66.6	73	100
Propylbenzene	(-70°)	(0.6)	-70°	0.2	8.5°	53.2
	(-50)	(2.8)	-50	1.5	10	53.6
	(-40)	(5.2)	-40	3.0	20	56.9
	(-30)	(8.8)	-30	5.5	30	60.6
	(-20)	(14.8)	-20	9.7	40	65.5
	(-10)	(25.1)	-10	16.2	50	72
	(-5)	(32.4)	-5	20.5	60	81
	(0)	(43.3)	0	26.2	65	86.8
	(1.5)	(50)	5	35.6	70	95.1
	(1)	(51.1)	7	41.6	73	100
Isoamylbenzene	-80°	3	(-45°)	(17.1)	0°	46.3
	-70	5.4	(-35)	(22.8)	10	46.8
	-60	8.4	-25	29.3	20	52.5
	-50	12.4	-15	36.6	30	57.3
	-40	17.9	-5	45.6	40	63.4
	(-30)	(27.3)	(0)	(52.8)	50	71.4
	(-25)	(34.4)	(5)	(60.3)	60	81.7
	(-22)	(40.7)	(7.5)	(66.6)	65	88
	(-20.5)	(50)	(-21)	(44.2)	70	95.5
	(-22)	(54)	(-10)	(44.9)	73	100

(Menschutkin, Ann. Inst. Pol. P.-le Gr., 19, 1.)

Antimony hydrogen trichloride, 2SbCl_3 , $\text{HCl} + 2\text{H}_2\text{O}$.

Deliquescent. Decomp. by H_2O . Melts in crystal H_2O at 16° . (Engel, C. R. 106, 1797.)

Antimony pentachloride, SbCl_5 .

Deliquesces to $\text{SbCl}_3 + 4\text{H}_2\text{O}$, which can be crystallized out of a little H_2O . Decomp. by more H_2O into SbO_2Cl . Sol. in a large amt. of H_2O , if it is added all at one time. Precipitation by H_2O is also hindered by presence of tartaric, or hydrochloric acid.

+ H_2O . Deliquescent. Sol. in chloroform. (Anschütz and Evans, A. 239, 285.)

+ $4\text{H}_2\text{O}$. Insol. in chloroform. (Anschütz and Evans.)

Antimony pentachloride with MCl .

See Chlorantimonate, M.

See also below.

Antimony hydrogen pentachloride, SbCl_5 , $\text{HCl} + 4\frac{1}{2}\text{H}_2\text{O}$.

"*Metachlorantimonic acid*" according to Weinland and Schmid, (Z. anorg. 1905, **44**. 43.)

Very easily sol. in H_2O , alcohol, acetone and glacial acetic acid. Aqueous solution decomp. on standing with separation of Sb_2O_3 , but remains clear in presence of 10% HCl . (Weinland and Schmid, Z. anorg. 1905, **44**. 43.)

SbCl_5 , $5\text{HCl} + 10\text{H}_2\text{O}$. Not deliquescent. Decomp. by H_2O . Melts in crystal H_2O at about 55° . (Engel, C. R. **106**. 1797.)

Antimony antimonyl chloride, SbCl_3 , SbOCl .

More easily attacked by H_2O than SbOCl . (Bemmelen, Z. anorg. 1903, **33**. 293.)

Antimony antimonyl potassium chloride, SbCl_3 , SbOCl , 2KCl .

Not deliquescent. Immediately decomp. by hot or cold H_2O ; sol. in hot glacial $\text{HC}_2\text{H}_3\text{O}_2$, or in HCl , or tartaric acid + Aq.

Insol. in KCl + Aq, hot or cold alcohol, CS_2 , or ligroine. (Benedikt, Proc. Am. Acad. **29**. 217.)

Antimony antimonyl rubidium chloride, SbCl_3 , SbOCl , 2RbCl .

Sol. in very dil. HCl + Aq. (Wells, Am. J. Sci. 1897, (4) **3**. 463.)

Antimony barium chloride, SbCl_3 , $\text{BaCl}_2 + \frac{1}{2}\text{H}_2\text{O}$.

Decomp. by H_2O .

Antimony caesium chloride, SbCl_3 , 6CsCl .

Decomp. by H_2O . Cryst. from dil. HCl + Aq. (Godeffroy, Arch. Pharm. (3) **12**. 47.)

2SbCl_3 , 3CsCl . Decomp. by H_2O ; sl. sol. in cold, easily in hot dil. HCl + Aq. This is identical with the above salt. (Saunders, Am. Ch. J. **14**. 152.)

SbCl_3 , 2CsCl . Sol. in boiling conc. HCl + Aq without decomp. (Setterberg, Oef. Vet. Akad. 1882, **6**. 23.)

SbCl_3 , CsCl . Cryst. from HCl + Aq without decomp. Decomp. by H_2O . (Setterberg, Oef. Vet. Akad. 1882, **6**. 27.)

Antimony calcium chloride, SbCl_3 , $\text{CaCl}_2 + 8\text{H}_2\text{O}$.

Easily decomp. (Benedict, Proc. Am. Acad. 1895, **30**. 9.)

SbCl_3 , $\text{CaSbCl}_6 \cdot \text{OH} + 9\text{H}_2\text{O}$. Deliquescent; sl. sol. in H_2O . (Weinland, B. 1901, **34**. 2635.)

Antimony chromium chloride,

CrCl_3 , $3\text{SbCl}_3 + 13\text{H}_2\text{O}$. (Weinland.) should be

$[\text{SbCl}_4]_3[\text{Cr}(\text{OH})_3] + 7\text{H}_2\text{O}$; and CrCl_3 , $\text{SbCl}_3 + 10\text{H}_2\text{O}$ should be

$[\text{SbCl}_4][\text{Cr}(\text{OH})_3\text{Cl}_2] + 6\text{H}_2\text{O}$.

(Pfeiffer, Z. anorg. 1903, **36**. 349.)

Antimony glucinum chloride, SbCl_3 , $\text{GlCl}_2 + 3\text{H}_2\text{O}$.

Very hygroscopic. Decomp. by H_2O . Very easily sol. in HCl . (Ephraim, B. 1903, **36**. 1822.)

+ $4\text{H}_2\text{O}$. Ppt. Decomp. by H_2O . Sol. in HCl . (Ephraim, B. 1903, **36**. 1822.)

Antimony hydrazine chloride, SbCl_3 , $3\text{N}_2\text{H}_4\text{Cl}$.

Sol. in conc. HCl + Aq; decomp. by H_2O . (Ferratini, C. A. **1912**, 1613.)

Antimony lithium chloride, SbCl_3 , $2\text{LiCl} + 5\text{H}_2\text{O}$.

Hygroscopic. Decomp. by H_2O . Very easily sol. in HCl . (Ephraim, B. 1903, **36**. 1821.)

+ $6\text{H}_2\text{O}$. Decomp. by H_2O ; easily sol. in HCl . (Ephraim, B. 1903, **36**. 1822.)

Antimony magnesium chloride, SbCl_3 , $\text{MgCl}_2 + 5\text{H}_2\text{O}$.

Hydroscopic. Decomp. by H_2O . Can be cryst. from HCl without decomp. (Ephraim, B. 1903, **36**. 1823.)

2SbCl_3 , MgCl_2 . Hygroscopic. Decomp. by H_2O . Very sol. in HCl . (Ephraim.)

SbCl_3 , $\text{MgSbCl}_6 \cdot \text{MgOH} + 17\text{H}_2\text{O}$. Hydroscopic. Sol. in H_2O with decomp. (Weinland, B. 1901, **34**. 2635.)

Antimony nitrosyl chloride, SbCl_3 , NOCl .

Very deliquescent; decomp. by pure H_2O ; sol. in H_2O containing tartaric acid. (Weber, Pogg. **123**. 347.)

2SbCl_3 , 5NOCl . Decomp. by H_2O . (Sudborough, Chem. Soc. **59**. 661.)

Antimony phosphorus chloride, SbCl_3 , PCl_3 .

Deliquescent. (Weber, Pogg. **125**. 78.)

Antimony phosphoryl chloride, SbCl_3 , POCl_3 .

Deliquescent. (Weber.)

Antimony platinum potassium chloride, $(\text{Sb}, \text{Pt})\text{Cl}_6\text{K}_2$.

Ppt. (Weinland, B. 1905, **38**. 1086.)

Antimony potassium chloride, SbCl_3 , 2KCl .

Sol. in H_2O without decomp. (Jacquelin, A. ch. (2) **66**. 128.)

Not deliquescent. Immediately decomp. by hot or cold H_2O . Sol. in HCl , or tartaric acid + Aq. (Benedikt, Proc. Am. Acad. **29**. 219.)

+ $2\text{H}_2\text{O}$. Very efflorescent.

SbCl_3 , 3KCl . Deliquescent. Decomp. by hot H_2O . (Poggiale.)

+ $2\text{H}_2\text{O}$. (Romanis, C. N. **49**. 273.)

Not obtained by Benedikt (*l.c.*)

10SbCl_3 , 23KCl . True composition of above salts. Sol. in H_2O . (Herty, Am. Ch. J. 1894, **16**. 495.)

SbCl_3 , 2KCl is the only true compound, all

others being isomorphous mixtures. (Jordis, B. 1903, **36**. 2539.)

$2\text{SbCl}_4, 3\text{KCl}$. Deliquescent. Decomp. by H_2O . (Bosek, Chem. Soc. 1895, **67**. 516.)

$\text{SbCl}_4, \text{KSbCl}_4, \text{KOH}$. Hydrosopic. Sol. in H_2O with decomp. (Weinland, B. 1901, **34**. 2635.)

See also Antimony antimonyl potassium chloride.

Antimony rubidium chloride, $\text{SbCl}_3, \text{RbCl}$.

Decomp. on air or with H_2O . (Saunders, Am. Ch. J. **14**. 162.)

$2\text{SbCl}_3, \text{RbCl} + \text{H}_2\text{O}$. Decomp. on air. (Wheeler, Z. anorg. **5**. 253.)

$\text{SbCl}_3, 6\text{RbCl}$. Decomp. by H_2O . (Godefroy, Arch. Pharm. (3) **9**. 343.)

Formula is $10\text{SbCl}_3, 23\text{RbCl}$ (?). (Saunders Am. Ch. J. **14**. 159.)

$10\text{SbCl}_3, 23\text{RbCl}$ (?). Decomp. by H_2O ; sol. in $\text{HCl} + \text{Aq}$. (Saunders.)

Formula is $3\text{SbCl}_3, 7\text{RbCl}$. (Wells and Foote, Am. J. Sci. 1897, (4) **3**. 461.)

Composition assigned to this salt by Saunders (Am. Ch. J. **14**. 155) is incorrect. (Ephraim, B. 1903, **36**. 1817.)

$3\text{SbCl}_3, 5\text{RbCl}$. As above. (Saunders.)

Formula is $2\text{SbCl}_3, 3\text{RbCl}$. (Wheeler.)

Rb_2SbCl_6 . Ppt. Decomp. by H_2O . (Weinland, B. 1905, **38**. 1083.)

$\text{Rb}_2\text{SbCl}_6, 2\text{Rb}_2\text{SbCl}_6$. Ppt. Decomp. by H_2O . (Weinland, B. 1901, **34**. 2635.)

Antimony selenium chloride, $\text{SbCl}_3, \text{SeCl}_4$.

Deliquescent. (Weber.)

Antimony selenyl chloride, $\text{SbCl}_3, \text{SeOCl}_2$.

Very deliquescent. (Weber, Pogg. **125**. 325.)

Antimony sodium chloride, $\text{SbCl}_3, 3\text{NaCl}$ (?).

Decomp. by much H_2O . (Poggiale.)

Antimony sulphur chloride, $2\text{SbCl}_3, 3\text{SCl}_2$.

Decomp. by H_2O .

$\text{SbCl}_3, \text{SCl}_2$. Sol. in dil. $\text{HNO}_3 + \text{Aq}$.

Mpt. $125-126^\circ$ in an atmos. of chlorine. Violently decomp. by H_2O . (Ruff, B. 1904, **37**. 4515.)

Antimony thallium chloride, $\text{SbCl}_3, 3\text{TlCl}$.

Ppt. (Ephraim, Z. anorg. 1909, **61**. 249.)

$\text{SbCl}_3, \text{TlCl}$. (Ephraim and Barteczko, Z. anorg. 1909, **61**. 251.)

$2\text{SbCl}_3, 2\text{TlCl}, \text{TlCl}_3$. Slowly decomp. by cold H_2O . (Ephraim and Barteczko, Z. anorg. 1909, **61**. 253.)

Antimony trichloride ammonia, $\text{SbCl}_3, \text{NH}_3$.

Not very deliquescent. Decomp. by H_2O .

Antimony pentachloride ammonia, $\text{SbCl}_5, 6\text{NH}_3$.

Decomp. by H_2O . (Persoz.)

Antimony pentachloride cyanhydric acid, $\text{SbCl}_5, 3\text{HCN}$.

Deliquescent; decomp. by H_2O . (Klein, A. **74**. 85.)

Antimony pentachloride nitric oxide, $2\text{SbCl}_5, \text{NO}$.

Decomp. by H_2O . (Besson, C. R. **108**. 1012.)

Antimony pentachloride nitrogen peroxide, $3\text{SbCl}_5, 2\text{NO}_2$.

Decomp. by H_2O . (Besson.)

Antimony pentachloride nitrogen sulphide, $\text{SbCl}_5, \text{N}_4\text{S}_4$.

Easily decomp. (Davis, Chem. Soc. 1906, **89**. 1577.)

Decomp. by cold H_2O , HCl , H_2SO_4 and warm alcohol, also by boiling with $\text{KOH} + \text{Aq}$. Almost insol. in organic solvents. (Wölbling, Z. anorg. 1908, **57**. 283.)

Antimony chloride potassium bromide, $\text{SbCl}_3, 3\text{KBr} + 1\frac{1}{2}\text{H}_2\text{O}$.

Very deliquescent. Decomp. by much H_2O . (Atkinson, Chem. Soc. **43**. 289.)

$2\text{SbCl}_3, 3\text{KBr} + 2\text{H}_2\text{O}$. (Atkinson.)

$\text{SbCl}_3, \text{KBr} + \text{H}_2\text{O}$. (Atkinson.)

Above are mixtures. (Herty, Am. Ch. J. 1894, **16**. 497.)

See Antimony bromide potassium chloride.

Antimony chlorofluoride, SbCl_2F_2 .

(Swarts, Z. anorg. 1896, **12**. 71.)

Antimony fluoiodide, SbF_2I .

Slowly decomp. by H_2O . (Ruff, B. 1906, **39**. 4321.)

$(\text{SbF}_2)_2\text{I}$. Sol. in H_2O with pptn. of I_2 . (Ruff, B. 1906, **39**. 4321.)

Antimony trifluoride, SbF_3 .

Deliquescent. Sol. in H_2O .

Solubility in H_2O at t° .

t°	100 g. of the solution contain g. SbF_3	100 g. H_2O contain g. SbF_3
0°	79.37	384.7
20	81.64	444.7
22.5	81.91	452.8
25	83.12	492.4
30	84.93	563.6

(Rosenheim, Z. anorg. 1909, **61**. 189.)

Solubility in $\text{HF} + \text{Aq}$ at 0° .

Normality of $\text{HF} + \text{Aq}$	100 g. H_2O of the HF solution dissolve g. SbF_3
2	474.9
1	432.5
0.5	404.0

(Rosenheim, Z. anorg. 1909, **61**. 192.)

Solubility of SbF_3 in salts + Aq at 0° .

Salt	Normality of salt solution	100 g. H_2O of the salt solution dissolve g. SbF_3
KCl	1	461.8
	0.5	448.3
	0.25	431.9
	0.125	407.3
KBr	1	448.7
	0.5	450.0
	0.25	455.6
	0.125	417.2
KNO_3	1	458.2
	0.5	451.9
	0.25	418.3
	0.125	401.4
$\frac{1}{2}\text{K}_2\text{SO}_4$	1	419.9
	0.5	408.5
	0.25	406.6
$\frac{1}{2}\text{K}_2\text{C}_2\text{O}_4$	1	465.7
	0.5	481.2
	0.25	451.3
	0.125	405.2
$\frac{1}{2}(\text{NH}_4)_2\text{C}_2\text{O}_4$	0.5	431.9
	0.25	442.3
	0.125	433.3
$\frac{1}{2}\text{K}_2\text{C}_4\text{H}_4\text{O}_6$	1	461.4
	0.5	430.5
	0.25	430.8
	0.125	435.2

(Rosenheim, Z. anorg. 1909, 61. 192.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 826.)

Antimony pentafluoride, SbF_5 .

Sol. in H_2O . (Marignac, A. 145. 239.)
 Very hygroscopic; bpt. 155° . Sol. in H_2O with hissing. (Ruff, B. 1904, 37. 678.)
 $+2\text{H}_2\text{O}$. (Ruff, B. 1904, 37. 679.)

Antimony pentafluoride diantimony trifluoride, $\text{Sb}_3\text{F}_{11} = 2\text{SbF}_5, \text{SbF}_3$.

Hygroscopic; bpt. 390° . Easily sol. in H_2O . (Ruff, B. 1904, 37. 680.)

Antimony pentafluoride pentaantimony trifluoride, $\text{SbF}_5, 5\text{SbF}_3$.

B pt. 384° (corr.). (Ruff, B. 1904, 37. 681.)

Antimony caesium fluoride,

$\text{CsF}, 2\text{SbF}_5$.

$\text{CsF}, 3\text{SbF}_5$.

$4\text{CsF}, 7\text{SbF}_5$.

CsF, SbF_5 .

$2\text{CsF}, \text{SbF}_5$.

(Wells, Am. J. Sci. 1901, (4) 11. 451.)

Antimony lithium fluoride, $\text{SbF}_3, 2\text{LiF}$.

Sol. in more than 20 pts. H_2O . (Flückinger, Pogg. 87. 245.)

SbF_3, LiF . Easily sol. in H_2O . (Stein, Chem. Z. 13. 357.)

Antimony potassium fluoride, $\text{SbF}_3, 2\text{KF}$.

Sol. in less than 2 pts. boiling, and in 9 pts. cold H_2O . Insol. in alcohol or ether.

SbF_3, KF . More sol. than $\text{SbF}_3, 2\text{KF}$. Sol. in 2.8 pts. H_2O . (Flückinger, Pogg. 87. 245.)

SbF_3, KF . Easily sol. in H_2O .

$\text{SbF}_3, 2\text{KF} + 2\text{H}_2\text{O}$. Easily sol. in H_2O . (Marignac, A. 145. 239.)

Antimony sodium fluoride, $\text{SbF}_3, 3\text{NaF}$.

Sol. in 14 pts. cold, and 4 pts. boiling H_2O . Sol. in HF . (Flückinger, Pogg. 87. 245.)

SbF_3, NaF . 100 pts. cold H_2O dissolve 93 pts. 100 pts. hot H_2O dissolve 166 pts. (Stein, Wagners' J. B. 1887. 1160.)

$4\text{SbF}_3, \text{NaF}$. As NH_4 salt. (Raad and Hauser, B. 1890, 23. R. 125.)

$\text{SbF}_3, 2\text{NaF}$. Easily sol. in H_2O . (Marignac, A. 145. 329.)

Antimony thallium fluoride, TlF, SbF_3 .

Sol. in H_2O without decomp. (Ephraim, B. 1909, 42. 4458.)

$\text{TlF}, 2\text{SbF}_3$. Sol. in H_2O without decomp. (Ephraim.)

$\text{TlF}, 3\text{SbF}_3$. Sol. in H_2O without decomp. Decomp. by cold conc. H_2SO_4 . (Ephraim.)

Antimony trifluoride ammonia, $\text{SbF}_3, 2\text{NH}_3$.

Sl. sol. in liquid NH_3 . (Ruff, B. 1906, 39. 4326.)

Antimony trifluoride ammonium chloride.

$\text{SbF}_3, \text{NH}_4\text{Cl}$.

Easily sol. in H_2O . (de Haen, B. 21. 901 R.)

Antimony trifluoride ammonium sulphate, $\text{SbF}_3, (\text{NH}_4)_2\text{SO}_4$.

More sol. than K or Na salt. 1 pt. H_2O dissolves 1.4 pts. at 24° and 15 pts. at 100° . (de Haen, B. 21. 902 R.)

Antimony fluoride lithium chloride, $\text{SbF}_3, \text{LiCl}$.

Sol. in H_2O . (Stein, Chem. Z. 13. 357.)

Antimony pentafluoride nitrosyl fluoride, SbF_5, NOF .

Hygroscopic. Decomp. by H_2O . Sol. in liquid NH_3 with decomp. Sl. sol. in NOCl , SiCl_4 , PCl_5 , AsCl_3 , SO_2Cl_2 and SOCl_2 . (Ruff, Z. anorg. 1908, 58. 334.)

Antimony trifluoride potassium chloride, SbF_3, KCl .

100 pts. H_2O dissolve 51 pts. at 24° , and 300 pts. at 100° . (de Haen, B. 21. 901 R.)

Antimony trifluoride potassium sulphate, $\text{SbF}_3, \text{K}_2\text{SO}_4$.

Sol. in H_2O . (de Haen.)
 $2\text{SbF}_3, \text{K}_2\text{SO}_4$. Very sol. in H_2O . (Mayer, B. 1894, 27. R. 922.)

Antimony trifluoride sodium chloride, $\text{SbF}_3, \text{NaCl}$.

Easily sol. in H_2O . (de Haen, B. 21. 901 R.)

Antimony trifluoride sodium sulphate, $\text{SbF}_3, \text{Na}_2\text{SO}_4$.

Sol. in H_2O . (de Haen.)

Antimony fluoiodide, SbF_2I .

Mpt. 80° ; slowly decomp. by H_2O . (Ruff, B. 1906, 39. 4321.)

$(\text{SbF}_2)_2\text{I}$. Mpt. $110-115^\circ$; decomp. by H_2O . (Ruff.)

Antimony fluosulphide, SbF_2S .

Very hygroscopic. Decomp. by H_2O . Sol. with decomp. in alcohol. Sol. in CCl_4 . (Ruff, B. 1906, 39. 4322.)

Antimony gold, Au_2Sb .

Insol. in equal pts. of HNO_3 and tartaric acids. (Roessler, Z. anorg. 1895, 9. 72.)

Antimony hydride, SbH_3 .

Scarcely sol. in H_2O . 1000 ccm. H_2O absorb 4.12 cc. SbH_3 at 10.5° . Decomp. by long contact with H_2O ; also by conc. H_2SO_4 or $\text{KOH} + \text{Aq}$. (Jones, Chem. Soc. 29. 641.)

Antimony trihydroxide, $\text{Sb}_2\text{O}_3, 2\text{H}_2\text{O} = \text{Sb}_2\text{O}(\text{OH})_4$.

(Schaffner, A. 51. 182.)

$\text{Sb}(\text{OH})_3$. Ppt. (Clarke and Stolla, B. 13. 1787.)

Does not exist. (Guntz, C. R. 102. 1472.)
 See **Antimonous acid and antimony trioxide**.

Antimony triiodide, SbI_3 .

Decomp. by H_2O or 80% alcohol. Sol. in $\text{HI} + \text{Aq}$; sol. in boiling CS_2 , and in boiling benzene, but separates out on cooling. Almost insol. in CHCl_3 . (Cooke, Proc. Am. Acad. (2) 5. 72.)

Easily sol. in AsBr_3 . (Walden, Z. anorg. 1902, 29. 374.)

Sol. in warm AsBr_3 . Sp. gr. of a solution sat. at 40° , which solidifies at 37° , = 3.720. This dissolves further AsI_3 , whereby the mpt. sinks to 31° and sp. gr. rises to 3.801. By mixing the latter solution with a solution of AsI_3 in CH_2I_2 , a liquid can be obtained with a sp. gr. of 3.702 at 20° . (Retgers, Z. phys. Ch. 1893, 11. 340.)

Sol. in PCl_3 . (Beckmann, Z. anorg. 1906, 51. 110.)

Sol. in SO_2Cl_2 . (Walden, Z. anorg. 1900, 25. 215.)

Sol. in SOCl_2 and S_2Cl_2 . (Walden, Z. anorg. 1900, 25. 216.)

Sol. in AsCl_3 . (Walden, Z. anorg. 1900, 25. 214.)

Sol. in SnCl_4 . (Walden, Z. anorg. 1900, 25. 218.)

Sol. in POCl_3 . (Walden, Z. anorg. 1900, 25. 212.)

Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, 25. 211.)

Partly sol. in, and partly decomp. by alcohol or ether. (M'Ivor, Chem. Soc. (2) 14. 328.)

Insol. in oil of turpentine and CCl_4 .

100 pts. methylene iodide dissolve 11.3 pts. SbI_3 at 12° ; sp. gr. of solution = 3.453. (Retgers, Z. anorg. 3. 343.)

Sol. in C_6H_6 . (Retgers, Z. phys. Ch. 1893, 11. 334.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Antimony pentaiodide, SbI_5 .

Very unstable. (Pendleton, C. N. 48. 97.)

Antimony barium iodide, $\text{SbI}_3, \text{BaI}_2 + 9\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in HCl , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{H}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq}$. CS_2 dissolves out SbI_3 . (Schäffer, Pogg. 109. 611.)

Antimony caesium iodide, $2\text{SbI}_3, 3\text{CsI}$.

Sl. sol. in $\text{HI} + \text{Aq}$. Exists in two distinct forms. (Wells, Am. J. Sci. 1901, (4) 11. 455.)

Antimony potassium iodide, $2\text{SbI}_3, 3\text{KI} + 3\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in HCl , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{H}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq}$. CS_2 dissolves out SbI_3 . (Schäffer, Pogg. 109. 611.)

$\text{SbI}_3, 2\text{KI} + 2\frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O . (Nicklès, J. Pharm. (3) 39. 116.)

Antimony rubidium iodide, $2\text{SbI}_3, 3\text{RbI}$.

Decomp. by H_2O . (Wheeler, Z. anorg. 5. 259.)

Antimony sodium iodide, $2\text{SbI}_3, 3\text{NaI} + 12\text{H}_2\text{O}$.

As $2\text{SbI}_3, 3\text{KI}$. (Schäffer, Pogg. 109. 611.)

Antimony thalious iodide, $2\text{SbI}_3, 3\text{TlI}$.

Decomp. by H_2O and by $\text{HCl} + \text{Aq}$, also by alcohol. (Ephraim, Z. anorg. 1908, 58. 354.)

Antimony nitride, SbN .

Decomp. by heat. (Franz Fischer, B. 1910, 43. 1471.)

Antimony trioxide, Sb_2O_3 .

Very sl. sol. in H_2O . Sol. in 8900–10,000 pts. H_2O at 100° ; 55,000–61,100 pts. at 15° . (Schulze, J. pr. (2) 27. 320.)

Sol. in $\text{HCl} + \text{Aq}$. Insol. in $\text{HNO}_3 + \text{Aq}$, but not as insol. as metastannic acid. Sol. in cold fuming HNO_3 or H_2SO_4 . Insol. in dil., but sol. in conc. alkalies, or alkali carbonates +

l. in cold NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$. 15 pts. boiling SbCl_3 . (Schneider, 8. 407.)

l. $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{H}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq}$, and from these solutions by H_2O . Easily sol. in benzoic acid. Insol. in pyrotartaric acid. Very sol. in $\text{KHC}_4\text{H}_4\text{O}_6 + \text{Aq}$. Sol. in

what sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$. (Köhler, 385, 258. 520.)

in liquid NH_3 . (Gore, Am. Ch. J. 826.)

l. lactic acid. (Kretzschmar, Ch. Z. 943.)

in grape sugar solution to which has been added. (Vogel, B. 1885, 3.)

in acetone. (Naumann, B. 1904, 37. dmann, C. C. 1899, II. 1014.)

l. glycerine in presence of alkalies. (Dingl. 1885, 258. 520.)

in a sol. colloidal modification. (B. 16. 1142.)

Valentinite, Senarmontite.

l. See Antimonous acid.

y tetroxide, Sb_2O_4 .

in H_2O . Slightly attacked by acids; $\text{HCl} + \text{Aq}$ acts only slightly. (Fresenius.)

Terzantite. Sl. sol. in $\text{HCl} + \text{Aq}$.

y pentoxide, Sb_2O_5 .

in H_2O . Easily sol. in $\text{HCl} + \text{Aq}$. Sl. sol. in conc. $\text{KOH} + \text{Aq}$.

"monoxyd" is sol. in glycerine in presence of alkalies.

glycerine, to which have been added $\text{NaOH} + \text{Aq}$ (1 : 1), dissolve 20.6 g. at b.-pt.; 20 g. $\text{NaOH} + \text{Aq}$ (1 : 1), dissolve 68.5 g. at b.-pt.; 40 g. $\text{NaOH} + \text{Aq}$ (1 : 1), dissolve 93.0 g. at b.-pt.; 80 g. $\text{NaOH} + \text{Aq}$ (1 : 1), dissolve 119.2 g. at b.-pt. (Dingl. 258. 520.)

See Antimonic acid.

y nitrogen pentoxide, $2\text{Sb}_2\text{O}_5, \text{N}_2\text{O}_5$. Decomp. by H_2O . (Thomas, C. R. 10. 1116.)

y oxybromide.

Antimonyl bromide.

y oxychloride.

Antimonyl chloride.

y oxyfluoride.

Antimonyl fluoride.

y oxysulphide, Sb_2OS_2 .

Antimony blende (kermesite).

in H_2O or dil. acids, except $\text{HCl} + \text{Aq}$. (Pogg. 110. 147.)

Antimony palladium, Sb_2Pd .

Sl. sol. in equal pts. of HNO_3 and tartaric acids. (Roessler, Z. anorg. 1895, 9. 69.)

Antimony platinum, Sb_2Pt .

Insol. in equal pts. of HNO_3 and tartaric acids. (Roessler, Z. anorg. 1895, 9. 67.)

Antimony phosphide, SbP .

Insol. in benzene, ether, or CS_2 . (M'Ivor, B. 6. 1362.)

Antimony selenide, SbSe .

(Chrétien, C. R. 1906, 142. 1341.)

Sb_2Se_4 . (Chrétien, l.c.)

Sb_4Se_6 . (Chrétien, l.c.)

Sb_2Se_3 . Sol. in $\text{KOH} + \text{Aq}$. (Hofacker, A. 107. 6.)

Sb_2Se_5 . (Hofacker.)

Antimony selenide, with M selenide.

See Selenoantimonates, M.

Antimony trisulphide, Sb_2S_3 (Kermes).

Insol. in H_2O and dil. acids.

1 l. H_2O dissolves 5.2×10^{-4} mols. pptd. Sb_2S_3 at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

Decomp. by conc. HNO_3 or H_2SO_4 . Sol. in conc. $\text{HCl} + \text{Aq}$. Easily sol. in dil. KOH , NaOH , $(\text{NH}_4)_2\text{S}$, and $\text{K}_2\text{S} + \text{Aq}$. Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$; very sl. sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$; insol. in $\text{KSH} + \text{Aq}$. (Fresenius.)

Sol. in a mixture of 50 pts. H_2O and 18 pts. HCl (sp. gr. 1.16) even when completely sat. with H_2S . (Lang and Carson, J. Soc. Chem. Ind. 1902, 21. 1018.)

Sl. sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Guerout, C. R. 1872, 75. 1276.)

Cryst. Sb_2S_3 is only sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$ (1 pt. in about 2000 pts. NH_3).

Pptd. amorphous Sb_2S_3 is appreciably more sol. (1 pt. in 600 pts. NH_3). (Garot, J. pr. 1843, 29. 83.)

Sl. sol. in hot 2% $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$, still less sol. in cold. (Materne, C. C. 1906, II. 557.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$.

Sol. in 14–15 pts. pure SbCl_3 . (Schneider, Pogg. 108. 407.)

Slowly sol. in $\text{H}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq}$.

Sol. in boiling $\text{Na}_2\text{SbS}_4 + \text{Aq}$.

Sol. in hot citric, tartaric and oxalic acids. Sl. sol. in malic, benzoic, picric and pyrogalllic acids. Insol. in formic and acetic acids. Especially easily sol. in citric and oxalic acids with addition of KNO_3 , KNO_2 or KClO_3 . (Bolton, C. N. 1878, 37. 86 and 99.)

Sol. in ethylamine sulphhydrate + Aq .

Min. *Stibnite*. Sol. in cold citric acid + Aq . (Bolton, C. N. 37. 14.)

Soluble modification. Sb_2S_3 may be obtained in a colloidal state in aqueous solution containing 1 pt. Sb_2S_3 to 200 pts. H_2O . This can be boiled without decomp., but Sb_2S_3 is pptd. by acids and salts.

Table of maximum dilution of solutions of acids and salts which cause pptn. of Sb_2S_3 .

HCl	1 : 270
H_2SO_4	1 : 140
$\text{H}_2\text{C}_2\text{O}_4$	1 : 45
K_2SO_4	1 : 65
$(\text{NH}_4)_2\text{SO}_4$	1 : 130
MgSO_4	1 : 1720
MnSO_4	1 : 2060
NaCl	1 : 135
BaCl_2	1 : 2050
MgCl_2	1 : 5800
CoCl_2	1 : 2500
KNO_3	1 : 75
Fe_2Cl_6	1 : 2500
$\text{Ba}(\text{NO}_3)_2$	1 : 1250
$\text{K}_2\text{Al}_2(\text{SO}_4)_4$	1 : 35,000
$(\text{NH}_4)_2\text{Fe}_2(\text{SO}_4)_4$	1 : 800
$\text{K}_2\text{Cr}_2(\text{SO}_4)_4$	1 : 40,000
$\text{KSbOC}_4\text{H}_4\text{O}_6$	1 : 18

(Schulze, J. pr. (2) 27. 320.)

Antimony trisulphide with M_2S .

See Sulphantimonites, M.

Antimony pentasulphide, Sb_2S_5 .

Insol. in H_2O , or H_2O containing H_2S . Sol. in conc. $\text{HCl} + \text{Aq}$. Completely sol. in $\text{NH}_4\text{OH} + \text{Aq}$; traces dissolve in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Easily sol. in KOH , or $\text{NaOH} + \text{Aq}$, or in alkali sulphides + Aq . Sol. in 50 pts. cold dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Geiger.)

Insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.

Insol. in cold, but sol. in hot alkali carbonates + Aq . (Berzelius.)

Insol. in $\text{Na}_2\text{SbS}_4 + \text{Aq}$.

When boiled with alcohol, ether, CS_2 , oil of turpentine, etc., portion of the S is dissolved out. (Berzelius.)

CS_2 dissolves about 5% of the sulphur. (Rammelsberg.)

Antimony pentasulphide with M_2S .

See Sulphantimonates, M.

Antimony sulphochloride, SbSCl_3 .

Decomp. by moist air or H_2O . (Cloeze, A. ch. (3) 30. 374.)

$\text{Sb}_2\text{S}_3\text{Cl}$. Easily attacked by acids; insol. in CS_2 . (Ouvrard, C. R. 116. 1516.)

$\text{Sb}_2\text{S}_3\text{Cl}$. (Ouvrard.)

2SbSCl , $3\text{Sb}_2\text{S}_3$. Decomp. by dil. $\text{HCl} + \text{Aq}$. (Schneider.)

SbSCl , 7SbCl_3 . Deliquescent; decomp. by H_2O . (Schneider, Pogg. 108. 407.)

Antimony sulphofluoride, SbF_3S .

See Antimony fluosulphide.

Antimony sulphoiodide, SbSI .

Not attacked by H_2O , and decomp. only by conc. acids. Insol. in CS_2 . (Schneider, Pogg. 110. 147.)

$\text{Sb}_2\text{S}_3\text{I}_4$. (Henry and Garot.)

$\text{Sb}_2\text{S}_3\text{I}_3$. Sol. in dry CS_2 . Very easily decomp. (Ouvrard, C. R. 117. 108.)

Antimony sulphur dioxide, SbSO_2 .

Ppt. (Faktor, C. C. 1900, I. 1211.)

Antimony telluride, SbTe .

Insol. in H_2O .

Sb_2Te_3 . Insol. in H_2O . (Oppenheim, J. pr. 71. 277.)

Antimonyl bromide, SbOBr .

Insol. in CS_2 . (Cooke, Proc. Am. Acad. 13. 104.)

Sl. sol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 826.)

$\text{Sb}_4\text{O}_5\text{Br}_2$. (M'Ivor, C. N. 29. 179.)

$10\text{Sb}_4\text{O}_5\text{Br}_2$, SbBr_3 .

Antimonyl chloride.

From SbCl_3 , SbOCl . Insol. in H_2O . Decomp. by boiling with H_2O ; sol. in $\text{HCl} + \text{Aq}$. Insol. in alcohol or ether; sol. in CS_2 , CHCl_3 , or C_6H_6 . (Sabanajew, Zeit. Ch. 1871. 204.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 826.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

$\text{Sb}_4\text{O}_5\text{Cl}_2$. *Algaroth powder*. Decomp. by H_2O . Sol. in $\text{HCl} + \text{Aq}$ (Cooke, Proc. Am. Acad. 13. 1); tartaric acid + Aq . (Schäffer, A. 152. 135.)

$\text{Sb}_2\text{O}_{11}\text{Cl}_2$. (Cooke.)

$\text{Sb}_2\text{OCl}_{12}$.

$\text{Sb}_{41}\text{O}_{10}\text{Cl}_{22}$.

From SbCl_3 , SbOCl_3 . Deliquescent. Decomposed by H_2O . Sol. in H_2O . (Daubrawa, A. 184. 118.)

Does not exist. (Anschütz and Evans, A. 239. 285.)

$\text{Sb}_2\text{OCl}_{13}$. Deliquescent. Insol. in CS_2 ; easily sol. in tartaric acid + Aq . (Williams, C. N. 24. 224.)

$\text{Sb}_2\text{O}_4\text{Cl}_7$. (Williams.)

SbO_2Cl . Decomp. by hot H_2O into HSbO_3 .

Antimonyl fluoride.

From SbF_3 , $\text{Sb}_4\text{O}_5\text{F}_6$. Not deliquescent. (Flückiger, Pogg. 87. 249.)

Antimonyl caesium fluoride, SbF_4OH , CsF .

(Wells, Am. J. Sci. 1901, (4) 11. 456.)

Antimonyl sodium fluoride, SbOF_3 , $\text{NaF} + \text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O . (Marignac, A. 145. 239.)

Antimonyl iodide, $\text{Sb}_4\text{O}_5\text{I}_2$.

Difficultly sol. in solution of tartaric acid or tartrates. Decomp. by HCl , HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq}$. Easily sol. in alkalies, or $(\text{NH}_4)_2\text{S} + \text{Aq}$.

SbOI . Insol. in CS_2 . (Cooke, Proc. Am. Acad. (2) 5. 72.)

Antimonyl sulphide.

See Antimony oxysulphide.

Argon, A.

100 cc. H_2O dissolve 4.05 cc. argon at 13.9° . Critical t.— 121.6° under 50.6 atmos. Bpt. — 186.9° . Sp. gr. 19.9. (Rayleigh, C. N. 1895, 71. 51–62; 299–302; C. C. 1895. 467.)

Coefficient of absorption in H_2O at $12^\circ = 0.0394$; at $13.9^\circ = 0.0405$. (Ramsay, Phil. Trans. 1895, 186. A. 225.)

Absorption by H_2O at t° .

t°	Coefficient of absorption
0°	0.0561
10	0.0438
20	0.0379
30	0.0348
40	0.0338
50	0.0343

(Antropoff, Roy. Soc. Proc. 1910, 83. A. 480.)

Absorption of argon by H_2O at t° and 760 mm. pressure.

t°	Coefficient of absorption
0°	0.05780
1	0.05612
5	0.05080
10	0.04525
15	0.04099
20	0.03790
25	0.03470
30	0.03256
35	0.03053
40	0.02865
45	0.02731
50	0.02567

(Estreicher, Z. phys. Ch. 1899, 31. 184.)

1 l. H_2O at 38° absorbs 25.7 cc. A.

1 l. blood absorbs 25.3 cc. A. (Regnard and Schloesing, C. R. 1897, 124. 303.)

Not absorbed by members of the fatty series of organic compounds; with members of the aromatic series absorption was observed varying from 8% of the volume employed for benzene to 1% for aniline. (Berthelot, C. R. 1899, 129. 71.)

Arsenamide, $As(NH_2)_3$.

Insol. in liquid NH_3 . Decomp. by H_2O . (Hugot, C. R. 1904, 139. 55.)

Arsenic, As.

Unaltered by pure H_2O . Insol. in $HCl + Aq$ if air is excluded, but sl. sol. in presence of air. Not attacked by dil. $H_2SO_4 + Aq$. Oxidized by conc. H_2SO_4 , HNO_3 , or aqua regia. Not attacked at 20° by HNO_3 , conc. or dil., or containing NO_2 ; nor by $HNO_3 + HCl$, as long as they do not act on each other; but if treated with the above mixture in extremely dilute state, and a few drops of $KNO_3 + Aq$

are added, the As is attacked at once. (Milon, A. ch. (3) 6. 101.)

Sol. in sea water; 0.009 mg. per liter off Brittany; 0.01 to 0.09 mg. per liter near Azores. (Gautier, C. R. 1903, 137. 232.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Insol. in liquid NH_3 . (Hugot, A. ch. 1900, (7) 21. 31.)

Insol. in $NaOH$, KOH , or $NH_4OH + Aq$.

Sol. in S_2Br_2 . (Hannay, Chem. Soc. (2) 11. 823.)

Insol. in alcohol and ether.

Sol. in certain fatty oils.

Insol. in methylene iodide. (Retgers, Z. anorg. 3. 343.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0032 g. As in 6 days. (Gates, J. phys. Ch. 1911, 15. 143.)

Yellow modification. Very unstable. (McLeod, C. N. 1894, 70. 139.)

Fairly stable in liquid air. (Thomson, Chem. Soc. 1906, 90. (2) 745.)

100 ccm. CS_2 dissolve at:

46° 20° 12° 0° -15° -60°
11 8 6 4 2.0–2.5 1.0 g. As.

Less sol. in benzene and ethyl acetate. (Erdmann, Z. anorg. 1902, 32. 448.)

Arsenic acid. See page 59.

Arsenic bromide, $AsBr_3$.

Decomp. by H_2O . Completely sol. in about 3 pts. boiling H_2O , and much less, in presence of HBr . (Wallace, Phil. Mag. (4) 17. 261.)

Sol. in CS_2 .

Sol. in $AlBr_3$. (Isbekow, Z. anorg. 1913, 84. 26.)

Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, 25. 211.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, 25. 217.)

Arsenic caesium bromide, $2AsBr_3, 3CsBr$.

Decomp. by H_2O ; can be recryst. from conc. $HBr + Aq$. (Wheeler, Z. anorg. 4. 451.)

Arsenic rubidium bromide, $2AsBr_3, 3RbCl$.

As the corresponding Cs comp.

Arsenic bromide ammonia, $AsBr_3, 3NH_3$.

Decomp. by H_2O . (Besson, C. R. 110. 1258.)

Arsenic bromide copper, $2AsBr_3, 7Cu$.

Stable toward hot H_2O . Decomp. by KOH . (Hilpert and Herrman, B. 1913, 46. 2224.)

Arsenic bromide silver, $AsBr_3, 3Ag$.

Scarcely decomp. by cold H_2O . (Hilpert and Herrmann.)

Arsenic chloride, $AsCl_3$.

Miscible with little H_2O , and with alcohol, ether, and volatile oils. Decomp. by much H_2O , or by boiling. (Gmelin.)

Miscible with oil of turpentine, and with olive oil. Somewhat sol. in $\text{HCl} + \text{Aq}$.

Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, **25**. 211.)

Sol. in liquid CN . (Centnerszwer, J. russ. phys. Ges. 1901, **38**. 545.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, **25**. 217.)

Arsenic pentachloride, AsCl_5 .

Fumes in the air with evolution of hydrogen chloride. Readily sol. in CS_2 , and absolute ether cooled to -30° . (Baskerville, J. Am. Chem. Soc. 1902, **24**. 1070.)

Arsenic caesium chloride, $2\text{AsCl}_3, 3\text{CsCl}$.

Decomp. by H_2O . 100 pts. $\text{HCl} + \text{Aq}$ (1.2 sp. gr.) dissolve 0.429 pt. salt. (Wheeler, Z. anorg. **4**. 451.)

Arsenic iridium phosphorus chloride.

See Iridium phosphorus chloride arsenic chloride.

Arsenic rubidium chloride, $2\text{AsCl}_3, 3\text{RbCl}$.

Decomp. by H_2O . 100 pts. $\text{HCl} + \text{Aq}$ (sp. gr. 1.2) dissolve 2.935 pts. salt. (Wheeler, Z. anorg. **4**. 451.)

Arsenic sulphur chloride, $2\text{AsCl}_3, 3\text{SCl}_2$.

Decomp. by H_2O . (Rose.)

Above compound is a mixture. (Nilson, C. N. **81**. 81.)

Arsenic chloride ammonia, $2\text{AsCl}_3, 7\text{NH}_3$.

Decomp. by cold H_2O , with evolution of NH_3 . From the solution crystallizes $\text{As}_4\text{Cl}_2, \text{N}_2\text{H}_{10}\text{O}_3$.

Sol. in alcohol without decomp. (Rose, Pogg. **52**. 62.)

Composition is $\text{AsCl}_3, 4\text{NH}_3$. (Besson, C. R. **110**. 1258.)

Arsenic chloride copper, $2\text{AsCl}_3, 7\text{Cu}$.

Somewhat decomp. by H_2O . Decomp. by KOH , or hot HCl . (Hilpert and Herrman, B. 1913, **46**. 2224.)

Arsenic chloride silver, $2\text{AsCl}_3, 7\text{Ag}$.

H_2O , NH_4OH and KOH split off Ag . (Hilpert and Herrmann.)

Arsenic trifluoride, AsF_3 .

Sol. in H_2O with evolution of heat and decomposition. (Berzelius.)

Easily sol. in benzene. (Moissan, C. R. **99**. 874.)

Miscible with alcohol and ether. (M'Ivor, C. N. **30**. 169.)

Arsenic pentafluoride, AsF_5 .

Sol. in H_2O , alkalis + Aq and liquid AsF_3 with evolution of heat. Absorbed by ether, alcohol and benzene with evolution of heat. (Ruff, B. 1906, **39**. 67.)

Arsenic potassium fluoride, $\text{AsF}_3, \text{KF} + \frac{1}{2}\text{H}_2\text{O}$.

$\text{AsF}_3, 2\text{KF} + \text{H}_2\text{O}$.

$\text{AsF}_3, \text{AsOF}_3, 4\text{KF} + 3\text{H}_2\text{O}$. (Marignac, A. **145**. 237.)

Arsenic fluoride ammonia, $2\text{AsF}_3, 5\text{NH}_3$.

Easily decomp. by H_2O . (Besson, C. R. **110**. 1258.)

Arsenic pentafluoride nitrosyl fluoride, AsF_5, NOF .

Decomp. by H_2O , fuming HCl , $\text{NaOH} + \text{Aq}$, dry ether and dry alcohol with evolution of NO . Sol. in conc. HNO_3 , hot conc. H_2SO_4 , boiling NOCl and AsF_3 . Insol. in CCl_4 and CS_2 . (Ruff, Z. anorg. 1908, **58**. 327.)

Arsenic trifluoride sulphur tetrachloride, $2\text{AsF}_3, \text{SCl}_4$.

Very hygroscopic. Decomp. by H_2O and NaOH . Decomp. by thionyl chloride, CCl_4 , CS_2 , abs. alcohol and ether. Decomp. by ligroin, benzene and toluene. (Ruff, B. 1904, **37**. 4520.)

Arsenic hydride, AsH_3 .

Sl. sol. in H_2O and alkali hydrates + Aq , with subsequent decomposition. H_2O absorbs $\frac{1}{2}$ vol. AsH_3 . Decomp. by conc. acids. Absorbed rapidly by oil of turpentine, slightly by fixed oils, and not at all by alcohol, ether, or $\text{KOH} + \text{Aq}$. (Gmelin.)

Insol. in $\text{KOH} + \text{Alcohol}$. (Meissner.)

Not more sol. in alkaline solutions than in pure H_2O . (Berzelius.)

AsH_3 . *Solid*. Insol. in H_2O , alcohol, ether, and CS_2 . (Wiederhold, Pogg. **118**. 615.)

Insol. in H_2O ; sol. in methylene iodide, xylene, or in conc. $\text{KOH} + \text{Aq}$. (Retgers, Z. anorg. **4**. 403.)

Arsenic hydride boron bromide, $\text{AsH}_3, \text{BBr}_3$.

Easily decomp. Decomp. by H_2O . Appreciably sol. in AsH_3 or BBr_3 . Insol. in CS_2 . (Stock, B. 1901, **34**. 949.)

Arsenic diiodide, As_2I_4 .

Decomp. by H_2O or alkalis; easily sol. in alcohol, ether, chloroform, or carbon disulphide. (Bamberger and Phillip, B. **14**. 2643.)

Not attacked by cold conc. H_2SO_4 , or by cold fuming HNO_3 . The latter oxidizes on warming. Decomp. by pyridine. Sol. in boiling acetic anhydride. (Hewitt and Winmill, Chem. Soc. 1907, **91**. 962.)

Arsenic triiodide, AsI_3 .

Sol. in 3.32 pts. boiling H_2O , and solution if boiled down deposits pure AsI_3 , but if left to cool slowly, deposits crystals of As_2O_3 and AsOI .

Sl. sol. in $\text{HCl} + \text{Aq}$.

Sol. in POCl_3 , PCl_3 , and PBr_3 . (Walden, Z. anorg. 1900, **25**. 212.)

Sol. in PCl_3 . (Beckmann, Z. anorg. 1906, 11. 110.)

Sol. in SOCl_2 , S_2Cl_2 and SO_2Cl_2 . (Walden, Z. anorg. 1900, 25. 216.)

Sol. in SnCl_4 . (Walden, l.c.)

Easily sol. in AsBr_3 . (Walden, Z. anorg. 1902, 29. 374.)

Sol. in AsCl_3 . (Walden, Z. anorg. 1900, 25. 214.)

Sol. in alcohol without decomp.

Sol. in ether, benzene, chloroform, and CS_2 .

100 pts. methylene iodide dissolve 17.4 pts. AsI_3 at 12° . (Retgers, Z. anorg. 3. 343.)

Arsenic pentaiodide, AsI_5 .

More or less sol. in H_2O , alcohol, CHCl_3 , ether and CS_2 . (Sloan, C. N. 1882, 46. 194.)

Arsenic caesium iodide, $2\text{AsI}_3 \cdot 3\text{CsI}$.

Decomp. by H_2O ; sol. in conc. $\text{HI} + \text{Aq}$. (Wheeler, Z. anorg. 4. 451.)

Arsenic rubidium iodide, $2\text{AsI}_3 \cdot 3\text{RbI}$.

As the corresponding Cs comp.

Arsenic sulphur iodide.

See Arsenic sulphoiodide.

Arsenic triiodide ammonia, $2\text{AsI}_3 \cdot 9\text{NH}_3$.

Insol. in benzene. (Bamberger and Phillip, B. 14. 2643.)

$\text{AsI}_3 \cdot 4\text{NH}_3$. (Besson, C. R. 110. 1258.)

Arsenic nitride, AsN .

Easily decomp. into As and N. (Hugot, C. R. 1904, 139. 56.)

Decomp. by heat. (Frans Fischer, B. 1910, 43. 1471.)

Arsenic suboxide, As_2O (?).

Insol. in H_2O ; decomp. by dil. acids or $\text{NH}_4\text{OH} + \text{Aq}$.

Does not exist. (Geuther, A. 240. 208.)

Arsenic trioxide, As_2O_3 .

"White arsenic" exists in two modifications: $\alpha\text{As}_2\text{O}_3$,—crystalline, octahedral, opaque, porcelainous, etc.; $\beta\text{As}_2\text{O}_3$,—amorphous, vitreous, "arsenic glass."

The data concerning the solubility of As_2O_3 are very contradictory, the reasons being that (1) the solubility of the two modifications is different; (2) that the length of time necessary to effect solution differs in the two modifications; and (3) that there is a tendency of the amorphous As_2O_3 to go over into the crystalline state during the process of solution. $\alpha\text{As}_2\text{O}_3$ is also not easily moistened, especially when in a pulverulent condition, which is not the case with the β modification. (Winkler, J. pr. (2) 31. 247.)

The older data are very unreliable, but possess a certain historical interest.

1 pt. As_2O_3 is sol. in 10.55 pts. (Wenzel); 11.34 pts. (Fischer); 11.86 pts. in $\frac{1}{2}$ hour (Klaproth); 12.2 pts. (Bucholz); 15.0 pts. (Brandt; Bergman); 16.0 pts. (Vogel); 24 pts. (Lametherie); 40 pts. (Pörner); 64 pts.

(Baumé); 80 pts. (Navier); 200 pts. (Aschof and Name, 1812); 640 pts. (Hagen, 1798) boiling H_2O .

1 pt. As_2O_3 is sol. in 7.72 pts. H_2O if α , or 9.33 pts. if β (Guibort); in 24 pts. H_2O if α , or 31 pts. if β (Taylor).

Sol. in 53.3 pts. H_2O at 18.75° . (Abl.)

Sol. in 30 pts. H_2O . (Nussebrook.)

After the solution in H_2O at 100° has been left standing at ordinary temperatures—

1 pt. As_2O_3 remains dissolved in 16 pts. H_2O at 16° , and 30 pts. H_2O at 7° (Bucholz); in 33 pts. H_2O at 7° (Klaproth); in 38.45 pts. H_2O after 3 days, 55 pts. H_2O after 8 days, 64.50 pts. H_2O after 2.3 weeks at 10° (Fischer); in 38.52 pts. if $\alpha\text{As}_2\text{O}_3$ was used, 55.06 pts. if $\beta\text{As}_2\text{O}_3$ was used (Guibort); in 38 pts. if $\alpha\text{As}_2\text{O}_3$ after 6 months, 53.71 pts. if $\beta\text{As}_2\text{O}_3$ after 48 hours (Taylor).

When an excess of pulverized As_2O_3 is left to digest for several days with cold H_2O —

1 pt. dissolves in 50 pts. (Bucholz), in 66 pts. (Fischer); in 80 pts. at 15° (Bergman); in 80 pts. if α , and 103 pts. if β (Guibort); 95 pts. at 10° (Spelman); 96 pts. at 35.5° (Hahnemann); 320 pts. H_2O at 20° (Aschof and Name, 1812.)

H_2O at 15.5° or below dissolves less than $\frac{1}{4}\%$ As_2O_3 . (Dalton.)

To dissolve 1 pt. As_2O_3 in 12 pts. H_2O , it is necessary to boil an excess of As_2O_3 with H_2O ; if 1 pt. As_2O_3 is boiled with 12 pts. H_2O , considerable remains undissolved, and even with 1 pt. As_2O_3 to 50–60 pts. H_2O long continued boiling is necessary to effect solution. If a clear solution saturated by long boiling with an excess of As_2O_3 is poured off and evaporated continuously to $\frac{1}{2}$ its original bulk, no As_2O_3 separates out, and the solution contains 1 pt. As_2O_3 to 6 pts. H_2O . (Fischer.)

100 pts. aqueous solution of $\beta\text{As}_2\text{O}_3$ sat. at 15° contain 0.96 pt. As_2O_3 , and 9.68 pts. when sat. at 100° . (Guibort.)

If 1 pt. pulverized As_2O_3 be digested 10 days at $19\text{--}25^\circ$ in 3–10 pts. H_2O , the solution contains 1 pt. As_2O_3 to 50 pts. H_2O . A solution of same strength is obtained in 25 days by digesting 1 pt. As_2O_3 in 40 pts. H_2O . If 1 pt. As_2O_3 be immersed in 80 pts. H_2O , the resulting solution contains 1 pt. As_2O_3 to 90 pts. H_2O ; if in 160 pts. H_2O , 1 pt. As_2O_3 to 180 pts. H_2O ; if in 240 pts. H_2O , 1 pt. As_2O_3 to 280 pts. H_2O ; if in 1000 pts. H_2O , 1 pt. As_2O_3 to 1200 pts. H_2O ; and even when 1 pt. As_2O_3 is digested at ordinary temperatures for several days with 16,000–100,000 pts. H_2O , a portion remains undissolved. Pulverized $\alpha\text{As}_2\text{O}_3$ was set aside with H_2O in closed bottles for 18 years; when 1 pt. As_2O_3 was present in 1000 pts. H_2O , a perfect solution was obtained; when 1 pt. As_2O_3 in 100 pts. H_2O , 0.017% As_2O_3 was undissolved, when 1 pt. As_2O_3 in 35 pts. H_2O , 0.35% As_2O_3 was undissolved, so that the solution contained 1 pt. As_2O_3 to 54 pts. H_2O . (Gmelin.)

Porcelainous modification ($\alpha\text{As}_2\text{O}_3$) is much more sol. in H_2O than the vitreous ($\beta\text{As}_2\text{O}_3$). 100 pts. H_2O at ordinary temperature dissolve 0.96 pt. $\beta\text{As}_2\text{O}_3$ and 1.25 pts. $\alpha\text{As}_2\text{O}_3$; 100 pts. boiling H_2O dissolve 9.68 pts. $\beta\text{As}_2\text{O}_3$ and 11.47 pts. $\alpha\text{As}_2\text{O}_3$; and when the temperature of this solution has fallen to 15° , the solution from $\beta\text{As}_2\text{O}_3$ retains 1.78 pts., and that from $\alpha\text{As}_2\text{O}_3$ retains 2.9 pts. (Berselius [citing Guibort].)

$\beta\text{As}_2\text{O}_3$ dissolves more quickly and abundantly than $\alpha\text{As}_2\text{O}_3$. The same amount H_2O which will take up 36–38 pts. $\beta\text{As}_2\text{O}_3$ at $12\text{--}13^\circ$ will dissolve only 12–14 pts. $\alpha\text{As}_2\text{O}_3$, or 100 pts. H_2O dissolve 4 pts. $\beta\text{As}_2\text{O}_3$ and 1.2–1.3 pts. $\alpha\text{As}_2\text{O}_3$. By long boiling with H_2O , $\alpha\text{As}_2\text{O}_3$ is converted into $\beta\text{As}_2\text{O}_3$, and thus acquires the solubility of the latter, so that 100 pts. boiling H_2O can take up 11 pts. As_2O_3 . But at low temperature $\beta\text{As}_2\text{O}_3$ is converted into $\alpha\text{As}_2\text{O}_3$ when in contact with H_2O , so that the solution becomes weaker after a while, and retains only the proportion of As_2O_3 corresponding to the solubility of $\alpha\text{As}_2\text{O}_3$. Comminution, which hastens the rate of solubility of $\alpha\text{As}_2\text{O}_3$ without increasing the amount dissolved, diminishes the solubility of $\beta\text{As}_2\text{O}_3$, as this is converted into $\alpha\text{As}_2\text{O}_3$.

by the friction or contact with H_2O . As_2O_3 , which has been rendered opaque by NH_4OH , and that which has been crystallized from an aqueous solution, are equally sol. in H_2O . (Bussy, C. R. 24. 774; A. 64. 286.)

100 pts. H_2O dissolve 1.707 pts. βAs_2O_3 in $2\frac{1}{2}$ years; 100 pts. boiling H_2O dissolve 11.46 pts. βAs_2O_3 in 3 hours, and 11.86 pts. in 12 hours; 10.14 pts. αAs_2O_3 in 3 hours, and 10.18 pts. in 12 hours. (Rose, Ann. Phys. (1) 36. 494.)

A cold sat. solution which stood over excess of As_2O_3 for 10 months at $10-20^\circ$ contains 1.2% As_2O_3 ; hot sat. solution a few days after saturation contains 2.25-2.50% As_2O_3 . If trace of HCl is present, the solution contains 3.8% As_2O_3 . Hot sat. solution of porcelain mod. of As_2O_3 contains 4 days after saturation 2.4% As_2O_3 at 24° ; after 82 days at 14° , 1.5%; after 4 months at 12° , 1.3% As_2O_3 . (Bacaloglo, J. pr. 83. 111.)

According to later experiments, 1 pt. αAs_2O_3 dissolves in 355 pts. H_2O in 1 day at 15° , while 1 pt. βAs_2O_3 dissolves in 108 pts. H_2O under the same conditions. 1 pt. αAs_2O_3 dissolves in 46 pts. H_2O , if solution is prepared at 100° , and allowed to stand 24 hours at 15° , while 1 pt. βAs_2O_3 dissolves in 30 pts. H_2O under the same conditions. (Büchner, N. Rep. Pharm. 22. 265.)

100 pts. H_2O dissolve pts. αAs_2O_3 and βAs_2O_3 at ordinary temperature:

Time	αAs_2O_3	βAs_2O_3
1 hour	0.023	1.589
3 hours	0.088	2.356
6 hours	0.353	3.666
12 hours	0.364	3.361
24 hours	0.956	3.306
2 days	1.627	2.629
4 days	1.814	2.429
1 week	1.673	1.763
3 weeks	1.776	1.713
$2\frac{1}{4}$ years	1.712	1.707

In the solution of βAs_2O_3 , octahedral crystals were deposited on the sides of the vessel after 12 hours, which continued to increase. There was no such deposit in the case of αAs_2O_3 .

From the maxima in the above table, 100 pts. H_2O can dissolve 3.7 pts. βAs_2O_3 and 1.7 pts. αAs_2O_3 at ordinary temperature.

100 pts. boiling H_2O dissolve 11.46 pts. βAs_2O_3 and 10.140 pts. αAs_2O_3 in 3 hours; 11.86 pts. βAs_2O_3 and 10.176 pts. αAs_2O_3 in 12 hours. (Cl. Winkler, J. pr. (2) 31. 247.)

100 pts. H_2O dissolve 1.75 pts. of a third modification (hexagonal crystalline) at ordinary temperature, and 2.75 pts. at 100° . (Claudet, Chem. Soc. (2) 6. 179.)

βAs_2O_3 dissolves more rapidly in $HCl + Aq$ than αAs_2O_3 . (Schultz-Sellac, B. 4. 109.)

While 100 ccm. H_2O dissolve 0.8507 g. βAs_2O_3 at 18.5° , 100 ccm. H_2O containing 1.3195 g. HCl dissolve 1.1513 g. βAs_2O_3 ; containing 6.09 g. HCl , 1.2724 g. βAs_2O_3 . (Chodounsky, Listy Chemické, 13. 114.)

100 ccm. H_2O dissolve 1.495 g. As_2O_3 at 15° . (Wood, Chem. Soc. 1908, 93. 412.)

Solubility of crystalline As_2O_3 in H_2O .
1 l. of the sat. solution contains at:

2°	15°	25°	39.8°	bpt.
12.006	16.566	20.384	29.302	60 + g. As_2O_3

(Bruner, Z. anorg. 1903, 37. 456.)

Much more easily sol. in many acids than in H_2O . Easily sol. in fuming H_2SO_4 . (Schultz-Sellac.)

100 pts. dilute $H_2SO_4 + Aq$ of various strengths dissolve at t° .

t°	Pts. βAs_2O_3	t°	Pts. βAs_2O_3	Ratios of amts. dissolved at $80^\circ : 18.5^\circ$
80°	1.0195	18.5°	0.5422	1.88 : 1
..	1.3664	0.7203	1.89 : 1
..	1.1933	0.6522	1.84 : 1

(Chodounsky, l.c.)
Decomp. by HNO_3 or aqua regia into As_2O_5 .
Sol. in $H_3PO_4 + Aq$. (Bergman.)
More sol. in $HCl + Aq$ than in H_2SO_4 , or $HNO_3 + Aq$, and still less in $HC_2H_3O_2 + Aq$.
Solubility in $HCl + Aq$.

Conc. of $HCl + Aq$	Grams of As_2O_3 per 100 cc. of solution
0.46N	1.52
0.98N	1.41
2.03N	1.17
3.13N	1.11
3.81N	1.13
5.32N	2.20
6.50N	5.11
7.85N	12.28
9.17N	18.16

As the concentration of the acid increases, the solubility of the oxide decreases, a minimum being reached when the concentration of the solvent is about 3.2N. Beyond this point, an increase in the concentration of the solvent leads to a corresponding increase in the solubility. (Wood, Chem. Soc. 1908, 93. 413.)

Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54. 674.)

Easily sol. in cold $H_2C_2O_4 + Aq$. (Bergman.)

When pulverized, it dissolves in hot $H_2C_2O_4 + Aq$, but separates out on cooling.

Easily sol. in hot benzoic acid + Aq .

Sol. in tartaric acid + Aq .

Easily sol. in alkali hydrates, or carbonates + Aq .

Easily sol. in NH_3 arsenite + Aq at 70–80°, crystallizing out on cooling. (Berzelius.)

Sol. in hot $\text{K}_2\text{C}_2\text{O}_4$ + Aq.

Sol. in AsCl_3 . (Penney and Wallace.)

More sol. in $\text{Na}_2\text{B}_4\text{O}_7$ + Aq than in H_2O .

Very al. sol. in absolute alcohol. (Vogel.)

Fol. in 80 pts. highly rectified spirit. (Wenzel.)

When 1 pt. powdered As_2O_3 is digested 30 days in 10–40 pts. alcohol, a solution is formed containing 1 pt. As_2O_3 to 60 pts. alcohol, when 1 pt. As_2O_3 is digested with 60–150 pts. alcohol, a solution is formed containing 1 pt. As_2O_3 to 124–140 pts. alcohol. (Fischer.)

Sol. in 70–80 pts. alcohol. (Thompson.)

Alcohol dissolves 0.446 pt. $\beta\text{As}_2\text{O}_3$. (Rose, A. Phys. (1) 52. 455.)

100 pts. alcohol dissolve pts. As_2O_3 :

Vol. % of alcohol	$\alpha\text{As}_2\text{O}_3$ at 15°	$\alpha\text{As}_2\text{O}_3$ at b.-pt. of alcohol	$\beta\text{As}_2\text{O}_3$ at 15°
56	1 680	4 895	0.504
79	1.430	4.551	0.540
84			0.565
86	0.715	3.197	
88			0.717
100	0.025	3.402	1.060

(Girardin, J. Pharm. (3) 46. 269.)

100 pts. absolute alcohol dissolve 0.446 pt. $\beta\text{As}_2\text{O}_3$ in 2½ years. (Winkler, J. pr. (2) 31. 347.)

Nearly insol. in ether.

100 pts. ether dissolve 0.454 pt. $\beta\text{As}_2\text{O}_3$. (Winkler.)

Ether extracts 1 mg. As_2O_3 from sat. As_2O_3 + Aq for every 15 cc. ether used; less is extracted when the solution is acidified with HCl , and almost none if acidified with H_2SO_4 or $\text{H}_3\text{C}_4\text{O}_4$. (Selmi, B. 13. 206.)

$\alpha\text{As}_2\text{O}_3$ is sol. in 50 pts. boiling nitrobenzol. $\beta\text{As}_2\text{O}_3$ is insol. in boiling nitrobenzol. (Auerbach, Z. anorg. 1903, 37. 353.)

$\beta\text{As}_2\text{O}_3$ dissolves in oil of turpentine, but $\alpha\text{As}_2\text{O}_3$ is insol. therean. $\alpha\text{As}_2\text{O}_3$ is very al. sol. in benzene or petroleum ether, but more sol. in methyl alcohol, ethyl alcohol, ether, or chloroform. (Selmi.)

100 pts. CS_2 dissolve 0.001 pt. $\beta\text{As}_2\text{O}_3$ in 2½ years. (Winkler.)

Sl. sol. in the fatty oils.

1000 pts. castor-oil dissolve 1.33 pts. As_2O_3 at ordinary temperature, and 9 pts. at boiling temperature. 1000 pts. other oils dissolve 0.6–0.8 pt. As_2O_3 in the cold, and about 1.7 pts. on boiling. (Berzelius.)

Insol. in chinoline or aniline. (Hoffmann, A. ch. (3) 9. 143, 169.)

Moderately sol. in^o chinolin. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329); (Eidmann, C. C. 1899, II 1014.)

Sol. in amyl alcohol and is divided between it and H_2O in the constant ratio of 1 : 5.47 at 25°. (Auerbach, Z. anorg. 1903, 37. 376.)

Min. *Arsenolite*.

Arsenic trioxide pentoxide, $3\text{As}_2\text{O}_3$, $2\text{As}_2\text{O}_5$ + $3\text{H}_2\text{O}$.

Decomp. by H_2O . (Joly, C. R. 100. 1221.)

$2\text{As}_2\text{O}_5$, As_2O_3 + H_2O . Decomp. by H_2O . (Joly.)

As_2O_3 , As_2O_5 + H_2O . (Joly.)

Arsenic tetroxide, As_2O_4 .

Sl. sol. in H_2O from which it is partially pptd. by alcohol. More easily sol. in alkali carbonates or HCl + Aq. Most easily sol. in NaOH or KOH + Aq. (Herbst, Dissert. 1894.)

Arsenic pentoxide, As_2O_5 .

Deliquescent in moist air; slowly sol. in H_2O , forming H_2AsO_4 , which see. Easily sol. in alcohol; much more sol. in alcohol than As_2O_3 . Very al. sol. in the fatty oils, 100 pts. of oil dissolving 0.2 pt. As_2O_5 in the cold, and 1 pt. with partial decomp. on boiling. (Berzelius.)

1000 pts. boiling poppy-oil dissolve 27 pts. As_2O_5 ; 1000 pts. boiling castor-oil dissolve 34 pts. As_2O_5 . (Heimpel and Grundner.)

+ $4\text{H}_2\text{O}$ Solubility in H_2O at t° .

t°	Pts. H_2AsO_4 in 100 pts. solution	t°	Pts. H_2AsO_4 in 100 pts. solution
-55°	69.9	-5°	80.0
50	70.9	0	81.0
-45	71.9	+5	82.1
-40	72.9	10	83.3
-35	73.9	15	84.7
-30	74.9	20	86.3
-25	75.9	25	88.0
-20	76.9	30	90.1
-15	77.9	35	92.8
-10	78.9		

(Menzies and Potter, J. Am. Chem. Soc. 1912, 34. 1464.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Solubility in H_2O at t° .

t°	Pts. H_2AsO_4 in 100 pts. of solution
+10°	88.4
20	89.1
30	89.8
40	90.5
50	91.2
60	91.9
70	92.6
80	93.2
90	93.8
100	94.4
110	95.0
120	95.6
130	96.2
140	96.8

Menzies and Potter, J. Am. Chem. Soc. 1912, 34. 1464.)

$\text{As}_2\text{O}_3 + 4\text{H}_2\text{O}$ and $3\text{As}_2\text{O}_3 + 5\text{H}_2\text{O}$ are the only hydrates that can be isolated. (Menzies and Potter.)

See also Arsenic Acid.

Arsenic trioxide, with alkali haloid.

See Arsenite, alkali haloid.

Arsenic sulphur trioxide, $\text{As}_2\text{O}_3, \text{SO}_3$.

Deliquescent; decomp. by H_2O . (Adie, Chem. Soc. 55. 157.)

$\text{As}_2\text{O}_3, 2\text{SO}_3$. As above. (Adie.)

$\text{As}_2\text{O}_3, 3\text{SO}_3$. (Weber, B. 19. 3186.)

$\text{As}_2\text{O}_3, 4\text{SO}_3$. As above. (Adie.)

$\text{As}_2\text{O}_3, 6\text{SO}_3$. (Weber.)

$\text{As}_2\text{O}_3, 8\text{SO}_3$. As above. (Adie.)

Arsenic oxychloride, etc.

See Arsenyl chloride, etc.

Arsenic phosphide, AsP .

Decomp. by H_2O . Not attacked by cold H_2SO_4 or HCl , and only sl. sol. therein on warming. Easily decomp. by HNO_3 , KOH , NaOH , $\text{BaO}_2\text{H}_2 + \text{Aq}$. Insol. in alcohol, ether, chloroform; sl. sol. in CS_2 .

$\text{P}_2\text{As}_2\text{O}_5$. Product of action of H_2O on above compound, which it resembles. (Janowsky, B. 6. 216.)

Arsenic monoselenide, As_2Se .

Insol. in most organic and inorganic solvents. Sol. very slowly in conc. HCl and H_2SO_4 . Sol. in boiling alkali hydroxides + Aq . (Szarvasy, B. 1897, 30. 1245.)

Arsenic triselenide, As_2S_3 .

Partially sol. in $\text{KOH} + \text{Aq}$ if boiled with it for a long time. (Uelsmann, A. 116. 123.)

Arsenic pentaselenide, As_2Se_5 .

Insol. in most solvents, as conc. HCl . Sol. in alkali hydrates and sulpho-hydrates + Aq . (Szarvasy, B. 1895, 28. 2655–2656.)

Insol. in H_2O , in dil. acids and in conc. HCl . Sl. sol. in warm $\text{HNO}_3 + \text{Aq}$. Oxidized by cold fuming HNO_3 . Sol. in alkalies and in hot alkali carbonates + Aq . Insol. in alcohol, ether, CS_2 , etc. (Clever, Z. anorg. 1895, 10. 129.)

Arsenic selenosulphide.

See Arsenic sulphoselenide.

Arsenic sulphide, As_2S_3 .

Ppt. Insol. in NH_4OH or in colorless $(\text{NH}_4)_2\text{S} + \text{Aq}$. Sol. in yellow $\text{NH}_4\text{SH} + \text{Aq}$. (Scott, Chem. Soc. 1900, 77. 652.)

Arsenic disulphide, As_2S_2 .

Min. *Realgar*. Difficultly sol. in alkali sulphides + Aq . Partly dissolved by $\text{KOH} + \text{Aq}$ with decomposition. Sol. at 150° in a sealed tube in $\text{NaHCO}_3 + \text{Aq}$, and crystallizes out on cooling. (Senarmont, A. ch. (3) 32. 158.)

Arsenic trisulphide, As_2S_3 .

Insol. in H_2O when prepared in the dry way, but when prepared moist is very liable to go into the colloidal modification mentioned below. Insol. in H_2O containing H_2SO_4 , HNO_3 , HCl , $\text{H}_2\text{C}_2\text{O}_4$, $\text{HC}_2\text{H}_3\text{O}_2$, $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$, CO_2 , NH_4Cl , KNO_3 , $(\text{NH}_4)_2\text{SO}_4$, MgSO_4 . (Bontigny.)

Insol. in H_2O . Traces are dissolved by $\text{H}_2\text{S} + \text{Aq}$. Sl. decomp. by boiling with H_2O , or long contact with cold H_2O . (Fresenius.)

1 l. H_2O dissolves 2.1×10^{-6} mols. pptd. As_2S_3 at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

Insol. in dil. acids. Insol. in cold, and scarcely attacked by hot conc. $\text{HCl} + \text{Aq}$.

Easily decomp. by HNO_3 or aqua regia.

Easily sol. in cold KOH , NaOH , or $\text{NH}_4\text{OH} + \text{Aq}$, also in alkali carbonates, or sulphates + Aq .

Sol. in hot $\text{KHSO}_3 + \text{Aq}$.

Sol. in citric acid, and alkali citrates + Aq . (Spiller.)

Slowly sol. in cold 2% $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$. Easily sol. on heating. (Materne, C. C. 1906, II. 557.)

Insol. in CS_2 .

Min. *Orpiment*.

As_2S_3 may also be obtained in a colloidal form, sol. in H_2O . Sat. solution contains 34.46% As_2S_3 ; it is decomp. by standing, but may be boiled without undergoing decomposition; most acids and many salts ppt. As_2S_3 . (Schulze, J. pr. (2) 25. 431.)

The following solutions cause pptn. of As_2S_3 in a solution of the colloidal modification, when added in the given state of dilution:—

$\text{HCl} + \text{Aq}$.	.	.	1 : 555
$\text{HNO}_3 + \text{Aq}$.	.	.	1 : 276
$\text{H}_2\text{SO}_4 + \text{Aq}$.	.	.	1 : 255
$\text{H}_2\text{SO}_3 + \text{Aq}$.	.	.	1 : 138
$\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$.	.	.	1 : 65
$\text{H}_3\text{PO}_4 + \text{Aq}$.	.	.	1 : 26
$\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.	.	.	1 : 0.18
$\text{K}_2\text{SO}_4 + \text{Aq}$.	.	.	1 : 76
$\text{Na}_2\text{SO}_4 + \text{Aq}$.	.	.	1 : 129
$(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$.	.	.	1 : 188
$\text{CaSO}_4 + \text{Aq}$.	.	.	1 : 2780
$\text{MgSO}_4 + \text{Aq}$.	.	.	1 : 2630
$\text{ZnSO}_4 + \text{Aq}$.	.	.	1 : 3330
$\text{MnSO}_4 + \text{Aq}$.	.	.	1 : 2860
$\text{NiSO}_4 + \text{Aq}$.	.	.	1 : 3440
$\text{FeSO}_4 + \text{Aq}$.	.	.	1 : 2380
$\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$.	.	.	1 : 52600
$\text{Ti}_2\text{SO}_4 + \text{Aq}$.	.	.	1 : 799
$\text{KCl} + \text{Aq}$.	.	.	1 : 137
$\text{KBr} + \text{Aq}$.	.	.	1 : 103
$\text{KI} + \text{Aq}$.	.	.	1 : 55
$\text{LiI} + \text{Aq}$.	.	.	1 : 127
$\text{NaCl} + \text{Aq}$.	.	.	1 : 212
$\text{NH}_4\text{Cl} + \text{Aq}$.	.	.	1 : 207
$\text{BaCl}_2 + \text{Aq}$.	.	.	1 : 2860
$\text{CaCl}_2 + \text{Aq}$.	.	.	1 : 4370
$\text{MgCl}_2 + \text{Aq}$.	.	.	1 : 10000

$\text{FeCl}_3 + \text{Aq}$	1 : 50000
$\text{AlCl}_3 + \text{Aq}$	1 : 83000
$\text{CrCl}_3 + \text{Aq}$	1 : 20000
$\text{KNO}_3 + \text{Aq}$	1 : 84
$\text{NaNO}_3 + \text{Aq}$	1 : 117
$\text{NH}_4\text{NO}_3 + \text{Aq}$	1 : 138
$\text{Ba}(\text{NO}_3)_2 + \text{Aq}$	1 : 2080
$\text{KClO}_3 + \text{Aq}$	1 : 88
$\text{CaH}_2(\text{CO}_3)_2 + \text{Aq}$	1 : 3120
$\text{K}_2\text{C}_2\text{H}_3\text{O}_4 + \text{Aq}$	1 : 85
$\text{K}_2\text{C}_2\text{O}_4 + \text{Aq}$	1 : 81
$\text{NaC}_2\text{H}_3\text{O}_4 + \text{Aq}$	1 : 78
$\text{Urea} + \text{Aq}$	1 : 25
$(\text{NH}_4)_2\text{Fe}(\text{SO}_4)_2 + \text{Aq}$	1 : 1160
$\text{K}_2\text{Al}_2(\text{SO}_4)_4 + \text{Aq}$	1 : 50000
$\text{K}_2\text{Fe}_2(\text{SO}_4)_4 + \text{Aq}$	1 : 55500
$\text{K}_2\text{Cr}_2(\text{SO}_4)_4 + \text{Aq}$	1 : 25000
$\text{K}_4\text{Fe}(\text{CN})_6 + \text{Aq}$	1 : 67
$\text{K}_4\text{Fe}(\text{CN})_6 + \text{Aq}$	1 : 81

Cold conc. solutions of boric, arsenious, tartaric, benzoic, and salicylic acids, also cane sugar, or chloral hydrate cause no pptn. Absolute alcohol and glycerine may also be mixed with the solutions without causing pptn. (Schulze, J. pr. (2) 25. 442.)

+6H₂O; decomp. completely into As₂S₃ under a pressure of 6000 to 7000 atmos. (Spring, Z. anorg. 1895, 10. 186.)

Arsenic pentasulphide, As₂S₅.

Insol. in H₂O. Sol. in NH₄OH, KOH, NaOH + Aq, and solutions of alkali sulphides and carbonates. Sol. in BaO₂H₂ and CaO₂H₂ + Aq.

Sol. in citric acid, and alkali citrates + Aq. (Spiller.)

Alcohol dissolves out S on boiling. (Berzelius.)

Sol. in alkali arsenates + Aq. (Nilson, J. pr. (2) 14. 155.)

+H₂O. (Nilson, l.c.)

Arsenic trisulphide, with M₂S.

See Sulpharsenites, M.

Arsenic pentasulphide, with M₂S.

See Sulpharsenates, M.

Arsenic sulphobromide, AsS₂Br₃ = AsSBr + SBr₂.

Decomp. by H₂O. (Hannay, Chem. Soc. 23. 284.)

Arsenic sulphochloride, As₂S₂Cl.

Slowly decomp. by boiling H₂O. Sol. in hot AsCl₃ without decomp. (Ouvrard, C. R. 116. 1516.)

AsS₂Cl. Decomp. by H₂O. Sol. in NH₄OH, and alkali carbonates + Aq. (Ouvrard.)

AsS₂Cl. Slowly decomp. by boiling H₂O. Sol. in alkali carbonates and in NH₄OH + Aq. (Ouvrard, C. R. 1893, 116. 1517.)

Arsenic sulphiodide, AsSI.

Insol. in alcohol, chloroform or carbon disulphide. (Schneider, J. pr. (2) 23. 486.)

Formula is probably As₂S₂, AsI₂.

Slowly attacked by HCl + Aq; somewhat more easily by HNO₃ + Aq. Easily sol. in KOH, or NH₄OH + Aq. (Schneider, J. pr. (2) 24. 505.)

2AsI₂, SI₂. Decomp. on air. (Schneider, J. pr. (2) 26. 509.)

As₂S₂I₂. Less sol. in CS₂ than AsI₂. (Ouvrard, C. R. 117. 107.)

As₂SI₂. (Ouvrard.)

See also Arsenyl sulphiodide.

Arsenic sulphoselenide, As₂SeS₂.

Easily sol. in cold NH₄SH + Aq. Nearly completely sol. in (NH₄)₂CO₃ + Aq. (v. Gerichten, B. 7. 29.)

As₂SSe₂. More difficultly sol. than the preceding comp. in NH₄SH + Aq. (v. Gerichten.)

As₂Se₂S₂. Sp. gr. = 6.402 at ca. 750°.

Insol. in most solvents. Easily sol. in alkali hydroxides and sulphides + Aq. (Szarvasky, B. 1895, 28. 2661.)

As₂Se₂S₂. Sp. gr. = 11.35 at 550–600°.

Insol. in most solvents. Easily sol. in alkali hydroxides and sulphides + Aq. (Szarvasky, B. 1895, 28. 2659.)

Arsenic telluride, As₂Te₂.

Sol. in HNO₃ and HNO₃ + HCl + Aq. (Oppenheim, J. pr. 71. 266.)

As₂Te₂. As above. (Oppenheim.)

Arsenic acid, anhydrous, As₂O₄.

See Arsenic pentoxide.

Metaarsenic acid, HAsO₃.

Slowly sol. in cold, quite easily sol. in hot H₂O, with considerable evolution of heat, and conversion into H₂AsO₄. (Kopp, A. ch. (3) 48. 196.)

Orthoarsenic acid, H₂AsO₄.

Sol. in H₂O, with absorption of heat.

1 pt. As₂O₄ dissolves in 0.405 pt. H₂O at 12.5°, or 100 pts. H₂O dissolve 244.81 pts. As₂O₄ at 12.5° (Vogel.)

Sol. in 0.5 pt. H₂O. (Thénard.)

Sol. in 6 pts. cold H₂O, and more quickly in 2 pts. hot H₂O. (Bucholz.)

100 pts. H₂O at 15.56° dissolve 150 pts. As₂O₄. (Ure's Dict.)

H₂AsO₄ + Aq sat. at 15° contains 15% As₂O₄.

Sp. gr. of H₂AsO₄ + Aq at 15°: a = sp. gr. if % is As₂O₄; b = sp. gr. if % is H₂AsO₄.

%	a	b	%	a	b
5	1.042	1.0337	45	1.540	1.3973
10	1.085	1.0690	50	1.635	1.4617
15	1.134	1.1061	55	1.742	1.5320
20	1.187	1.1457	60	1.6086
25	1.245	1.1882	65	1.6919
30	1.306	1.2342	70	1.7827
35	1.378	1.2840	75
40	1.453	1.3382

(Schiff, A. 113. 183, calculated by Gerlach, Z. anal. 27. 303.)

Sp. gr. of $H_3AsO_4 + Aq$ at 15° : a = sp. gr. if % is As_2O_5 ; b = sp. gr. if % is H_3AsO_4 .

%	a	b	%	a	b
1	1.008	1.006	47	1.564	1.412
2	1.016	1.013	48	1.582	1.425
3	1.023	1.019	49	1.601	1.437
4	1.031	1.026	50	1.620	1.450
5	1.039	1.032	51	1.642	1.464
6	1.048	1.039	52	1.663	1.478
7	1.057	1.046	53	1.685	1.491
8	1.065	1.052	54	1.706	1.505
9	1.074	1.059	55	1.728	1.519
10	1.083	1.066	56	1.752	1.534
11	1.092	1.073	57	1.777	1.549
12	1.102	1.081	58	1.801	1.564
13	1.111	1.088	59	1.825	1.579
14	1.121	1.096	60	1.850	1.594
15	1.130	1.103	61	1.880	1.610
16	1.140	1.111	62	1.910	1.626
17	1.150	1.119	63	1.940	1.643
18	1.160	1.126	64	1.970	1.659
19	1.170	1.134	65	2.000	1.675
20	1.180	1.142	66	2.030	1.693
21	1.191	1.150	67	2.060	1.712
22	1.203	1.158	68	2.090	1.730
23	1.214	1.167	69	2.120	1.749
24	1.226	1.175	70	2.150	1.767
25	1.237	1.183	71	...	1.786
26	1.249	1.192	72	...	1.809
27	1.261	1.201	73	...	1.830
28	1.274	1.210	74	...	1.851
29	1.286	1.219	75	...	1.872
30	1.298	1.228	76	...	1.897
31	1.312	1.238	77	...	1.921
32	1.325	1.248	78	...	1.946
33	1.339	1.257	79	...	1.970
34	1.352	1.267	80	...	1.995
35	1.366	1.277	81	...	2.020
36	1.381	1.288	82	...	2.045
37	1.396	1.299	83	...	2.070
38	1.411	1.309	84	...	2.095
39	1.426	1.320	85	...	2.120
40	1.441	1.331	86	...	2.149
41	1.458	1.342	87	...	2.178
42	1.475	1.353	88	...	2.207
43	1.492	1.366	89	...	2.236
44	1.509	1.376	90	...	2.265
45	1.526	1.387	91	...	2.295
46	1.545	1.400

(Kopp, calculated by Gerlach, Z. anal. 27. 316.)

See also Arsenic pentoxide.

Pyroarsenic acid, $H_4As_2O_7$.

Very deliquescent; easily sol. in H_2O with evolution of much heat, and conversion into H_3AsO_4 .

Arsenates.

Arsenates of the alkali metals, and acid arsenates of the alkaline-earth metals are sol. in H_2O . Neutral and basic arsenates are easily sol. in mineral acids, including H_3AsO_4 ;

less sol. in $HC_2H_3O_2 + Aq$. The neutral alkaline-earth arsenates are less sol. in $NH_4OH + Aq$ than in H_2O , but more sol. in $NH_4Cl + Aq$ (Field). The alkali arsenates are sol. in hot glycerine. (Lefèvre, C. R. 100. 1058.)

Aluminum arsenate, $Al_2(AsO_4)_3$.

Ppt. Insol. in H_2O ; difficultly sol. in acids. (Coloriano, C. R. 103. 273.)

Insol. in acetone. (Naumann, B. 1904, 37. 4328.)

$2Al_2O_3, 3As_2O_5$. Nearly unattacked by boiling H_2O ; sol. in dil. acids. (Lefèvre, A. ch. (6) 27. 5.)

Aluminum potassium arsenate, $2Al_2O_3, 3K_2O, 3As_2O_5$.

(Lefèvre.)

Aluminum sodium arsenate, $2Al_2O_3, 3Na_2O, 3As_2O_5$.

(Lefèvre.)

Ammonium arsenate, $(NH_4)_3AsO_4 + 8H_2O$.

Difficultly sol. in H_2O . Less sol. in H_2O than $(NH_4)_2HAsO_4$. (Mitscherlich.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.)

Ammonium hydrogen arsenate,

$(NH_4)_2HAsO_4$.

Effloresces, giving off NH_3 ; more sol. in H_2O than $(NH_4)_3AsO_4$. (Salkowsky, J. pr. 104. 129.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

Ammonium dihydrogen arsenate,

$NH_4H_2AsO_4$.

Not efflorescent. Very sol. in H_2O .

Ammonium barium arsenate, $NH_4BaAsO_4 + \frac{1}{2}H_2O$.

Sol. by 10 days' contact in 1391 pts. H_2O ; in 18,832 pts. of a mixture of 1 pt. $NH_4OH + Aq$ and 3 pts. H_2O ; in 227 pts. of a solution of 1 pt. NH_4Cl in 10 pts. H_2O ; and in 2169 pts. of a solution of 1 pt. NH_4Cl in 10 pts. $NH_4OH + Aq$ and 60 pts. H_2O . (Lefèvre, A. ch. 1892, (6) 27. 13.)

$(NH_4)_2BaH_2(AsO_4)_2$. Efflorescent. Insol. in H_2O ; easily sol. in dil. $HNO_3 + Aq$. (Baumann, Arch. Pharm. 36. 36.)

Ammonium calcium arsenate, $NH_4CaAsO_4 + \frac{1}{2}H_2O$.

1000 pts. pure H_2O dissolve 0.20 pt. this salt; 1000 pts. $NH_4Cl + Aq$ (containing 50 pts. NH_4Cl) dissolve 4.15 pts. this salt; 900 pts. $H_2O + 100$ pts. NH_4OH (sp. gr. = 0.880) dissolve 0.01 pt. this salt. (Field, Chem. Soc. 11. 6.)

Soluble by 10 days' contact in 2167 pts. H_2O at 15° ; in 381 pts. $NH_4Cl + Aq$ (1 : 7); in

43478 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (1 : 3); in 10570 pts. $\text{NH}_4\text{Cl} + \text{NH}_4\text{OH} + \text{Aq}$ (1 : 10 : 60). (Lefèvre, A. ch. 1892, (6) 27. 13.)

+6 H_2O . Sol. in hot, very sl. sol. in cold H_2O ; sl. sol. in NH_4Cl , and $\text{NH}_4\text{OH} + \text{Aq}$. (Wach, Schw. J. 12. 285.)

+7 H_2O . (Bloxam, C. N. 54. 163.)

$(\text{NH}_4)_2\text{CaH}_2(\text{AsO}_4)_2$. Efflorescent. Insol. in H_2O ; easily sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Baumann, Arch. Pharm. 36. 36.)

$(\text{NH}_4)\text{Ca}_2\text{H}_2(\text{AsO}_4)_2 + 3\text{H}_2\text{O}$.

$(\text{NH}_4)\text{Ca}_2\text{H}_2(\text{AsO}_4)_2 + 3\text{H}_2\text{O}$. (Bloxam, C. N. 54. 163.)

Ammonium glucinum arsenate, $\text{NH}_4\text{GlAsO}_4 + 4\frac{1}{2}\text{H}_2\text{O}$.

More stable than the corresponding potassium salt. (Bleyer, Z. anorg. 1912, 75. 291.)

Ammonium iron (ferric) dihydrogen arsenate, $\text{NH}_4\text{H}_2\text{AsO}_4, \text{FeAsO}_4$.

Hydrolyzed by H_2O .

Sol. in cold conc. HCl , hot HNO_3 , hot dil. H_2SO_4 , and in hot arsenic acid + Aq containing 75% arsenic pentoxide.

Sol. in hot conc. $\text{NH}_4\text{OH} + \text{Aq}$. Completely hydrolyzed by caustic alkalis.

Insol. in conc. $\text{NH}_4\text{Cl} + \text{Aq}$ and in 50% acetic acid. (Curtman, J. Am. Chem. Soc. 1910, 32. 628.)

Ammonium magnesium arsenate, $\text{NH}_4\text{MgAsO}_4$.

Sl. sol. in H_2O . Sol. in acids.

Anhydrous salt is sol. in 2784 pts. H_2O at 15°; in 15,904 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (1 : 3) (0.96 sp. gr.); in 1386 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (1 : 70); in 886.7 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (1 : 7); in 3014 pts. NH_4Cl (1 pt.) + NH_4OH (0.96 sp. gr.) (10 pts.) + Aq (60 pts.); in 32,827 pts. magnesia mixture. (Fresenius, Z. anal. 3. 206.)

Anhydrous salt is sol. in 4389 pts. $\text{NH}_4\text{NO}_3 + \text{Aq}$ (1 : 50); in 2561.5 pts. $\text{KCl} + \text{Aq}$ (1 : 165); in 1422 pts. ammoniacal solution of 3.5 g. tartaric acid in 250 cc. H_2O ; in 933.5 pts. ammoniacal solution of 2.5 g. citric acid in 250 cc. H_2O . (Puller, Z. anal. 10. 62.)

+ $\frac{1}{2}\text{H}_2\text{O}$.

Sol. in 2656 pts. H_2O at 15°; in 15,038 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (1 : 3) (0.96 sp. gr.); in 844 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (1 : 7); in 1315 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (1 : 70); in 2871 pts. NH_4Cl (1 pt.) + NH_4OH (0.96 sp. gr.) (10 pts.) + Aq (60 pts.). (Fresenius.)

1000 pts. pure H_2O dissolve 0.14 pt. salt; 1000 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (containing 100 pts. NH_4Cl) dissolve 0.95 pt. salt; 900 pts. H_2O + 100 pts. NH_4OH (sp. gr. 0.880) dissolve 0.07 pt. salt. (Field, Chem. Soc. 11. 6.)

+6 H_2O . Sl. efflorescent. Sl. sol. in H_2O . Very sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Solubility of $\text{NH}_4\text{MgAsO}_4 + 6\text{H}_2\text{O}$ in H_2O and NH_4 salts + Aq .
Grams salt dissolved in 100 g. solvent.

t°	H_2O	5% $\text{NH}_4\text{NO}_3 + \text{Aq}$	5% $\text{NH}_4\text{Cl} + \text{Aq}$	$\text{NH}_4\text{OH} + \text{Aq}$ 1 pt. $\text{NH}_4\text{OH} + \text{Aq}$ (0.96) + 4 pts. H_2O	4% $\text{NH}_4\text{OH} + \text{Aq} + 5\%$ $\text{NH}_4\text{Cl} + \text{Aq}$	4% $\text{NH}_4\text{OH} + \text{Aq} + 10\%$ $\text{NH}_4\text{Cl} + \text{Aq}$
0°	0.03388	0.09216	0.08397	0.00874
20	0.02066	0.11358	0.12284	0.00958	0.01331	0.03165
30	0.11758	0.11264
40	0.02746	0.13936	0.19016	0.01173
50	0.02261	0.18945	0.18889	0.01005
60	0.02103	0.21115	0.21952	0.00902	0.04691	0.05353
70	0.01564	0.18880	0.22092	0.00949
80	0.02364	0.18945	0.23144	0.00912

(Wenger, Dissert. 1911.)

Ammonium manganous arsenate, $\text{NH}_4\text{MnAsO}_4 + 6\text{H}_2\text{O}$.

Nearly insol. in cold H_2O ; easily sol. in dil. acids; insol. in alcohol. (Otto, J. pr. 2. 414.)

Ammonium sodium arsenate, $\text{NH}_4\text{NaHAsO}_4 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Uelsmann, Zeit. f. ges. Nat. 23. 347.)

Ammonium sodium hydrogen arsenate, $(\text{NH}_4)_2\text{Na}_2\text{H}_2(\text{AsO}_4)_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Filhol and Senderens, C. R. 94. 649.)

Ammonium strontium arsenate, $\text{NH}_4\text{SrAsO}_4 + \frac{1}{2}\text{H}_2\text{O}$.

Sol. by 10 days' contact in 3229 pts. H_2O , in 11,586 pts. dil. $\text{NH}_4\text{OH} + \text{Aq}$, in 199 pts.

of a mixture of 1 pt. NH_4Cl in 7 pts. H_2O , and in 1519 pts. of a solution of 1 pt. NH_4Cl in 10 pts. $\text{NH}_4\text{OH} + \text{Aq}$ and 60 pts. H_2O . (Lefèvre, A. ch. 1892, (6) 27. 13.)

Ammonium uranyl arsenate, $\text{NH}_4(\text{UO}_2)\text{AsO}_4 + x\text{H}_2\text{O}$.

Insol. in H_2O , $\text{HC}_2\text{H}_3\text{O}_2$, and saline solutions as $\text{NH}_4\text{Cl} + \text{Aq}$; sol. in mineral acids. (Puller, Z. anal. 10. 72.)

Ammonium vanadium arsenate,

$\text{NH}_4(\text{VO}_2)_2\text{AsO}_4$, and $(\text{NH}_4)_2\text{HAsO}_4 + 2(\text{VO}_2)_2\text{H}_2\text{AsO}_4$.

See Arseniovanadate, ammonium.

Ammonium arsenate tellurate.

See Arseniotellurate, ammonium.

Antimony arsenate (?)

Insol. in H_2O ; insol. in acids after ignition, but when fresh is sol. in conc. boiling $HCl + Aq$, and sl. sol. in $HNO_3 + Aq$. (Dumas.)

Barium arsenate, $Ba_3(AsO_4)_2$.

1000 pts. pure H_2O dissolve 0.55 pt. $Ba_3(AsO_4)_2$; 1000 pts. $NH_4Cl + Aq$ (containing 50 pts. NH_4Cl) dissolve 1.95 pts. $Ba_3(AsO_4)_2$; 900 pts. $H_2O + 100$ pts. $NH_4OH + Aq$ (sp. gr. = 0.88) dissolve 0.03 pt. $Ba_3(AsO_4)_2$. (Field, Chem. Soc. 11. 6.)

Sol. in cold HNO_3 , and $HCl + Aq$ (Berzelius); $H_2C_4H_4O_6$, and $HC_2H_3O_2 + Aq$. (Anthon.)

Solubility in H_2O is not increased by presence of NH_4 , Na, or K salts. (Laugier.)

Not pptd. in presence of Na citrate. (Spiller.)

+ $1\frac{1}{2}H_2O$. (Salkowsky, J. pr. 104. 129.)

Barium hydrogen arsenate, $BaHAsO_4 + 1\frac{1}{2}H_2O$.

Very sl. sol. in H_2O , but decomp. thereby into $Ba_3(AsO_4)_2$ and $BaH_4(AsO_4)_2$. (Berzelius.)

Sl. sol. in cold acids.

+ H_2O . Sl. sol. in either $BaCl_2 + Aq$ or $Na_2HAsO_4 + Aq$. (Maumené, J. B. 1864. 237.)

Barium tetrahydrogen arsenate, $BaH_4(AsO_4)_2 + 2H_2O$.

Easily sol. in H_2O . (Setterberg, Berz. J. B. 26. 206.)

Difficultly sol. in little, but decomp. by much H_2O . Easily sol. in $HCl + Aq$, less easily in $HC_2H_3O_2 + Aq$ (Hörmann, Dissert, 1879.)

Barium arsenate, acid, $BaO, 2As_2O_5 + 4H_2O$.

Very sl. sol. in H_2O . (Mitscherlich.)

Barium pyroarsenate, $Ba_2As_2O_7$.

Insol. in H_2O , but decomp. thereby into $BaHAsO_4 + H_2O$. (Lefèvre, C. R. 108. 1058.)

Barium potassium arsenate, $BaKAsO_4$.

Sl. decomp. by cold H_2O ; rapidly sol. in dil. acids. (Lefèvre, A. ch. (6) 27. 1.)

Barium sodium arsenate, $BaNaAsO_4 + 9H_2O$.

(Joly, C. R. 1887, 104. 1702.)

Barium arsenate chloride, $3Ba_3(AsO_4)_2, BaCl_2$.

Insol. in H_2O ; sol. in dil. $HNO_3 + Aq$. (Lechartier, C. R. 65. 172.)

Bismuth arsenate, basic, $BiAsO_4, 3Bi_2O_3$.

Insol. in H_2O . Sol. in mineral acids. (Cavazzi, Gazz. ch. it. 14. 289.)

$5Bi_2O_3, 2As_2O_5 + 8H_2O$. Min. *Rhagite*.

Easily sol. in $HCl + Aq$; sl. sol. in $HNO_3 + Aq$.

Bismuth arsenate, $BiAsO_4 + \frac{1}{2}H_2O$.

Insol. in H_2O . Insol. in $HNO_3 + Aq$ in presence of H_3AsO_4 , or alkali arsenates + Aq ; sol. in $HCl + Aq$. (Salkowsky, J. pr. 104. 129.)

Not wholly insol. in $HNO_3 + Aq$. (Schneider, J. pr. (2) 20. 418.)

Very sol. in $H_3AsO_4 + Aq$. (Dumas.)

Insol. in $Bi(NO_3)_3 + Aq$. (Dumas.)

Sol. in $Bi(NO_3)_3 + Aq$. (Salkowsky.)

Insol. in conc. $Bi(NO_3)_3 + Aq$ containing a small quantity of HNO_3 . (Schneider.)

Bismuth copper arsenate, $Bi_2Cu_{20}As_{10}H_{44}O_{70} = Bi_2O_3, 20CuO, 5As_2O_5 + 22H_2O$.

Min. *Mixite*. Decomp. by dil. $HNO_3 + Aq$ into insol. $BiAsO_4$, and $Cu_3(AsO_4)_2$, which goes into solution. (Dana.)

Bismuth uranyl arsenate, $Bi_2(AsO_4)_2, 8BiO_2H_2, (UO_2)_2(AsO_4)_2$.

Min. *Walpurgite*.

Cadmium arsenate, $Cd_3(AsO_4)_2$.

Ppt. (Salkowsky, J. pr. 104. 129.)

$2CdO, As_2O_5$. (Lefèvre, C. R. 110. 405.)

$5CdO, 2As_2O_5 + 5H_2O$. Ppt. (Salkowsky.)

Cadmium pyroarsenate, $Cd_2As_2O_7$.

(de Schulten.)

Cadmium hydrogen arsenate, $CdHAsO_4 + H_2O$.

Decomp. by H_2O . (Demel, B. 12. 1279.)

$CdH_4(AsO_4)_2 + 2H_2O$. Decomp. by excess of H_2O . (de Schulten, Bull. Soc. (3) 1. 473.)

Cadmium potassium arsenate, $2CdO, K_2O, As_2O_5$.

(Lefèvre, C. R. 110. 405.)

Cadmium sodium arsenate, $CdO, 2Na_2O, As_2O_5$.

Slowly sol. in dil. acids. (Lefèvre, C. R. 110. 405.)

$2CdO, 4Na_2O, 3As_2O_5$. (Lefèvre.)

Cadmium arsenate bromide, $3Cd_3(AsO_4)_2, CdBr_2$.

Sol. in very dil. $HNO_3 + Aq$. (de Schulten, Bull. Soc. (3) 1. 472.)

Cadmium arsenate chloride, $3Cd_3(AsO_4)_2, CdCl_2$.

Sol. in very dil. $HNO_3 + Aq$. (de Schulten.)

Cæsium arsenate, $Cs_2O, 2As_2O_5 + 5H_2O$.

Ppt. (Ephraim, Z. anorg. 1910, 65. 246.)

Calcium arsenate, $Ca_3(AsO_4)_2 + 3H_2O$.

Ppt. Insol. in H_2O ; sol. in $H_3AsO_4 + Aq$. (Kotschoubey, J. pr. 49. 182.)

Calcium pyroarsenate, $Ca_2As_2O_7$.

Slowly decomp. by cold H_2O into $CaHAsO_4 + 1\frac{1}{2}H_2O$. (Lefèvre.)

Calcium hydrogen arsenate, $\text{CaHAsO}_4 + \frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O . (Debray, A. ch. (3) 61. 419.)
+ H_2O . Min. *Haidingerite*. Easily sol. in acids.

+ $2\frac{1}{2}\text{H}_2\text{O}$. Min. *Pharmacolite*. Easily sol. in acids.

+ $3\text{H}_2\text{O}$. Insol. in H_2O ; sol. in HCl , HNO_3 , or $\text{H}_2\text{AsO}_4 + \text{Aq}$; also in $(\text{NH}_4)_2\text{SO}_4$, NH_4NO_3 , $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$, and $\text{NH}_4\text{Cl} + \text{Aq}$. (Pfaff.)

Calcium tetrahydrogen arsenate,
 $\text{CaH}_4(\text{AsO}_4)_2$.

Sol. in H_2O . (Graham.)

+ H_2O . Sl. sol. in H_2O . Decomp. by much hot H_2O into H_2AsO_4 and $\text{Ca}_3(\text{AsO}_4)_2$. (Hörmann, Dissert. 1879.)

Calcium iron (ferric) arsenate, 6CaO , $4\text{Fe}_2\text{O}_3$, $5\text{As}_2\text{O}_5 + 15\text{H}_2\text{O}$ (?).

Min. *Arseniosiderite*. Sol. in acids.

Calcium magnesium arsenate, $\text{Ca}_3\text{H}_2(\text{AsO}_4)_4$,
 $\text{Mg}_3\text{H}_2(\text{AsO}_4)_4 + 10\text{H}_2\text{O}$.

Min. *Picropharmacolite*. Easily sol. in acids.

$\text{Ca}_3(\text{AsO}_4)_2$, $\text{Mg}_3(\text{AsO}_4)_2$. Sol. in $\text{HNO}_3 + \text{Aq}$. (Kühn.)

Min. *Berzelite*. Sol. in $\text{HNO}_3 + \text{Aq}$.

$\text{Ca}_3\text{Mg}_3\text{H}_{14}(\text{AsO}_4)_{14} + 49\text{H}_2\text{O}$. Min. *Wapplerite*.

Calcium potassium arsenate, CaKAsO_4 .

(Lefèvre, A. ch. (6) 27. 5.)

Calcium sodium arsenate, CaNaAsO_4 .

(Lefèvre, A. ch. (6) 27. 1.)

4CaO , $2\text{Na}_2\text{O}$, $3\text{As}_2\text{O}_5$. Not attacked by boiling H_2O ; easily sol. in dil. acids. (Lefèvre.)

Calcium uranyl arsenate, $\text{Ca}(\text{UO}_2)_2(\text{AsO}_4)_2 + 8\text{H}_2\text{O}$.

Min. *Uranospinite*.

Calcium vanadium arsenate, CaHAsO_4 ,
 $2(\text{VO}_2)\text{H}_2\text{AsO}_4 + 8\text{H}_2\text{O}$.

See *Arseniovanadate*, calcium.

Calcium arsenate chloride, $\text{Ca}_3(\text{AsO}_4)_2$, CaCl_2 .

Insol. in H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Lechartier, C. R. 65. 172.)

$3\text{Ca}_3(\text{AsO}_4)_2$, CaCl_2 . As above. (Lechartier.)

Cerous arsenate, CeHAsO_4 .

Insol. in H_2O . Sol. in arsenic acid + Aq . (Berzelius.)

Ceric hydrogen arsenate, $\text{Ce}(\text{HAsO}_4)_2 + 6\text{H}_2\text{O}$.

Ppt. Insol. in H_2O and dil. acids. (Barbieri, B. 1910, 43. 2216.)

Ceric dihydrogen arsenate, $\text{Ce}(\text{H}_2\text{AsO}_4)_4 + 4\text{H}_2\text{O}$.

Sol. in conc. HNO_3 . (Barbieri l. c.)

Chromic arsenate, $2\text{Cr}_2\text{O}_3$, $3\text{As}_2\text{O}_5$.

Insol. in H_2O and conc. boiling acids. (Lefèvre, A. ch. (6) 27. 5.)

Chromic potassium arsenate, $2\text{Cr}_2\text{O}_3$, $3\text{K}_2\text{O}$,
 $3\text{As}_2\text{O}_5$.

(Lefèvre.)

Chromic sodium arsenate, $2\text{Cr}_2\text{O}_3$, $3\text{Na}_2\text{O}$,
 $3\text{As}_2\text{O}_5$.

(Lefèvre.)

Cobaltous arsenate, basic, 4CoO , As_2O_5 .

Easily sol. in acids. (Gentile, J. B. 1851. 359.)

$\text{Co}(\text{CoOH})\text{AsO}_4$. Insol. in H_2O ; difficultly sol. in acids. (Coloriano.)

Cobaltous arsenate, $\text{Co}_3(\text{AsO}_4)_2 + 8\text{H}_2\text{O}$.

Ppt. Insol. even in boiling H_2O ; easily sol. in HNO_3 , HCl , and $\text{NH}_4\text{OH} + \text{Aq}$; sol. in $\text{H}_2\text{AsO}_4 + \text{Aq}$ (Proust); sol. in dil. $\text{FeSO}_4 + \text{Aq}$. (Karsten, Pogg. 60. 266.)

Min. *Cobalt bloom*, *Erythrite*. Easily sol. in acids.

5CoO , $2\text{As}_2\text{O}_5 + 3\text{H}_2\text{O}$. Insol. in H_2O ; difficultly sol. in acids. (Coloriano, C. R. 103. 273.)

2CoO , As_2O_5 . Sl. attacked by boiling H_2O ; easily sol. in dil. acids. (Lefèvre.)

Cobaltous hydrogen arsenate, $\text{CoH}_4(\text{AsO}_4)_2$.

Sol. in H_2C .

Cobaltous potassium arsenate, CoKAsO_4 .

(Lefèvre.)

Cobaltous sodium arsenate, CoNaAsO_4 .

(Lefèvre.)

4CoO , $2\text{Na}_2\text{O}$, $3\text{As}_2\text{O}_5$. (Lefèvre.)

Cobaltous vanadium arsenate,

$\text{Co}(\text{VO}_2)_2\text{H}_2(\text{AsO}_4)_2 + 8\text{H}_2\text{O}$.

See *Arseniovanadate*, cobaltous.

Cobaltous arsenate ammonia, $\text{Co}_3(\text{AsO}_4)_2$,
 $\text{NH}_3 + 7\text{H}_2\text{O}$.

(Ducru, A. ch. 1901, (7) 22. 185.)

$\text{Co}_3(\text{AsO}_4)_2$, $2\text{NH}_3 + 6\text{H}_2\text{O}$. (Ducru, l. c.)

$\text{Co}_3(\text{AsO}_4)_2$, $3\text{NH}_3 + 5\text{H}_2\text{O}$. (Ducru, l. c.)

Cuprous arsenate, $2\text{Cu}_2\text{O}$, As_2O_5 .

(Hampe, Dissert. 1874.)

$4\text{Cu}_2\text{O}$, As_2O_5 . (Hampe, l. c.)

Cuprous pyroarsenate, $\text{Cu}_4\text{As}_2\text{O}_7$.

Ppt. Sol. in NH_4OH or $\text{KOH} + \text{Aq}$. (Reichard, B. 1898, 31. 2166.)

Cupric arsenate, basic, 8CuO , $\text{As}_2\text{O}_5 + 12\text{H}_2\text{O}$.

Min. *Chalcophyllite*. Easily sol. in acids and $\text{NH}_4\text{OH} + \text{Aq}$.

6CuO , $\text{As}_2\text{O}_5 + 3\text{H}_2\text{O}$. Min. *Aphanesite*, *Clioclasite*. Sol. in acids and ammonia.

5CuO , $\text{As}_2\text{O}_5 + 2\text{H}_2\text{O}$. Min. *Erinite*. Sol. in $\text{HNO}_3 + \text{Aq}$.

+5H₂O. Min. *Cornwallite*. Sol. in acids, and NH₄OH + Aq.

+9H₂O. Min. *Tirolite*.

4CuO, As₂O₃ + H₂O. Insol. in H₂O. (Debray, A. ch. (3) 61. 423.)

Min. *Olivenite*. Sol. in acids, and NH₄OH + Aq; decomp. by hot KOH + Aq.

+7H₂O. Min. *Euchroite*. Sol. in HNO₃ + Aq.

+4½H₂O. (Hirsch, C. C. 1891, I. 15.)

Cupric arsenate, Cu₂(AsO₄)₂.

Insol. in H₂O. Easily sol. in HCl + Aq; sl. sol. in other acids; sol. in NH₄OH + Aq. (Coloriano, C. R. 103. 273.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

+4H₂O. Decomp. by hot H₂O. (Debray.)

+5H₂O. Min. *Trichalcite*. Easily sol. in cold HCl + Aq.

Cupric arsenate, acid, 5CuO, 2As₂O₃.

Sol. in H₂SO₄ + Aq. (Vogel.)

+3H₂O. (Salkowsky.)

+8, 9½, and 12½H₂O. (Hirsch.)

CuHAsO₄ + H₂O. Insol. in H₂O. (Coloriano.)

+1½H₂O. Insol. in H₂O. (Debray, A. ch. (3) 61. 419.)

8CuO, 3As₂O₃ + 12H₂O. (Hirsch.)

Cupric lead arsenate, 3CuO, PbO, As₂O₃ + 2H₂O.

Min. *Bayldonite*. Nearly insol. in HNO₃ + Aq.

Cupric potassium arsenate, CuKAsO₄.

Slowly sol. in NH₄OH + Aq; easily sol. in acids. (Lefèvre, A. ch. (6) 27. 5.)

8CuO, K₂O, As₂O₃. Easily sol. in dil. acids. (Lefèvre.)

Cupric sodium arsenate, CuNaAsO₄.

(Lefèvre.)

3CuO, Na₂O, 2As₂O₃. Very sol. in dil. acids. (Lefèvre.)

2Cu₂(AsO₄)₂, NaH₂AsO₄ + 5H₂O. Ppt. (Hirsch, C. C. 1891, I. 15.)

6Cu₂(AsO₄)₂, 2NaH₂AsO₄, Na₂HAsO₄ + 13½H₂O, or 16H₂O. Ppt. (Hirsch.)

3Cu₂(AsO₄)₂, Na₂HAsO₄ + 9½H₂O. Ppt. (Hirsch.)

4Cu₂(AsO₄)₂, Na₂HAsO₄ + 11H₂O. Ppt. (Hirsch.)

Cupric uranyl arsenate, Cu(UO₂)₂(AsO₄)₂ + 8H₂O.

(Werther, A. 68. 312.)

Min. *Zeunerite*.

Cupric vanadium arsenate,

Cu(VO₂)₂H₂(AsO₄)₂ + 3H₂O.

See *Arseniovanadate, cupric*.

Cupric arsenate ammonia, Cu₂(AsO₄)₂, 3NH₃ + 4H₂O.

Insol. in cold or hot H₂O. (Damour, J. pr. 37. 485.)

2CuO, As₂O₃, 4NH₃ + 3H₂O. Decomp. by H₂O. (Schiff, A. 123. 42.)

Cupric arsenate calcium carbonate, 5CuO, As₂O₃, CaCO₃ + 4H₂O, or 9H₂O.

Min. *Tyrolite*. Easily sol. in acids, and NH₄OH + Aq.

Cupric arsenate sodium chloride, 2Cu₂(AsO₄)₂, NaCl + 7½H₂O.

Decomp. by hot H₂O. (Hirsch, Dissert. 1891.)

3Cu₂(AsO₄)₂, 2NaCl + 13½H₂O.

+17½H₂O. (Hirsch, l.c.)

5Cu₂(AsO₄)₂, 3NaCl + 23H₂O. (Hirsch.)

Didymium arsenate, Di₂H₂(AsO₄)₂.

Ppt. Insol. in H₂O; sl. sol. in weak acids. (Marignac, A. ch. (3) 38. 164.)

5Di₂(AsO₄)₂, As₂O₃ + 3H₂O. Ppt.

Glucinum arsenate, Gl₂(AsO₄)₂.

Insol. in H₂O; sol. in H₂AsO₄ + Aq. (Berzelius.)

Glucinum hydrogen arsenate, GlHAsO₄.

Obtained in impure state by heating As₂O₃ with Gl(OH)₃ in a sealed tube at 220°. (Bleyer, Z. anorg. 1912, 75. 287.)

Glucinum tetrahydrogen arsenate, GlH₄(AsO₄)₂.

Very hygroscopic. (Bleyer, Z. anorg. 1912, 75. 287.)

Glucinum potassium arsenate, KGlAsO₄, ½GlO + 5H₂O.

Unstable. Amorphous. Easily hydrolyzed, giving more basic salts. (Bleyer, Z. anorg. 1912, 75. 289.)

Glucinum sodium arsenate, NaGlAsO₄, ½GlO + 6H₂O.

Unstable. Easily hydrolyzed. (Bleyer, Z. anorg. 1912, 75. 290.)

Iron (ferrous) arsenate, Fe₂(AsO₄)₂ + 6H₂O (?).

Ppt. Sl. sol. in NH₄OH + Aq. Insol. in (NH₄)₂AsO₄ + Aq or other NH₄ salts + Aq. (Wittstein.)

+8H₂O. Min. *Symplectite*. Sol. in HCl + Aq.

Iron (ferric) arsenate, basic, 16Fe₂O₃, As₂O₃ + 24H₂O.

Insol. in NH₄OH + Aq. (Berzelius.)

2Fe₂O₃, As₂O₃ + 12H₂O. Insol. in NH₄OH + Aq.

3Fe₂O₃, 2As₂O₃.

3Fe₂(AsO₄)₂, Fe₂O₃H₂ + 12H₂O. Min. *Pharmacosiderite*. Easily sol. in acids; decomp. by KOH + Aq.

- Iron (ferric) arsenate, $\text{Fe}_2\text{O}_3, \text{As}_2\text{O}_5$.**
 Ppt. Insol. in H_2O . Decomp. by hot H_2O .
 Sol. in HCl , H_2SO_4 , and HNO_3 . (Metzke, Z. anorg. 1898, 19. 473.)
 $+4\text{H}_2\text{O}$. Min. *Scorodite*. Easily sol. in $\text{HCl} + \text{Aq}$; insol. in $\text{HNO}_3 + \text{Aq}$.
 $+8\text{H}_2\text{O}$. Insol. in H_2O . When freshly pptd., sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in HCl , or $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{HC}_2\text{H}_3\text{O}_2$, or NH_4 salts + Aq . (Wittstein.)
 Sol. in warm $\text{H}_2\text{SO}_4 + \text{Aq}$ or $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Berthier, A. ch. (3) 7. 79.)
- Iron (ferric) arsenate, acid, $\text{Fe}_2\text{O}_3, 3\text{As}_2\text{O}_5, +16.7\text{H}_2\text{O}$.**
 Ppt.; sl. sol. in acids with a yellow color, and in $\text{NH}_4\text{OH} + \text{Aq}$ with a red color. (Metzke, Z. anorg. 1898, 19. 476.)
 $2\text{Fe}_2\text{O}_3, 3\text{As}_2\text{O}_5 + 12\text{H}_2\text{O}$. Insol. in H_2O or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.
 Sol. in mineral acids.
 Sol. only in conc. $\text{H}_3\text{AsO}_4 + \text{Aq}$.
 Sol. in $(\text{NH}_4)_2\text{AsO}_4$, and other NH_4 salts + Aq . (Wittstein.)
 Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
 $+22\frac{1}{2}\text{H}_2\text{O}$. Ppt. Sl. sol. in acids with a yellow color, and in $\text{NH}_4\text{OH} + \text{Aq}$ with a red color. (Metzke, Z. anorg. 1898, 19. 475.)
- Iron (ferroferric) arsenate, $6\text{FeO}, 3\text{Fe}_2\text{O}_3, 4\text{As}_2\text{O}_5 + 32\text{H}_2\text{O}$.**
 Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$. (Wittstein, J. B. 1866. 243.)
- Iron (ferric) lead arsenate, $5\text{Fe}_2(\text{AsO}_4)_2, \text{Pb}_3(\text{AsO}_4)_2$.**
 Min. *Carminite Spar. Carminite*. Sol. in acids; $\text{KOH} + \text{Aq}$ dissolves out As_2O_5 . (Sandberger.)
- Iron (ferric) potassium arsenate, $2\text{Fe}_2\text{O}_3, 3\text{K}_2\text{O}, 3\text{As}_2\text{O}_5$.**
 Not attacked by boiling H_2O ; easily sol. in dil. acids. (Lefèvre.)
 $\text{Fe}_2\text{O}_3, \text{K}_2\text{O}, 2\text{As}_2\text{O}_5$. (Lefèvre.)
- Iron (ferric) sodium arsenate, $\text{Fe}_2\text{O}_3, \text{Na}_2\text{O}, 2\text{As}_2\text{O}_5$.**
 (Lefèvre.)
 $2\text{Fe}_2\text{O}_3, 3\text{Na}_2\text{O}, 3\text{As}_2\text{O}_5$. (Lefèvre.)
- Lanthanum arsenate, $\text{La}_2\text{H}_3(\text{AsO}_4)_3$.**
 (Frerichs and Smith.)
 Doubtful. (Cleve, B. 11. 910.)
- Lead arsenate, basic, $15\text{PbO}, 2\text{As}_2\text{O}_5$ (?).**
 Ppt. (Strömholm, Z. anorg. 1904, 38. 446.)
- Lead arsenate, $\text{Pb}_3(\text{AsO}_4)_2$.**
 Insol. in H_2O , NH_4OH , or NH_4 salts + Aq . (Wittstein.)
 Sol. in 2703.5 pts. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ containing 38.94% $\text{HC}_2\text{H}_3\text{O}_2$. (Bertrand, Monit. Scient. (3) 10. 477.)
 Sol. in sat. $\text{NaCl} + \text{Aq}$. (Becquerel, C. R. 20. 1523.)
- Not pptd. in presence of Na citrate. (Spiller.)
- Lead pyroarsenate, $\text{Pb}_2\text{As}_2\text{O}_7$.**
 Insol. in H_2O or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in HCl , or $\text{HNO}_3 + \text{Aq}$. (Rose.)
 Decomp. by cold H_2O . (Lefèvre.)
 $+ \text{H}_2\text{O} = \text{PbHAsO}_4$. Ppt. (Salkowsky, pr. 104. 109.)
- Lead potassium arsenate, PbKAsO_4 .**
 (Lefèvre, A. ch. (6) 27. 5.)
- Lead sodium arsenate, PbNaAsO_4 .**
 (Lefèvre.)
 $4\text{PbO}, 2\text{Na}_2\text{O}, 3\text{As}_2\text{O}_5$. Superficially decomp. by cold H_2O . (Lefèvre.)
- Lead arsenate chloride, $3\text{Pb}_3(\text{AsO}_4)_2, \text{PbCl}_2$.**
 Sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Lechartier.)
 Min. *Mimetite*. Sol. in HNO_3 , and $\text{KOH} + \text{Aq}$.
- Lithium arsenate, Li_3AsO_4 .**
 Ppt. Sol. in dil. acids and in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (de Schulten, Bull. Soc. (3) 1. 479.)
 $\text{LiH}_2\text{AsO}_4 + \frac{3}{2}\text{H}_2\text{O}$. Decomp. by H_2O into H_3AsO_4 and Li_3AsO_4 . (Rammelsberg, Pogg. 128. 311.)
- Magnesium arsenate, $\text{Mg}_3(\text{AsO}_4)_2$.**
 Ppt.
 Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)
 $+7\text{H}_2\text{O}, +8\text{H}_2\text{O}, +10\text{H}_2\text{O}$, and $+22\text{H}_2\text{O}$. (Grühl, Dissert. 1897.)
 $+8\text{H}_2\text{O}$. Min. *Hörnseite*. Insol. in H_2O ; easily sol. in acids.
- Magnesium hydrogen arsenate, MgHAsO_4 .**
 $+ \frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O . (de Schulten, R. 100. 263.)
 $+5\text{H}_2\text{O}$. (Schiefer.)
 $+6\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O . 1000 pts. boiling H_2O dissolve 1.5 pts. (Thompson.)
 Sol. in $\text{HNO}_3 + \text{Aq}$ before ignition, but insol. in acids after ignition. (Graham, A. 29. 29.)
 $+7\text{H}_2\text{O}$. Min. *Roesslerite*. Sol. in $\text{HCl} + \text{Aq}$.
- Magnesium tetrahydrogen arsenate, $\text{MgH}_4(\text{AsO}_4)_2$.**
 Very deliquescent; sol. in H_2O . (Schiefer.)
- Magnesium potassium arsenate, MgKAsO_4 .**
 Insol. in, but decomp. by cold H_2O . (Rose.)
 Easily sol. in dil. acids. (Lefèvre.)
 $+7\text{H}_2\text{O}$. (Kinkelin, Dissert. 1893.)
 $4\text{MgO}, 2\text{K}_2\text{O}, 3\text{As}_2\text{O}_5$. Not attacked by boiling H_2O ; slowly sol. in dil. acids. (Lefèvre.)
- Magnesium potassium hydrogen arsenate, $\text{KMgH}(\text{AsO}_4)_2 + x\text{H}_2\text{O}$.**
 Decomp. by H_2O . (Kinkelin, Dissert. 1883.)

$\text{Mg}_3\text{KH}_2(\text{AsO}_4)_3 + 5\text{H}_2\text{O}$. (Chevron and Droixhe, J. B. 1888, 523.)

Magnesium potassium sodium arsenate, $\text{Mg}_2\text{KNa}(\text{AsO}_4)_2 + 10\text{H}_2\text{O}$.
(Kinkelin, Dissert. 1883.)

Magnesium sodium arsenate, MgNaAsO_4 .
Insol. in H_2O . Very sl. sol. in dil. acids. (Lefèvre.)
 4MgO , $2\text{Na}_2\text{O}$, $3\text{As}_2\text{O}_5$. (Lefèvre.)

Magnesium vanadium arsenate,
 $\text{MgH}_2(\text{VO}_2)_2(\text{AsO}_4)_2 + 9\text{H}_2\text{O}$ and
 $\text{MgHAsO}_4, 2(\text{VO}_2)\text{H}_2\text{AsO}_4 + 9\text{H}_2\text{O}$.
See Arseniovanadate, magnesium.

Magnesium arsenate chloride, $\text{Mg}_3(\text{AsO}_4)_2, \text{MgCl}_2$.
Insol. in H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Lechartier, C. R. 65. 172.)

Magnesium arsenate fluoride, $\text{Mg}_3(\text{AsO}_4)_2, \text{MgF}_2$.
Insol. in H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Lechartier.)

Manganous arsenate, basic, 6MnO , $\text{As}_2\text{O}_5 + 3\text{H}_2\text{O}$ (?).
Min. *Chondroarsenite*. Easily and completely sol. in dil. HCl , and $\text{HNO}_3 + \text{Aq}$.

Manganous arsenate, $\text{Mn}_3(\text{AsO}_4)_2 + \text{H}_2\text{O}$.
Insol. in H_2O ; sl. sol. in acids. (Coloriano, C. R. 103. 273.)
 5MnO , $2\text{As}_2\text{O}_5 + 5\text{H}_2\text{O}$. Insol. in H_2O . (Coloriano.)

2MnO , As_2O_5 . Sl. decomp. by cold H_2O , but rapidly on heating. (Lefèvre.)
 $\text{MnHAsO}_4 + \text{H}_2\text{O}$. Decomp. by boiling H_2O into 5MnO , $2\text{As}_2\text{O}_5 + 5\text{H}_2\text{O}$. Sol. in HNO_3 , H_2SO_4 , or $\text{H}_3\text{AsO}_4 + \text{Aq}$.

Manganous tetrahydrogen arsenate, $\text{MnH}_4(\text{AsO}_4)_2$.
Deliquescent. Easily sol. in H_2O . (Schiefer.)

Manganous potassium arsenate, MnKAsO_4 .
(Lefèvre, A. ch. (6) 27. 5.)

Manganous sodium arsenate, MnNaAsO_4 .
Very sol. in dil. acids. (Lefèvre.)
 2MnO , $4\text{Na}_2\text{O}$, $3\text{As}_2\text{O}_5$. Not attacked by boiling H_2O ; very sol. in dil. acids. (Lefèvre.)

Manganous arsenate chloride, $\text{Mn}_3(\text{AsO}_4)_2, \text{MnCl}_2$.
Insol. in H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Lechartier, A. 58. 259.)

Manganic arsenate, $\text{Mn}_2(\text{AsO}_4)_2 + 2\text{H}_2\text{O}$.
Insol. in H_2O ; sol. in acids.

Mercurous arsenate, $(\text{Hg}_2)_2(\text{AsO}_4)_2$.
Insol. in H_2O ; difficultly sol. in acids. (Coloriano, C. R. 103. 273.) Ppt. (Haack, C. C. 1890, II. 736.)

$\text{Hg}_2(\text{AsO}_4)_2$. Insol. in H_2O , $\text{HC}_2\text{H}_3\text{O}_2$, or alcohol. Decomp. by cold $\text{HCl} + \text{Aq}$. Sl. sol. in cold $\text{HNO}_3 + \text{Aq}$, from which it is precipitated by NH_4OH as Hg_2HAsO_4 . (Simon, Pogg. 41. 424.)

Mercurous hydrogen arsenate, Hg_2HAsO_4 .
Insol. in H_2O , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by cold $\text{HCl} + \text{Aq}$; sol. in cold $\text{HNO}_3 + \text{Aq}$ without decomp; very sl. sol. without decomp. in $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Simon, Pogg. 41. 424.)

Mercuric arsenate, $\text{Hg}_2(\text{AsO}_4)_2$.
Ppt. Sol. in H_3AsO_4 or $\text{HNO}_3 + \text{Aq}$. (Bergman.) Very sl. sol. in H_2O . Easily sol. in $\text{HCl} + \text{Aq}$. Sl. sol. in $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{H}_2\text{AsO}_4 + \text{Aq}$. (Haack, C. C. 1890, II. 736.)

Mercurous silver arsenate, $\text{Hg}_2\text{AgAsO}_4$.
Sol. in hot conc. HNO_3 . (Jacobsen, Bull. Soc. 1909, (4) 5. 948.)

Mercurous arsenate nitrate, $\text{Hg}_2\text{AsO}_4, \text{HgNO}_3 + \text{H}_2\text{O}$.
Insol. in H_2O or $\text{HC}_2\text{H}_3\text{O}_2$; sol. in $\text{HNO}_3 + \text{Aq}$. (Simon, Pogg. 41. 424.)
 $3\text{Hg}_2\text{AsO}_4, 2\text{HgNO}_3, 2\text{Hg}_2\text{O}$. Ppt. (Haack.)

Molybdenum arsenate.
Ppt.

Nickel arsenate, basic, 5NiO , As_2O_5 .
Min. — (Bergemann.)
 $\text{Ni}(\text{NiOH})\text{AsO}_4$. Difficultly attacked by acids or alkalis. (Coloriano, Bull. Soc. (2) 45. 241.)
 5NiO , $2\text{As}_2\text{O}_5 + 3\text{H}_2\text{O}$. As above.

Nickel arsenate, $\text{Ni}_3(\text{AsO}_4)_2$.
Min. — (Bergemann.)
 $+x\text{H}_2\text{O}$. Insol. in H_2O . Sol. in H_3AsO_4 , and conc. mineral acids. Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
 $+2\text{H}_2\text{O}$. Insol. in H_2O ; difficultly sol. in acids. (Coloriano, Bull. Soc. 45. 241.)
 $+8\text{H}_2\text{O}$. Min. *Nickel-bloom*, *Annabergite*. Easily sol. in acids.

$\text{NiHAsO}_4 + \text{H}_2\text{O}$. Sol. in H_2O . Difficultly attacked by acids. (Coloriano, C. R. 103. 274.)

Nickel potassium arsenate, 12NiO , $3\text{K}_2\text{O}$, $5\text{As}_2\text{O}_5$.
(Lefèvre.)
 2NiO , K_2O , As_2O_5 . Rapidly sol. in dil. acids. (Lefèvre.)

Nickel sodium arsenate, NiNaAsO_4 .
Very slowly sol. in dil. acids. (Lefèvre.)
 4NiO , $2\text{Na}_2\text{O}$, $3\text{As}_2\text{O}_5$. (Lefèvre.)

Nickel arsenate ammonia,
 $\text{Ni}_3(\text{AsO}_4)_2, \text{NH}_3 + 7\text{H}_2\text{O}$.
 $\text{Ni}_3(\text{AsO}_4)_2, 2\text{NH}_3 + 6\text{H}_2\text{O}$.
 $\text{Ni}_3(\text{AsO}_4)_2, 3\text{NH}_3 + 5\text{H}_2\text{O}$. (Ducru, C. R. 1900, 131. 703.)

n arsenate (?).

i arsenate (?).

Sol. in $\text{HNO}_3 + \text{Aq.}$

m arsenate, K_2AsO_4 .

escent. Very sol. in H_2O . (Graham, 47.)

in ethyl acetate. (Naumann, B. 3601.)

m hydrogen arsenate, K_2HAsO_4 .

H_2O .

m dihydrogen arsenate, KH_2AsO_4 .

5.3 pts. H_2O at 6° , forming a solution. gr. 1.1134. Much more sol. in hot sol. in alcohol.

6,666 pts. boiling conc. alcohol. (Wenzel.)

m sodium hydrogen arsenate,

$\text{NaHAsO}_4 + 16\text{H}_2\text{O}$.

H_2O .

$\text{H}_4(\text{AsO}_4)_4 + 9\text{H}_2\text{O}$. Sol. in H_2O , and y decomp. thereby into its constituents. (Wohl and Senderens, C. R. 95. 343.)

m strontium arsenate, K_2SrAsO_4 .

re, C. R. 108. 1058.)

m vanadium arsenate, $\text{K}(\text{VO}_2)_2\text{AsO}_4$

$\frac{1}{2}\text{H}_2\text{O}$.

seniovanadate, potassium.

m zinc arsenate, KZnAsO_4 .

re.)

m arsenate sulphate.

seniosulphate, potassium.

a arsenate (?).

m metaarsenate, RbAsO_3 .

H_2O . (Bouchonnet, C. R. 1907, .)

m arsenate, $\text{Rb}_2\text{AsO}_4 + 2\text{H}_2\text{O}$.

hygroscopic; sol. in H_2O to give an solution. Absorbs CO_2 from the air. (Bouchonnet, l.c.)

m pyroarsenate, $\text{Rb}_4\text{As}_2\text{O}_7$.

Bouchonnet, l.c.)

m hydrogen arsenate, $\text{Rb}_2\text{HAsO}_4 + \text{H}_2\text{O}$.

abs CO_2 from the air. Very hydro-sol. in H_2O . Insol. in alcohol. (Bouchonnet, l.c.)

m dihydrogen arsenate, RbH_2AsO_4 .

hygroscopic. Very sol. in H_2O ; aq. is acid to litmus. (Bouchonnet, l.c.)

arsenate, Ag_3AsO_4 .

in H_2O . Sol. in acids; easily sol. in $\text{H}_2\text{O} + \text{Aq.}$ (Joly, C. R. 103. 1071.)

1 l. H_2O dissolves 0.0085 g. Ag_3AsO_4 at 20° . (Whitby, Z. anorg. 1910, 67. 108.)

Much less sol. in H_2AsO_4 than Ag_3PO_4 . (Graham.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Scheele.)

Sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ Insol. in NH_4 sulphate, nitrate, or succinate + Aq. (Wittstein.)

Very sl. sol. in $\text{NH}_4\text{NO}_3 + \text{Aq.}$ more easily in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ (Graham.)

Sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq.}$ but not so easily as Ag_3PO_4 .

Not pptd. in presence of Na citrate. (Spiller.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Silver hydrogen arsenate, Ag_2HAsO_4 .

Decomp. by H_2O , with formation of Ag_3AsO_4 . (Setterberg, Berz. J. B. 26. 208.)

AgH_2AsO_4 . Decomp. by H_2O . (Joly, C. R. 103. 1071.)

Ag_2O , $2\text{As}_2\text{O}_3$. Decomp. by H_2O . Rather sl. sol. in $\text{HNO}_3 + \text{Aq.}$ Very easily sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Hurtzig and Geuther, A. 111. 168.)

Silver arsenate ammonia, $\text{Ag}_3\text{AsO}_4 \cdot 4\text{NH}_3$.

Easily sol. in H_2O . (Widmann, Bull. Soc. (2) 20. 64.)

Silver arsenate sulphate, $3\text{Ag}_2\text{O}$, As_2O_3 , SO_3 .

Decomp. by H_2O , with separation of Ag_3AsO_4 ; decomp. by dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Setterberg, Berz. J. B. 26. 209.)

Sodium arsenate, $\text{Na}_2\text{AsO}_4 + 12\text{H}_2\text{O}$.

Permanent in dry air. Sol. in 3.57 pts. H_2O at 15.5° . (Graham.) 100 pts. H_2O at 15.5° dissolve 28 pts. $\text{Na}_2\text{AsO}_4 + 12\text{H}_2\text{O}$. (Berzelius.) Sol. in 3.75 pts. H_2O at 17° ; or 100 pts. H_2O at 17° dissolve 26.7 pts.; or sat. $\text{Na}_2\text{AsO}_4 + \text{Aq}$ at 17° contains 21.1% $\text{Na}_2\text{AsO}_4 + 12\text{H}_2\text{O}$ or 10.4% Na_2AsO_4 , and has sp. gr. 1.1186. (Schiff, A. 113. 350.)

Melts in crystal H_2O at 85.5° .

Sp. gr. of $\text{Na}_2\text{AsO}_4 + \text{Aq}$ at 17° .

% = % $\text{Na}_2\text{AsO}_4 + 12\text{H}_2\text{O}$.

%	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.0053	9	1.0490	17	1.0945
2	1.0107	10	1.0547	18	1.1003
3	1.0161	11	1.0603	19	1.1061
4	1.0215	12	1.0659	20	1.1121
5	1.0270	13	1.0716	21	1.1179
6	1.0325	14	1.0773	22	1.1238
7	1.0380	15	1.0830
8	1.0435	16	1.0887

(Schiff, calculated by Gerlach, Z. anal. 8. 286.)

"Arsenate of soda" dissolves in 60 pts. boiling alcohol. (Wenzel.)

+ $4\frac{1}{2}\text{H}_2\text{O}$. (Hall, Chem. Soc. 51. 93.)

+ $10\text{H}_2\text{O}$. Efflorescent. (Hall.)

Sodium hydrogen arsenate, $\text{Na}_2\text{HAsO}_4 + 7\text{H}_2\text{O}$.

Not efflorescent. (Schiff.)

Solubility in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$. A table is given which records the g. of As_2O_3 in 100 cc. of the filtrate. (Curry, J. Am. Chem. Soc. 1915, 37, 1685.) $+7\frac{1}{2}\text{H}_2\text{O}$. (Lescocur, C. R. 104, 1171.) $+12\text{H}_2\text{O}$. Efflorescent. Sol. in H_2O ; sol. in 1.79 pts. H_2O at 14° ; or 100 pts. H_2O at 14° dissolve 56 pts. $\text{Na}_2\text{HAsO}_4 + 12\text{H}_2\text{O}$. Sat. $\text{Na}_2\text{HAsO}_4 + \text{Aq}$ contains 35.9% $\text{Na}_2\text{HAsO}_4 + 12\text{H}_2\text{O}$, or 16.5% Na_2HAsO_4 , and has sp. gr. = 1.1722. (Schiff, A. 113, 350.)100 pts. H_2O at 7.2° dissolve 22.268 pts. (Thompson.)100 pts. H_2O dissolve 17.2 pts. $\text{Na}_2\text{HAsO}_4 + 12\text{H}_2\text{O}$ at 0° , and 140.7 pts. at 30° . (Tilden, Chem. Soc. 45, 409.)Melts in crystal H_2O at 28° . (Tilden.)Sp. gr. of $\text{Na}_2\text{HAsO}_4 + \text{Aq}$ at 14° .
 $\% = \frac{c}{c} \text{Na}_2\text{HAsO}_4 + 12\text{H}_2\text{O}$.

%	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.0042	15	1.0665	29	1.1358
2	1.0064	16	1.0712	30	1.1410
3	1.0126	17	1.0759	31	1.1463
4	1.0168	18	1.0807	32	1.1516
5	1.0212	19	1.0855	33	1.1569
6	1.0256	20	1.0904	34	1.1623
7	1.0300	21	1.0953	35	1.1677
8	1.0344	22	1.1003	36	1.1731
9	1.0389	23	1.1052	37	1.1786
10	1.0434	24	1.1103	38	1.1841
11	1.0479	25	1.1153	39	1.1896
12	1.0525	26	1.1204	40	1.1952
13	1.0571	27	1.1255		
14	1.0618	28	1.1306		

(Schiff, calculated by Gerlach, Z. anal. 8, 280.)

Insol. in alcohol.

 $+13\frac{1}{2}\text{H}_2\text{O}$ (Setterberg.)**Sodium dihydrogen arsenate, $\text{NaH}_2\text{AsO}_4 + \text{H}_2\text{O}$.**More sol. in H_2O than Na_2AsO_4 or Na_2HAsO_4 (Schiff.) $+2\text{H}_2\text{O}$ Efflorescent (Joly and Duffet, C. R. 102, 1301.)**Sodium trihydrogen diarsenate, $\text{Na}_3\text{H}_3\text{AsO}_4 + 3\text{H}_2\text{O}$** Sol. in H_2O (Fihol and Senderens, C. R. 95, 343.)**Sodium strontium arsenate, NaSrAsO_4 .**Not attacked by boiling H_2O . (Lefèvre.) $+9\text{H}_2\text{O}$. Scarcely sol. in H_2O (Joly, C. R. 104, 905.) $+18\text{H}_2\text{O}$ (Joly.)**Sodium uranyl arsenate, $\text{Na}_2\text{UO}_2\text{AsO}_4$**

Ppt. (Werther, A. 68, 312.)

Sodium zinc arsenate, NaZnAsO_4 .

Slowly sol. in dil. acids. (Lefèvre.)

 $\text{Na}_2\text{ZnAs}_2\text{O}_7$. As above. (Lefèvre.)**Sodium arsenate fluoride, $\text{Na}_2\text{AsO}_4, \text{NaF} + 12\text{H}_2\text{O}$.**Sol. in 9.5 pts. H_2O at 25° , and 2 pts. at 75° . (Briegleb, A. 97, 95.)**Sodium arsenate stannate, $6\text{Na}_2\text{O}, 2\text{As}_2\text{O}_3, \text{SnO}_2 + 50\text{H}_2\text{O}$.**

More difficultly sol. than sodium stannate. (Haefely, Phil. Mag. (4) 10, 290.)

 $5\text{Na}_2\text{AsO}_4, \text{Na}_2\text{SnO}_3 + 60\text{H}_2\text{O}$. (Prandtl, B. 1907, 40, 2133.)**Sodium arsenate sulphate, $\text{Na}_3\text{As}_2\text{O}_7, 2\text{Na}_2\text{SO}_4$.**Sol. in H_2O . (Mitscherlich.) $\text{Na}_4\text{As}_2\text{O}_7, \text{Na}_2\text{SO}_4$. (Setterberg.)**Sodium arsenate tungstate, $\text{Na}_4\text{As}_2\text{O}_7, \text{Na}_2\text{W}_2\text{O}_{10} + 20\text{H}_2\text{O}$.**

See Arseniotungstate, sodium.

Strontium arsenate, $\text{Sr}_2(\text{AsO}_4)_2$.Not attacked by boiling H_2O ; easily sol. in dil. acids. (Lefèvre, A. ch. (6) 27, 5.)**Strontium pyroarsenate, $\text{Sr}_2\text{As}_2\text{O}_7$.**Decomp. by cold H_2O into $\text{SrHAsO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$. (Lefèvre.)**Strontium hydrogen arsenate, $\text{SrHAsO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$** Insol. in cold, but decomp. by hot H_2O into a basic, and a sol. acid salt. 100 pts. H_2O at 15.5° dissolve 0.284 pt. (Thompson, 1831.)Sol. in $\text{HC}_2\text{H}_3\text{O}_2$, and very easily in $\text{HCl} + \text{Aq}$. (Kotschoubey, J. pr. 49, 182.)Sol. in $\text{HNO}_3 + \text{Aq}$. $\text{SrH}_4(\text{AsO}_4)_2 + 2\text{H}_2\text{O}$. Partly sol. in H_2O . (Hörmann, Dissert. 1879.)**Strontium vanadium arsenate, $\text{SrHAsO}_4, 2(\text{VO})_2\text{H}_2\text{AsO}_4 + 7\frac{1}{2}\text{H}_2\text{O}$**

See Arseniovanadate, strontium.

Strontium arsenate chloride, $3\text{Sr}_2(\text{AsO}_4)_2, \text{SrCl}_2$ Insol. in H_2O , easily sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Lechartier, C. R. 65, 172.)**Thallous arsenate, Tl_2AsO_4** Sol. in H_2O Willm, A. ch. (4) 5, 5.)**Thallous hydrogen arsenate, Tl_2HAsO_4** Very easily sol. in H_2O (Willm.)**Thallous dihydrogen arsenate, TlH_2AsO_4** Easily sol. in H_2O . (Willm.)**Thallic arsenate, $\text{TlAsO}_4 + 2\text{H}_2\text{O}$** Insol. in H_2O , sol. in $\text{HCl} + \text{Aq}$; decomp. by NH_4OH , or $\text{KOH} + \text{Aq}$. (Willm.)

Thorium hydrogen arsenate, $\text{Th}(\text{HAsO}_4)_2 + 6\text{H}_2\text{O}$.

Insol. in H_2O or $\text{H}_2\text{AsO}_4 + \text{Aq}$. (Berzelius.)
Ppt.; insol. in H_2O . (Barbieri, C. A. 1911. 3385.)

$\text{Th}(\text{H}_2\text{AsO}_4)_4 + 4\text{H}_2\text{O}$. Decomp. by H_2O . (Barbieri, l. c.)

Tin (stannous) arsenate, $\text{SnHAsO}_4 + \frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O . (Lenssen, A. 114. 113.)

Tin (stannic) arsenate, $2\text{SnO}_2, \text{As}_2\text{O}_3$.

Ppt. Insol. in H_2O and dil. $\text{HNO}_3 + \text{Aq}$. (Haefely, Phil. Mag. (4) 10. 290.)

$\text{Sn}_3(\text{AsO}_4)_4 + 6\text{H}_2\text{O}$. Insol. in H_2O ; sol. in conc. $\text{HCl} + \text{Aq}$, and in aqua regia; insol. in $\text{HNO}_3 + \text{Aq}$ or H_2SO_4 . (Williams, Proc. Soc. Manchester, 15. 67.)

Colloidal. Very slowly sol. in H_2O , from which it is pptd. by HCl , HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq}$; also by BaCl_2 , CaCl_2 , NH_4Cl , and $\text{FeCl}_3 + \text{Aq}$, and by AgNO_3 , or $\text{KI} + \text{Aq}$. Not pptd. by alcohol, $\text{HC}_2\text{H}_3\text{O}_2$, HgCl_2 , Na_2CO_3 , K_2CO_3 , or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. The pptd. jelly is readily sol. in conc. acids, and KOH , or $\text{NaOH} + \text{Aq}$. (Williams, l. c.)

Tin (stannous) arsenate chloride, $\text{Sn}_3(\text{AsO}_4)_2, \text{SnCl}_2 + 2\text{H}_2\text{O}$.

Decomp. on air. (Lenssen, A. 114. 113.)

Titanium arsenate (?).

Insol. in H_2O . Sol. in titanio acid, arsenic acid, or $\text{HCl} + \text{Aq}$. Sol. in Ti salts + Aq . (Rose.)

Titanyl arsenate, $5\text{TiO}_2, 2\text{As}_2\text{O}_3$.

Sol. in acids without decomp. Scarcely attacked by KOH or by $\text{NH}_4\text{OH} + \text{Aq}$. (Reichard, B. 1894, 27. 1026.)

Uranous arsenate, $\text{U}_3(\text{AsO}_4)_2$.

Ppt.

Uranous hydrogen arsenate, $\text{UH}_2(\text{AsO}_4)_2 + 3\text{H}_2\text{O}$.

Ppt. Sol. in $\text{HCl} + \text{Aq}$.

Uranyl arsenate, $(\text{UO}_2)\text{HAsO}_4 + 4\text{H}_2\text{O}$.

Insol. in H_2O , $\text{HC}_2\text{H}_3\text{O}_2$, and saline solutions, as $\text{NH}_4\text{Cl} + \text{Aq}$; sol. in the mineral acids; sol. in $\text{K}_2\text{CO}_3 + \text{Aq}$. (Werther, A. 68. 313.)

$(\text{UO}_2)_2\text{H}_4(\text{AsO}_4)_2 + 3\text{H}_2\text{O}$. (Werther.)

$(\text{UO}_2)_2\text{As}_2\text{O}_7$. Insol. in H_2O ; sol. in acids.

$(\text{UO}_2)_3(\text{AsO}_4)_2 + 12\text{H}_2\text{O}$.

Min. *Troegerite*.

Vanadium dihydrogen arsenate, $(\text{VO}_2)\text{H}_2\text{AsO}_4 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O . (Friedheim, B. 23. 2600.)

See *Arseniovanadic acid*.

Vanadium zinc arsenate, $(\text{VO}_2)_2\text{ZnH}_2(\text{AsO}_4)_2 + 5\frac{1}{2}\text{H}_2\text{O}$, and $2(\text{VO}_2)\text{H}_2\text{AsO}_4 + 6\frac{1}{2}\text{H}_2\text{O}$.

See *Arseniovanadate, zinc*.

Vanadyl arsenate, $(\text{VO})_2\text{HAsO}_4 + \text{H}_2\text{O}$.

Very slowly sol. in H_2O ; insol. in alcohol; easily sol. in $\text{HCl} + \text{Aq}$. (Berzelius.)

Composition given by Friedheim (B. 23. 2600).

Yttrium arsenate, YtHAsO_4 .

Ppt. Insol. in acetic, easily sol. in mineral acids.

Zinc arsenate, basic, $4\text{ZnO}, \text{As}_2\text{O}_3 + \text{H}_2\text{O}$.

(Friedel, J. B. 1866. 949.)

Min. *Adumite*. Easily sol. in dil. $\text{HCl} + \text{Aq}$, and is attacked by $\text{HC}_2\text{H}_3\text{O}_2$.

Zinc arsenate, $\text{Zn}_3(\text{AsO}_4)_2$.

(deSchulten, Bull. Soc. (3) 2. 300.)

$+ 3\text{H}_2\text{O}$. Ppt. Sol. in HNO_3 , and $\text{H}_3\text{AsO}_4 + \text{Aq}$. (Köttig, J. pr. 48. 182.)
 $+ 8\text{H}_2\text{O}$.

Min. *Köttigite*.

Zinc arsenate, acid, $\text{Zn}_5\text{H}_2(\text{AsO}_4)_4$.

Easily sol. in cold $\text{HCl} + \text{Aq}$, less easily in cold HNO_3 . Sol. in KOH , or $\text{NaOH} + \text{Aq}$ (Gorguel, Dissert, 1894.)

$+ 3\text{H}_2\text{O}$. Insol. in H_2O ; sol. in H_3AsO_4 , or $\text{HNO}_3 + \text{Aq}$. (Mitscherlich.)

$+ 5\text{H}_2\text{O}$. Sol. in dil. $\text{HCl} + \text{Aq}$. (Demel, B. 12. 1279.) Could not be obtained, (Coloriano, Bull. Soc. (2) 45. 709.)

$2\text{ZnO}, \text{As}_2\text{O}_3$. Very slowly decomp. by cold, rapidly by boiling H_2O . (Lefèvre.)

$\text{ZnHAsO}_4 + \text{H}_2\text{O}$. Insol. in H_2O . (Debray, Bull. Soc. (2) 2. 14.)

Decomp. by hot H_2O into $4\text{ZnO}, \text{As}_2\text{O}_3 + \text{H}_2\text{O}$. (Coloriano, C. R. 103. 273.)

$\text{Zn}(\text{ZnOH})_2\text{As}_2\text{O}_7 + 7\text{H}_2\text{O}$ (Gorgeul.)

Zinc arsenate ammonia, $\text{Zn}_3(\text{AsO}_4)_2, 2\text{NH}_3 + 3\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in acids, NH_4OH , or $\text{KOH} + \text{Aq}$. (Bette, A. 15. 141.)

Zirconium arsenate, $2\text{ZrO}_2, \text{As}_2\text{O}_3 + \frac{1}{2}\text{H}_2\text{O} = (\text{ZrO})\text{HAsO}_4 + \frac{3}{2}\text{H}_2\text{O}$.

Ppt. Insol. in H_2O or $\text{HCl} + \text{Aq}$. (Paykull, B. 6. 1467.)

Perarsenic acid.

See *Perarsenic acid*.

Arsenicotungstic Acid.

Ammonium vanadium arsenicotungstate.

See *Arsenicovanadicotungstate, ammonium*.

Arsenicovanadicotungstic acid.

Ammonium arsenicovanadicotungstate,
 $16(\text{NH}_4)_2\text{O}, 5\text{As}_2\text{O}_3, 15\text{V}_2\text{O}_3, 26\text{WO}_3 + 101\text{H}_2\text{O}$.

Sl. sol. in cold, readily sol. in hot H_2O . (Rogers, J. Am. Chem. Soc. 1903, 25. 308.)

Arsenimide, $\text{As}_2(\text{NH})_3$.

Decomp. by H_2O . (Hugot, C. R. 1904, 139. 56.)

Arsenioarsenic acid, $3\text{As}_2\text{O}_3$, $2\text{As}_2\text{O}_5 + 3\text{H}_2\text{O}$.

Decomp. by H_2O . (Joly, C. R. 100. 1221.)
 $3\text{As}_2\text{O}_3$, $\text{As}_2\text{O}_5 + \text{H}_2\text{O}$. Decomp. by H_2O . (Joly.)
 As_2O_3 , $\text{As}_2\text{O}_5 + \text{H}_2\text{O}$. Decomp. by H_2O . (Joly.)

See also *Arsenic trioxide pentoxide*.

Arseniochromic acid.

Ammonium arseniochromate, $2(\text{NH}_4)_2\text{O}$, As_2O_5 , $4\text{CrO}_3 + \text{H}_2\text{O}$.

Insol. in H_2O . (Friedheim and Mozkin, Z. anorg. 1894, 6. 280.)

$3(\text{NH}_4)_2\text{O}$, As_2O_5 , 8CrO_3 . Decomp. by recryst. from H_2O . (Friedheim and Mozkin, Z. anorg. 1894, 6. 281.)

Potassium arseniochromate, $2\text{K}_2\text{O}$, As_2O_5 , 4CrO_3 .

Decomp. by recryst. from H_2O . (Friedheim and Mozkin, Z. anorg. 1894, 6. 275.)

$2\text{K}_2\text{O}$, As_2O_5 , $4\text{CrO}_3 + \text{H}_2\text{O}$. Decomp. by recryst. from H_2O . (Friedheim and Mozkin, l. c.)

Arseniomolybdic acid, As_2O_5 , $6\text{MoO}_3 + 10\text{H}_2\text{O}$.

By recryst. from H_2O the comp. with $18\text{H}_2\text{O}$ is formed. (Pufahl, Dissert. 1888.)

$+16\text{H}_2\text{O}$. Sol. in H_2O . (Debray.)

$+18\text{H}_2\text{O}$. Completely sol. in H_2O . Sp. gr. of sat. solution at 18.8° is 2.21. Easily sol. in abs. alcohol. Insol. in CS_2 , liq. hydrocarbons and CHCl_3 . (Pufahl, l. c.)

As_2O_5 , $7\text{MoO}_3 + 14\text{H}_2\text{O}$. (Seyberth, B. 7. 391.)

As_2O_5 , $18\text{MoO}_3 + 28\text{H}_2\text{O}$. Very sol. in H_2O . Sp. gr. of sat. solution at $18.3^\circ = 2.45$ and 1 cc. contains 2.16 g. acid. Easily sol. in absolute alcohol; insol. in CS_2 , liquid hydrocarbons and CHCl_3 . (Pufahl, l. c.)

Sol. in ether with subsequent separation into two layers. See Phosphotungstic acid. (Drechsel, B. 20. 1452.)

$+38\text{H}_2\text{O}$. Efflorescent. When recryst. comp. with $28\text{H}_2\text{O}$ is formed. (Pufahl, l. c.)

As_2O_5 , $20\text{MoO}_3 + 27\text{H}_2\text{O}$. Sl. sol. in $\text{HNO}_3 + \text{Aq}$. (Debray, C. R. 78. 1408.)

Ammonium arseniomolybdate, $(\text{NH}_4)_2\text{O}$, As_2O_5 , $2\text{MoO}_3 + 3\text{H}_2\text{O}$.

(Friedheim, Z. anorg. 1894, 6. 28.)

$+4\text{H}_2\text{O}$. (Friedheim, l. c.)

$(\text{NH}_4)_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 2\text{H}_2\text{O}$. Sl. sol. in cold H_2O ; sol. in acids. (Debray.)

$+4\text{H}_2\text{O}$. Sl. sol. in cold, very easily sol. in hot H_2O . (Pufahl, l. c.)

$2(\text{NH}_4)_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 6\text{H}_2\text{O}$. Sl. sol. in H_2O . Cannot be recryst. therefrom. (Pufahl.)

$+12\text{H}_2\text{O}$. (Friedheim, Z. anorg. 1894, 6. 31.)

$3(\text{NH}_4)_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 4\text{H}_2\text{O}$. (Friedheim, l. c.)

$+8\text{H}_2\text{O}$. (Friedheim, l. c.)

$(\text{NH}_4)_2\text{O}$, $2\text{H}_2\text{O}$, 7MoO_3 , $\text{As}_2\text{O}_5 + 4\text{H}_2\text{O}$

Sol. in hot H_2O . (Seyberth, B. 7. 391.)

Not obtained. (Pufahl.)

$7(\text{NH}_4)_2\text{O}$, $2\text{As}_2\text{O}_5$, $14\text{MoO}_3 + 28\text{H}_2\text{O}$. (Friedheim, l. c.)

$5(\text{NH}_4)_2\text{O}$, As_2O_5 , $16\text{MoO}_3 + 5\text{H}_2\text{O}$. (Friedheim, Z. anorg. 1894, 6. 31.)

$5(\text{NH}_4)_2\text{O}$, As_2O_5 , $16\text{MoO}_3 + 9\text{H}_2\text{O}$. Nearly insol. in cold, sol. in boiling H_2O . Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Gibbs, Am. Ch. J. 3. 402.)

$+12\text{H}_2\text{O}$. (Pufahl, l. c.)

$2(\text{NH}_4)_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 17\text{H}_2\text{O}$. (Pufahl, l. c.)

$3(\text{NH}_4)_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 14\text{H}_2\text{O}$. Very sol. in H_2O and alcohol. (Kehrmann, Z. anorg. 1894, 7. 421.)

$3(\text{NH}_4)_2\text{O}$, As_2O_5 , 20MoO_3 . Easily sol. in H_2O . (Debray, C. R. 78. 1408.)

$3(\text{NH}_4)_2\text{O}$, As_2O_5 , $24\text{MoO}_3 + 12\text{H}_2\text{O}$. Decomposed by H_2O , especially when boiling. Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$, less easily sol. in warm H_2SO_4 and boiling $\text{H}_2\text{AsO}_4 + \text{Aq}$. Sl. sol. in molybdic acid + Aq , HNO_3 , and conc. $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Pufahl, l. c.)

Barium arseniomolybdate, BaO , As_2O_5 , $6\text{MoO}_3 + 10\text{H}_2\text{O}$.

Sl. sol. in H_2O . Partially decomp. by boiling. (Pufahl, l. c.)

3BaO , As_2O_5 , 6MoO_3 . Sl. sol. in H_2O . (Pufahl, l. c.)

3BaO , As_2O_5 , 7MoO_3 . Ppt. (Seyberth.)

3BaO , As_2O_5 , 18MoO_3 . Decomp. by H_2O . (Pufahl, l. c.)

Cadmium arseniomolybdate, CdO , $2\text{H}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 11\text{H}_2\text{O}$.

(Pufahl.)

3CdO , $3\text{H}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 33\text{H}_2\text{O}$. (Pufahl.)

Cæsium arseniomolybdate, Cs_2O , As_2O_5 , 6MoO_3 .

Sl. sol. in H_2O . (Pufahl, l. c.)

$4\text{Cs}_2\text{O}$, As_2O_5 , $26\text{MoO}_3 + 15\text{H}_2\text{O}$. Ppt. (Ephraim, Z. anorg. 1910, 65. 246.)

Calcium arseniomolybdate, CaO , As_2O_5 , $6\text{MoO}_3 + 10\text{H}_2\text{O}$.

Rather difficultly sol. in cold H_2O . (Pufahl, l. c.)

3CaO , As_2O_5 , 6MoO_3 . As Ba salt. (Pufahl, l. c.)

3CaO , As_2O_5 , $18\text{MoO}_3 + 32\text{H}_2\text{O}$. Very sol. in H_2O . Solution sat. at 18° has sp. gr. = 2.163. (Pufahl, l. c.)

Cobalt arseniomolybdate, CoO , $2\text{H}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 11\text{H}_2\text{O}$.

(Pufahl.)

3CoO , $3\text{H}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 33\text{H}_2\text{O}$. (Pufahl.)

Cupric arseniomolybdate, CuO , $2\text{H}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 15\text{H}_2\text{O}$. (Pufahl.)

3CuO , $3\text{H}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 34\text{H}_2\text{O}$. (Pufahl.)

Lithium arseniomolybdate, Li_2O , As_2O_5 , $6\text{MoO}_3 + 14\text{H}_2\text{O}$.

Very sol. in H_2O . (Pufahl, *l.c.*)

$3\text{Li}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 34\text{H}_2\text{O}$. Solution sat. at 15° has sp. gr. of 2.481. (Pufahl, *l.c.*)

Magnesium arseniomolybdate, MgO , As_2O_5 , $6\text{MoO}_3 + 13\text{H}_2\text{O}$.

Very sol. in H_2O . (Pufahl, *l.c.*)

3MgO , As_2O_5 , $18\text{MoO}_3 + 36\text{H}_2\text{O}$. Sol. in H_2O . (Pufahl, *l.c.*)

Manganese arseniomolybdate, MnO , $2\text{H}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 11\text{H}_2\text{O}$.

(Pufahl.)

3MnO , $3\text{H}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 33\text{H}_2\text{O}$. (Pufahl.)

Nickel arseniomolybdate, NiO , $2\text{H}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 11\text{H}_2\text{O}$.

(Pufahl.)

3NiO , $3\text{H}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 34\text{H}_2\text{O}$. (Pufahl.)

Potassium arseniomolybdate, K_2O , As_2O_5 , $2\text{MoO}_3 + 5\text{H}_2\text{O}$.

Sol. in H_2O . (Friedheim, *Z. anorg.* 2. 314.)
 K_2O , As_2O_5 , $6\text{MoO}_3 + 5\text{H}_2\text{O}$. Sol. in hot H_2O without decomp. (Friedheim, *Z. anorg.* 1892, 2. 330.)

K_2O , As_2O_5 , $18\text{MoO}_3 + 25\text{H}_2\text{O}$. Easily sol. in cold H_2O . Decomp. on dilution. (Pufahl, *l.c.*)

$3\text{K}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 26\text{H}_2\text{O}$. Easily sol. in H_2O . (Pufahl, *l.c.*)

$3\text{K}_2\text{O}$, As_2O_5 , 20MoO_3 . Insol. in H_2O . (Debray, *C. R.* 78. 1408.)

$3\text{K}_2\text{O}$, As_2O_5 , $24\text{MoO}_3 + 12\text{H}_2\text{O}$. Somewhat sol. in H_2O acidified with HNO_3 . (Pufahl, *l.c.*)

Rubidium arseniomolybdate, $3\text{Rb}_2\text{O}$, $3\text{As}_2\text{O}_5$, $5\text{MoO}_3 + 9\text{H}_2\text{O}$.

Easily sol. in H_2O . (Ephraim, *Z. anorg.* 1910, 65. 241.)

Rb_2O , As_2O_5 , 6MoO_3 . Sl. sol. in H_2O . (Pufahl, *l.c.*)

$4\text{Rb}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 40\text{H}_2\text{O}$. Pptd. (Ephraim, *Z. anorg.* 1910, 65. 241-4.)

Silver arseniomolybdate, $3\text{Ag}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + x\text{H}_2\text{O}$.

(Pufahl, Leipzig, 1888.)

$6\text{Ag}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 22\text{H}_2\text{O}$. Sl. sol. in H_2O . Very sol. in NH_4OH and in dil. HNO_3 . (Pufahl, *l.c.*)

$7\text{Ag}_2\text{O}$, $2\text{As}_2\text{O}_5$, $36\text{MoO}_3 + 30\text{H}_2\text{O}$. Sl. sol. in cold, easily sol. in hot H_2O strongly acidified with HNO_3 . (Pufahl, *l.c.*)

Sodium arseniomolybdate, Na_2O , As_2O_5 , $2\text{MoO}_3 + 8\text{H}_2\text{O}$.

(Friedheim, *Z. anorg.* 1892, 2. 357.)

Na_2O , As_2O_5 , $6\text{MoO}_3 + 12\text{H}_2\text{O}$. Very sol. in H_2O . Solution sat. at 19.8° has sp. gr. = 1.678. (Friedheim, *l.c.*)

$3\text{Na}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 11\text{H}_2\text{O} + 12\text{H}_2\text{O}$, and $+13\text{H}_2\text{O}$. Sl. sol. in cold H_2O . (Pufahl, *l.c.*)

$3\text{Na}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 24\text{H}_2\text{O}$. Easily sol. in H_2O . (Pufahl, *l.c.*)

$+30\text{H}_2\text{O}$. Sl. sol. in cold H_2O . (Pufahl, *l.c.*)

Strontium arseniomolybdate, SrO , As_2O_5 , $6\text{MoO}_3 + 10\text{H}_2\text{O}$.

As Ba salt. (Pufahl, *l.c.*)

3SrO , As_2O_5 , 6MoO_3 . As Ba salt. (Pufahl, *l.c.*)

3SrO , As_2O_5 , $18\text{MoO}_3 + 32\text{H}_2\text{O}$. Very sol. in H_2O . (Pufahl, *l.c.*)

Thallium arseniomolybdate, $6\text{Tl}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + x\text{H}_2\text{O}$.

Ppt. (Pufahl.)

$3\text{Tl}_2\text{O}$, $3\text{H}_2\text{O}$, As_2O_5 , $18\text{MoO}_3 + 3\text{H}_2\text{O}$. Ppt. (Pufahl.)

Zinc arseniomolybdate, ZnO , $2\text{H}_2\text{O}$, As_2O_5 , $6\text{MoO}_3 + 11\text{H}_2\text{O}$.

(Pufahl.)

3ZnO , As_2O_5 , $18\text{MoO}_3 + 37\text{H}_2\text{O}$. Very sol. in H_2O . (Pufahl.)

Arseniophosphovanadicotungstic acid.

Ammonium arseniophosphovanadicotungstate, $88(\text{NH}_4)_2\text{O}$, $2\text{As}_2\text{O}_5$, $12\text{P}_2\text{O}_5$, $69\text{V}_2\text{O}_5$, $148\text{WO}_3 + 484\text{H}_2\text{O}$.

Sol. in H_2O . Insol. in alcohol and ether. (Rogers, *J. Am. Chem. Soc.* 1903, 25. 313.)

Arseniophosphovanadicovanadiotungstic acid.

Ammonium arseniophosphovanadicovanadiotungstate, $99(\text{NH}_4)_2\text{O}$, $2\text{As}_2\text{O}_5$, $12\text{P}_2\text{O}_5$, $6\text{V}_2\text{O}_5$, $66\text{V}_2\text{O}_5$, $191\text{WO}_3 + 522\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . (Rogers, *J. Am. Chem. Soc.* 1903, 25. 314.)

Arseniophosphovanadiotungstic acid.

Ammonium arseniophosphovanadiotungstate, $82(\text{NH}_4)_2\text{O}$, $3\text{As}_2\text{O}_5$, $12\text{P}_2\text{O}_5$, $52\text{V}_2\text{O}_5$, $201\text{WO}_3 + 567\text{H}_2\text{O}$.

Very sol. in warm H_2O . Insol. in organic solvents. (Rogers, *J. Am. Chem. Soc.* 1903, 25. 312.)

Arseniosulphuric acid.

Ammonium arseniosulphate, $2(\text{NH}_4)_2\text{O}$, As_2O_5 , $2\text{SO}_3 + 3\text{H}_2\text{O}$.

Can be recryst. from H_2O . (Friedheim and Mozkin, *Z. anorg.* 1894, 6. 290.)

Potassium arseniosulphate, $2\text{K}_2\text{O}$, As_2O_5 , $2\text{SO}_3 + 3\text{H}_2\text{O}$.

(Friedheim and Mozkin, *Z. anorg.* 1894, 6. 289.)

$5\text{K}_2\text{O}$, As_2O_5 , $8\text{SO}_3 + 6\text{H}_2\text{O}$. (Friedheim and Mozkin, *Z. anorg.* 1894, 6. 291.)

Sodium arseniosulphate, $2\text{Na}_2\text{O}$, As_2O_3 , $2\text{SO}_3 + 3\text{H}_2\text{O}$.

(Friedheim and Mozkin, Z. anorg. 1894, 6. 290.)

Arseniotelluric acid.

Ammonium arseniotellurate, $2(\text{NH}_4)_2\text{O}$, As_2O_3 , $\text{TeO}_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Weinland, Z. anorg. 1901, 28. 65.)

$4(\text{NH}_4)_2\text{O}$, $3\text{As}_2\text{O}_3$, $2\text{TeO}_3 + 11\text{H}_2\text{O}$. Sol. in H_2O . (Weinland.)

Sodium arseniotellurate, $2\text{Na}_2\text{O}$, As_2O_3 , $2\text{TeO}_3 + 9\text{H}_2\text{O}$.

Ppt. (Weinland, l.c.)

Arseniotungstic acid, $3\text{H}_2\text{O}$, As_2O_3 , $16\text{WO}_3 + 32\text{H}_2\text{O} = \text{H}_3\text{AsW}_8\text{O}_{28} + 16\text{H}_2\text{O}$ (α -anhydroarsenioluteotungstic acid).

Sol. in H_2O . (Kehrmann, A. 245. 45.)

$3\text{H}_2\text{O}$, As_2O_3 , 19WO_3 (?). Sp. gr. of sat. solution in H_2O is 3.279. (Fremery, B. 17. 296.)

Is a mixture containing principally $\text{H}_3\text{AsW}_8\text{O}_{28} + 16\text{H}_2\text{O}$. (Kehrmann.)

As_2O_3 , $18\text{WO}_3 + x\text{H}_2\text{O}$. Sol. in H_2O . (Kehrmann, Z. anorg. 1899, 22. 292.)

Aluminum ammonium arseniotungstate.

See **Aluminicoarseniotungstate**, ammonium.

Ammonium arseniotungstate, $4(\text{NH}_4)_2\text{O}$, $2\text{H}_2\text{O}$, As_2O_3 , $6\text{WO}_3 + 3\text{H}_2\text{O}$.

Sl. sol. in cold H_2O or $\text{HNO}_3 + \text{Aq}$; easily sol. in boiling H_2O . (Gibbs, Proc. Am. Acad. 16. 135.)

$7(\text{NH}_4)_2\text{O}$, As_2O_3 , $14\text{WO}_3 + 17\text{H}_2\text{O}$. Very sl. sol. even in boiling H_2O . (Fremery, l.c.)

$3(\text{NH}_4)_2\text{O}$, As_2O_3 , $16\text{WO}_3 + 16\text{H}_2\text{O} = (\text{NH}_4)_3\text{AsW}_8\text{O}_{28} + 8\text{H}_2\text{O}$. Sol. in H_2O . (Kehrmann.)

$5(\text{NH}_4)_2\text{O}$, As_2O_3 , $17\text{WO}_3 + 8\text{H}_2\text{O}$. Can be recryst. from H_2O without decomp. Decomp. by long boiling with H_2O . (Kehrmann, Z. anorg. 1899, 22. 294.)

$3(\text{NH}_4)_2\text{O}$, As_2O_3 , $18\text{WO}_3 + 14$, or $18\text{H}_2\text{O}$. Very sol. in cold H_2O . Can be recryst. from H_2O . (Kehrmann, l.c.)

$3(\text{NH}_4)_2\text{O}$, As_2O_3 , $21\text{WO}_3 + x\text{H}_2\text{O}$. Easily sol. in H_2O . Easily decomp. on recryst. (Kehrmann, l.c.)

$3(\text{NH}_4)_2\text{O}$, As_2O_3 , $24\text{WO}_3 + 12\text{H}_2\text{O}$. More sol. in H_2O than corresponding phosphotungstate. (Kehrmann, l.c.)

Barium arseniotungstate, 2BaO , As_2O_3 , $16\text{WO}_3 + x\text{H}_2\text{O}$.

Sol. in H_2O . (Péchar, A. ch. (6) 22. 262.)

7BaO , As_2O_3 , $22\text{WO}_3 + 54\text{H}_2\text{O}$. Sol. in H_2O . Can be recryst. therefrom. (Kehrmann, l.c.)

Potassium arseniotungstate, $3\text{K}_2\text{O}$, $3\text{H}_2\text{O}$, As_2O_3 , 6WO_3 .

Insol. in H_2O . Readily sol. in alkali hydroxides + Aq. (Gibbs.)

$3\text{K}_2\text{O}$, As_2O_3 , $16\text{WO}_3 + 16\text{H}_2\text{O} = \text{K}_3\text{AsW}_8\text{O}_{28} + 8\text{H}_2\text{O}$. Sol. in H_2O . (Kehrmann.)

$5\text{K}_2\text{O}$, As_2O_3 , $17\text{WO}_3 + 22\text{H}_2\text{O}$. Scarcely sol. in cold H_2O . (Kehrmann, Z. anorg. 1899, 22. 295.)

$3\text{K}_2\text{O}$, As_2O_3 , $18\text{WO}_3 + 14\text{H}_2\text{O}$. Efflorescent. (Kehrmann, l.c.)

$3\text{K}_2\text{O}$, As_2O_3 , $19\text{WO}_3 + 16\text{H}_2\text{O}$ (?). Sol. in H_2O . (Fremery.)

Silver arseniotungstate, $\text{Ag}_3\text{AsW}_8\text{O}_{28}$.

Insol. in H_2O (Kehrmann, A. 245. 55); perhaps identical with—

$6\text{Ag}_2\text{O}$, As_2O_3 , $16\text{WO}_3 + 11\text{H}_2\text{O}$. Insol. in H_2O . (Gibbs.)

Sodium arseniotungstate, $3\text{Na}_2\text{O}$, As_2O_3 , $3\text{WO}_3 + 20\text{H}_2\text{O}$.

Very sol. in H_2O . (Lefort, C. R. 92. 1461.)

Arsenious acid, HAsO_2 .

Solubility of HAsO_2 in amyl alcohol + Aq. at 25° .

a_w = mol. of HAsO_2 in 1 l. of H_2O .

a_a = mol. of HAsO_2 in 1 l. of amyl alcohol.

h = partition coefficient.

a_w	a_a	h
0.0449	0.0082	5.48
0.0446	0.0083	5.38
0.0887	0.0164	5.41
0.0892	0.0161	5.53
0.1800	0.0324	5.55

(Auerbach, Z. anorg. 1903, 37. 356.)

Solubility of HAsO_2 in sat. $\text{H}_3\text{BO}_3 + \text{Aq}$ and amyl alcohol.

a_w = mol. of HAsO_2 in 1 l. of H_2O .

a_a = mol. of HAsO_2 in 1 l. of amyl alcohol.

h = partition coefficient.

a_w	a_a	h
0.0859	0.0161	5.33
0.1720	0.0321	5.35

(Auerbach, l.c.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

See **Arsenic trioxide**.

Arsenites.

All arsenites, except those of the alkali metals, are partially or wholly insol. in H_2O , but easily sol. in acids; several are sol. in $(\text{NH}_4)_2\text{SO}_4$, NH_4NO_3 , or $\text{NH}_4\text{Cl} + \text{Aq}$.

All basic arsenites are sol. in acids except those that give an insol. salt with the bases. Many are sol. in excess of $\text{As}_2\text{O}_3 + \text{Aq}$.

Aluminum arsenite, $\text{Al}_2\text{O}_3, \text{As}_2\text{O}_3$.

Sl. sol. in boiling H_2O . Easily sol. in $\text{NaOH} + \text{Aq}$ and in acids. (Reichard, B. 1894, **27. 1029.**)

Aluminum arsenite iodide, $\text{AlI}_3, 6\text{As}_2\text{O}_3 + 16\text{H}_2\text{O}$.

(Grühl, Dissert. 1897.)

Ammonium arsenite, NH_4AsO_2 .

Very sol. in H_2O . (Luynes, J. pr. **72. 180.**)
 Insol. in acetone. (Eidmann, C. C. **1899, II. 1014**); (Naumann, B. 1904, **37. 4328.**)
 $(\text{NH}_4)_2\text{AsO}_3$ (?). Sol. in H_2O . (Stavenhagen, J. pr. 1895, (2) **51. 11.**)

$(\text{NH}_4)_4\text{As}_2\text{O}_5$. Very sol. in H_2O . Insol. in alcohol or ether. (Stein, A. **74. 218.**)

Could not be obtained. (Stavenhagen.)

Ammonium arsenite bromide, $2\text{As}_2\text{O}_3, \text{NH}_4\text{Br}$.

Sl. sol. in H_2O . (Rüdorff, B. **19. 2679.**)

Ammonium arsenite chloride, $\text{As}_2\text{O}_3, \text{NH}_4\text{Cl}$.

Sl. sol. in H_2O . Sol. in warm dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Rüdorff.)

Ammonium arsenite iodide, $2\text{As}_2\text{O}_3, \text{NH}_4\text{I}$.

Sl. sol. in boiling H_2O . Sol. in warm dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Rüdorff.)

Antimony arsenite (?).

Ppt. Sol. in a small amount H_2O , but insol. in a large quantity. (Berzelius.)

Completely sol. in $\text{KOH} + \text{Aq}$. (Reynolds.)

Barium arsenite, $\text{Ba}(\text{AsO}_2)_2$.

Easily sol. in H_2O when recently pptd., but insol. after being dried. Pptd. from aqueous solution by boiling. (Filhol, A. **68. 308.**)

Only sl. sol. in H_2O . (Stavenhagen, J. pr. 1895, (2) **51. 18.**)

$\text{Ba}_3(\text{AsO}_3)_2$. Sl. sol. in cold H_2O ; sol. in hot H_2O and dil. acids. (Stavenhagen, J. pr. 1895, (2) **51. 17.**)

$\text{BaH}_4(\text{AsO}_3)_2$. Ppt. (Bloxam, Chem. Soc. **15. 281.**)

$+34\text{H}_2\text{O}$. Moderately sol. in cold, more easily sol. in hot H_2O . Insol. in alcohol. (Perper, Dissert. **1894.**)

$\text{Ba}_2\text{As}_2\text{O}_5 + 2\text{H}_2\text{O}$. Easily sol. in H_2O . (Stavenhagen, J. pr. 1895, (2) **51. 18.**)

$+4\text{H}_2\text{O}$. Sl. sol. in H_2O ; also somewhat sol. in alcohol. (Stein, A. **74. 218.**)

Sl. sol. in $\text{H}_2\text{AsO}_4 + \text{Aq}$ and $\text{BaO}_2\text{H}_2 + \text{Aq}$. (Dumas.)

Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wackenroder, A. **41. 316.**)

Not pptd. from solutions containing Na citrate. (Spiller.)

BaAs_4O_7 . Sol. in H_2O . Less sol. in alcohol. (Reichard, B. 1894, **27. 1033.**)

Bismuth arsenite, $\text{BiAsO}_3 + 5\text{H}_2\text{O}$ (?).

Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Schneider, J. p. (2) **20. 419.**)

Sl. sol. in H_2O . (Stavenhagen, J. pr. 1895, (2) **51. 35.**)

Cadmium arsenite, $\text{Cd}_3(\text{AsO}_3)_2$.

Sl. sol. in H_2O ; easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$ and dil. acids. (Stavenhagen, *l.c.*)

$\text{Cd}_2\text{As}_2\text{O}_5$. Ppt. (Reichard, B. 1898, **31. 2168.**)

Sol. in acids without decomp.; insol. in alkalis. (Reichard, B. 1894, **27. 1033.**)

$5\text{CdO}, \text{As}_2\text{O}_3 + 12\text{H}_2\text{O}$. Not attacked by KOH , $\text{Ba}(\text{OH})_2$ or alkali carbonates $+ \text{Aq}$. Insol. in $\text{KCN} + \text{Aq}$. (Reichard, Ch. Z. 1902, **26. 1145.**)

Cæsium arsenite bromide, $\text{As}_2\text{O}_3, \text{CsBr}$.

Sol. in H_2O . (Wheeler, Z. anorg. **4. 451.**)

Cæsium arsenite chloride, $\text{As}_2\text{O}_3, \text{CsCl}$.

As above.

Cæsium arsenite iodide, $\text{As}_2\text{O}_3, \text{CsI}$.

As above.

Calcium arsenite, $\text{Ca}(\text{AsO}_2)_2$.

Somewhat sol. in H_2O ; sol. in $\text{Ca}(\text{OH})_2 + \text{Aq}$ or $\text{As}_2\text{O}_3 + \text{Aq}$. (Simon, Pogg. **47. 417.**)

$\text{Ca}_3(\text{AsO}_3)_2$. Ppt. (Kühn, J. B. **1852. 379.**)

Only sl. sol. H_2O ; readily sol. in dil. acids. (Stavenhagen, *l.c.*)

Sol. in H_2O , insol. in alcohol. (Reichard, B. 1894, **27. 1036.**)

$3\text{CaO}, 2\text{As}_2\text{O}_3 + 3\text{H}_2\text{O}$. Sl. sol. in H_2O ; easily sol. in $\text{NH}_4\text{Cl} + \text{Aq}$; sol. in $\text{As}_2\text{O}_3 + \text{Aq}$. (Stein.)

$\text{CaH}_4(\text{AsO}_3)_2 + x\text{H}_2\text{O}$. Moderately sol. in H_2O . Insol. in abs. alcohol. (Perper, Dissert. **1894.**)

$\text{Ca}_2\text{As}_2\text{O}_5$. Sl. sol. in H_2O ; 1 pt. in 3000–4000 pts. H_2O . Alkali chlorides increase solubility slightly. (Stavenhagen, *l.c.*)

Sl. sol. in H_2O ; insol. in H_2O containing CaO_2H_2 . (Berzelius.)

Not pptd. in presence of 4000–5000 pts. H_2O . (Harting, Lassaigue.)

Not pptd. from solutions containing NH_4 salts; and when pptd. is sol. in $(\text{NH}_4)_2\text{SO}_4$, NH_4NO_3 , $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$, and $\text{NH}_4\text{Cl} + \text{Aq}$. (Gieseke and Schweigger.)

Sol. in $\text{NH}_4\text{AsO}_2 + \text{Aq}$. (Schweigger.)

Sol. in $\text{CaCl}_2 + \text{Aq}$. (Ordway.)

Easily sol. in dil. acids. Not pptd. from solutions containing sodium citrate. (Spiller.)

Calcium arsenite iodide, $\text{CaI}_2, 3\text{As}_2\text{O}_3 + 12\text{H}_2\text{O}$.

Sl. sol. in H_2O . Decomp. on heating. (Grühl, Dissert. **1897.**)

Chromic arsenite, CrAsO_3 .

Sol. in H_2O , but slowly decomp. by boiling. (Neville, C. N. **34. 220.**)

Sol. in HCl ; repptd. by $\text{NH}_4\text{OH} + \text{Aq}$; sol. in $\text{KOH} + \text{Aq}$. (Reichard, B. 1894, **27. 1028.**)

Cobaltous arsenite basic, $7\text{CoO}, \text{As}_2\text{O}_3$.

Very sol. in dil., difficultly sol. in conc. H_2SO_4 . Sol. in conc. NaOH and in conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Reichard, Z. anal. 1903, **42. 10.**)

Cobaltous arsenite, $3\text{CoO}, \text{As}_2\text{O}_3$.

Sol. $\text{KOH} + \text{Aq}$ with decomp. (Identical with salt of Girard). (Reichard, B. 1894, **27**. 1031.)

$+4\text{H}_2\text{O}$. Sl. sol. in H_2O ; easily sol. in acids. (Stavenhagen, J. pr. 1895, (2) **51**. 39.)

$3\text{CoO}, 2\text{As}_2\text{O}_3 + 4\text{H}_2\text{O}$. Sol. in HNO_3 . (Girard, C. R. 1852, **34**. 918.)

$\text{Co}_2\text{H}_6(\text{AsO}_3)_4$. Insol. in H_2O ; sol. in HNO_3 , HCl , or $\text{NH}_4\text{OH} + \text{Aq}$. (Proust.)

Only sol. in KOH , or $\text{NaOH} + \text{Aq}$ when formed in a solution containing an excess of those reagents. (Reynoso, C. R. **31**. 68.)

$\text{Co}_2\text{As}_2\text{O}_5$. Ppt. (Reichard, B. 1898, **31**. 2165.)

Sol. in HNO_3 and $\text{HCl} + \text{Aq}$. (Proust.)

Cupric arsenite, $\text{Cu}(\text{AsO}_2)_2$.

(Avery, J. Am. Chem. Soc. 1906, **28**. 1161.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 827.)

$+ \text{H}_2\text{O}$. Sl. sol. in H_2O . (Stavenhagen, l.c.)

$+ 2\text{H}_2\text{O}$. Sl. sol. in H_2O ; insol. in alcohol. (Stavenhagen, l.c.)

$3\text{CuO}, \text{As}_2\text{O}_3$. Ppt. (Stavenhagen, l.c.)

$2\text{CuO}, \text{As}_2\text{O}_3$. (Scheele's green.) Insol. in H_2O ; sol. in $\text{KOH} + \text{Aq}$, $\text{NH}_4\text{OH} + \text{Aq}$, and in most acids. Formula is $\text{Cu}_3(\text{AsO}_3)_2 + 2\text{H}_2\text{O}$. (Sharples, C. N. **35**. 89.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ without decomp. Sol. in $\text{KOH} + \text{Aq}$ with decomp. (Reichard, B. 1894, **27**. 1026.)

Insol. in pyridine. (Schroeder, Dissert. **1901**.)

$5\text{CuO}, \text{As}_2\text{O}_3$. Insol. in H_2O , sol. in acids, $\text{NH}_4\text{OH} + \text{Aq}$ and conc. $\text{MOH} + \text{Aq}$. (Reichard, Ch. Z. 1902, **26**. 1142.)

$x\text{CuO}, y\text{As}_2\text{O}_3$. Min. *Trippkëite*. Easily sol. in HNO_3 and in $\text{HCl} + \text{Aq}$.

Didymium arsenite, $\text{Di}_2\text{H}_3(\text{AsO}_3)_3$.

Ppt. (Frerichs and Smith, A. **191**. 355.)

Does not exist. (Cleve, B. **11**. 910.)

Glucinum arsenite iodide, $\text{GlI}_2, 3\text{As}_2\text{O}_3 + 8\text{H}_2\text{O}$.

Decomp. by H_2O . (Grühl, Dissert. **1897**.)

Gold (aurous) arsenite, $3\text{Au}_2\text{O}, \text{As}_2\text{O}_3$.

Decomp. by light. (Reichard, B. 1894, **27**. 1027.)

Gold (auric) arsenite, $\text{AuAsO}_3 + \text{H}_2\text{O}$.

Very sol. in H_2O , $\text{NH}_4\text{OH} + \text{Aq}$ and dil. acids. (Stavenhagen, J. pr. 1895, (2) **51**. 28.)

Iron (ferrous) arsenite, $\text{FeO}, \text{As}_2\text{O}_3$.

Decomp. in the air when moist; sol. in $\text{NH}_4\text{OH} + \text{Aq}$ when freshly pptd. (Reichard, B. 1894, **27**. 1029-30.)

$\text{Fe}_2\text{As}_2\text{O}_5$. Ppt. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; insol. in NH_4 arsenite, or other NH_4 salts + Aq . (Wittstein.)

Iron (ferric) arsenite, basic, $4\text{Fe}_2\text{O}_3, \text{As}_2\text{O}_3 + 5\text{H}_2\text{O}$.

Ppt. H_2O extracts As_2O_3 . Sol. in conc. acids with separation of As_2O_3 . Acetic acid is without action. (Bunsen and Berthold, **1834**.)

Sol. in KOH , or $\text{NaOH} + \text{Aq}$.

Iron (ferric) arsenite, $\text{Fe}_2\text{O}_3, \text{As}_2\text{O}_3$.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ when freshly pptd. (Reichard, B. 1894, **27**. 1030.)

$\text{Fe}_4\text{As}_2\text{O}_9$. Ppt. (Reichard, B. 1898, **31**. 2170.)

$+ 7\text{H}_2\text{O}$. Sol. in NaOH , and $\text{KOH} + \text{Aq}$.

"Ferric arsenite" is sl. sol. in $\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$. (Kynaston, Dingl. **235**. 326.)

Lanthanum arsenite, $\text{La}_2\text{H}_3(\text{AsO}_3)_3$.

Ppt. (Frerichs and Smith, A. **191**. 355.)

Does not exist. (Cleve, B. **11**. 910.)

Lead arsenite, $\text{Pb}(\text{AsO}_2)_2 + x\text{H}_2\text{O}$.

Sl. sol. in H_2O . Insol. in KOH , but sol. in $\text{NaOH} + \text{Aq}$. (Berzelius.)

$\text{Pb}_2\text{As}_2\text{O}_5$. Insol. in H_2O , NH_4OH , NH_4 arsenite, or other NH_4 salts + Aq . (Wittstein.)

$\text{Pb}_3(\text{AsO}_3)_2$. Scarcely sol. in H_2O ; easily sol. in HNO_3 , or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Boiling H_2O dissolves some As_2O_3 . Not completely insol. in $\text{KOH} + \text{Aq}$. (Streng, A. **129**. 238.)

Sol. in acetic acid; insol. in H_2O in the presence of ammonium salts; sol. in $\text{NaOH} + \text{Aq}$; sl. sol. in $\text{KOH} + \text{Aq}$. (Reichard, B. 1894, **27**. 1024.)

$+ \text{H}_2\text{O}$. Sl. sol. in H_2O ; easily sol. in dil. acids. (Stavenhagen, J. pr. 1895, (2) **51**. 33.)

Lead arsenite chloride, $\text{Pb}_3\text{As}_2\text{O}_5, 2\text{PbCl}_2$.

Min. *Ekdemite*. Easily sol. in $\text{HNO}_3 + \text{Aq}$, and warm $\text{HCl} + \text{Aq}$.

Magnesium arsenite, $\text{Mg}_3(\text{AsO}_3)_2$.

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$, but sol. in a large excess of $\text{NH}_4\text{Cl} + \text{Aq}$. (Rose.)

Very sol. in boiling H_2O and in dil. acids. Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Reichard, B. 1894, **27**. 1032.)

Very sol. in H_2O and dil. acids. (Stavenhagen, l.c.)

$\text{Mg}_3\text{As}_2\text{O}_5 + 4\text{H}_2\text{O}$. Hydroscopic. Very sol. in H_2O and acids. (Stavenhagen, l.c.)

$3\text{MgO}, 2\text{As}_2\text{O}_3 + 3\text{H}_2\text{O}, + 15\text{H}_2\text{O}$, and $+ 18\text{H}_2\text{O}$. (Perper, Dissert. **1894**.)

Magnesium arsenite iodide, $\text{MgI}_2, 3\text{As}_2\text{O}_3 + 12\text{H}_2\text{O}$.

Moderately sol. in H_2O . (Grühl, Dissert. **1897**.)

Manganous arsenite, $\text{Mn}_2(\text{AsO}_3)_2 + 3\text{H}_2\text{O}$.

Sol. in H_2O ; insol. in alcohol; easily oxidized by moist air. (Stavenhagen, l.c.)

$3\text{MnO}, 2\text{As}_2\text{O}_3$. (Reichard, B. 1894, **27**. 1032.)

$\text{Mn}_3\text{H}_2\text{As}_4\text{O}_{10} + 4\text{H}_2\text{O}$. Sl. sol. in H_2O . Very sol. in acids and alkali. (Stavenhagen, *l.c.*)

$\text{Mn}_3\text{As}_2\text{O}_8$. Ppt. (Reichard, B. 1898, 31. 2165.)

Mercurous arsenite, $\text{Hg}_2\text{O}, \text{As}_2\text{O}_3$.

Decomp. by light. Decomp. by H_2O . (Reichard, B. 1894, 27. 1022.)

Hg_3AsO_5 . Only sl. sol. in H_2O ; sol. in dil. acids. (Stavenhagen, J. pr. 1895, (2) 51. 24.)

Gradually and completely decomposed by H_2O . (Reichard, Ch. Z. 1902, 26. 1143.)

Mercuric arsenite, $\text{Hg}_2(\text{AsO}_3)_2$.

Sl. sol. in H_2O . (Stavenhagen, *l.c.*)

Decomp. more easily by H_2O than is the mercurous comp. (Reichard, Ch. Z. 1902, 26. 1143.)

$2\text{HgO}, \text{As}_2\text{O}_3$. Not decomp. by boiling with H_2O . Undecomp. by boiling acids. Decomp. by $\text{KOH} + \text{Aq}$, $\text{K}_2\text{CO}_3 + \text{Aq}$ and $\text{NH}_4\text{OH} + \text{Aq}$. (Reichard, B. 1894, 27. 1021.)

$\text{Hg}_3\text{As}_2\text{O}_8$. Ppt. Decomp. by boiling H_2O . Very sl. sol. in $\text{H}_2\text{SO}_4 + \text{HCl}$. (Reichard, B. 1898, 31. 2170.)

Nickel arsenite, $\text{Ni}_3(\text{AsO}_3)_2$.

Insol. in H_2O ; easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$ (Proust.)

Ppt. (Reichard, B. 1898, 31. 2165.)

$3\text{NiO}, 2\text{As}_2\text{O}_3$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ (identical with salt of Girard). (Reichard, B. 1894, 27. 1031.)

$+ 4\text{H}_2\text{O}$. Insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Proust.)

Sol. in $\text{KOH} + \text{Aq}$. (Girard, C. R. 34. 918.)

$2\text{NiO}, \text{As}_2\text{O}_3$. Insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$; sol. in $\text{KOH} + \text{Aq}$. (Reynoso, C. R. 31. 68.)

Platinum arsenite, $\text{Pt}_3(\text{AsO}_3)_4$.

Sol. in H_2O and alcohol; very unstable. (Stavenhagen, *l.c.*)

Potassium arsenite, KAsO_2 .

Sol. in H_2O ; sl. sol. in alcohol. (Pasteur, A. 68. 309.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Does not exist. (Stavenhagen, *l.c.*)

K_2AsO_3 . Very sol. in H_2O ; sol. in alcohol. (Stavenhagen, *l.c.*)

$\text{K}_4\text{As}_2\text{O}_8 + 6\text{H}_2\text{O}$. Very sol. in H_2O ; sol. in alcohol. (Stavenhagen, *l.c.*)

$\text{K}_2\text{As}_4\text{O}_7 + 2\text{H}_2\text{O}$. Sol. in H_2C ; sl. sol. in alcohol. (Pasteur, A. 68. 309.)

Potassium arsenite bromide, $4\text{As}_2\text{O}_3, 2\text{KBr}$.

More sol. in H_2O than iodide. (Schiff and Sestini, A. 228. 72.)

$2\text{As}_2\text{O}_3, \text{KBr}$. (Rüdorff, B. 19. 2675.)

Potassium arsenite chloride, $2\text{As}_2\text{O}_3, \text{KCl}$.

Much more quickly sol. in hot H_2O than bromide or iodide. (Rüdorff, B. 19. 2675.)

$\text{As}_2\text{O}_3, \text{KCl}$. Decomp. by H_2O .

Potassium arsenite iodide, $3\text{As}_2\text{O}_3, 2\text{KI} + \text{H}_2\text{O}$.

Sl. sol. in cold H_2O ; sol. in 20 pts. boiling, and 40 pts. cold H_2O . (Emmet, Sill. Am. J. (2) 18. 583.)

$6\text{KAsO}_2, 2\text{KI} + 3\text{H}_2\text{O}$. Sol. in H_2O and alcohol. Decomp. by acids. (Harms.)

$2\text{KH}(\text{AsO}_2)_2, \text{As}_2\text{O}_3, 2\text{KI}$. Sl. sol. in H_2O . (Harms, A. 91. 371.)

$2\text{As}_2\text{O}_3, \text{KI}$. Very difficultly sol. even in boiling H_2O . Very easily sol. in $\text{KOH} + \text{Aq}$, but much less so in $\text{K}_2\text{CO}_3 + \text{Aq}$. (Rüdorff, B. 19. 2670.)

Sol. in 40 pts. cold, 20 pts. hot H_2O ; sol. in alkalies. (Schiff and Sestini, A. 228. 72.)

Potassium arsenite sulphate, $\text{K}_3\text{AsO}_3, 10\text{K}_2\text{SO}_4$.

(Stavenhagen, Zeit. angew. ch. 1894, 8. 166.)

Rubidium arsenite, RbAsO_2 .

Sol. in H_2O ; aq. solution is alkaline to litmus. Insol. in alcohol. (Bouchonnet, C. R. 1907, 144. 641.)

Rubidium arsenite bromide, $\text{As}_2\text{O}_3, \text{RbBr}$.

Decomp. by H_2O . (Wheeler, Z. anorg. 4. 451.)

Rubidium arsenite chloride, $\text{As}_2\text{O}_3, \text{RbCl}$.

As above.

Rubidium arsenite iodide, $\text{As}_2\text{O}_3, \text{RbI}$.

As above.

Silver arsenite, Ag_3AsO_3 .

Insol. in H_2O . Not pptd. in presence of 20,000 pts. H_2O . (Harting.)

1 l. H_2O dissolves 0.0115 g. Ag_3AsO_3 at 20° . (Whitby, Z. anorg. 1910, 67. 108.)

Only sl. sol. in H_2O and in dil. acids; readily sol. in $\text{NH}_4\text{OH} + \text{Aq}$ and conc. acids. (Stavenhagen, *l.c.*)

Decomp. by light, by $\text{KOH} + \text{Aq}$ and by $\text{NH}_4\text{OH} + \text{Aq}$. (Reichard, B. 1894, 27. 1022-23.)

Easily sol. in $\text{HNO}_3 + \text{Aq}$ and other acids. (Marcet.)

More easily sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ than Ag_3PO_4 ; sl. sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Santos, C. N. 38. 94.)

Insol. in $\text{KOH} + \text{Aq}$. (Kühn, Arch. Pharm. (2) 69. 267.)

Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Marcet.)

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$, but sol. therein in presence of alkali nitrates. (Santos, *l.c.*)

Incompletely sol. in $(\text{NH}_4)_2\text{CO}_3$, $(\text{NH}_4)_2\text{SO}_4$, or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Wittstein, Repert. 51. 41.)

Decomp. by $\text{NH}_4\text{Cl} + \text{Aq.}$ Sol. in $\text{KAsO}_2 + \text{Aq.}$ (Kühn, *l.c.*)

Not pptd. in solutions containing sol. citrates. (Spiller.)

Sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

Sl. sol. in methyl acetate. (Bezold, Dissert. 1908.)

Insol. in ethyl acetate. (Hamers, Dissert. 1906); (Naumann, B. 1910, **43**. 314.)

+ H_2O . Very sol. in H_2O , $\text{NH}_4\text{OH} + \text{Aq.}$ and in dil. acids. (Stavenhagen, J. pr. 1895, (2) **51**. 29.)

$2\text{Ag}_2\text{O}$, As_2O_3 . Ppt. (Pasteur, J. Pharm. (3) **13**. 395.)

Could not be obtained. (Stavenhagen, *l.c.*)

$3\text{Ag}_2\text{O}$, $2\text{As}_2\text{O}_3$. Sol. in cold $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ (Santos.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ and in potassium arsenite + Aq. (Girard, C. R. **34**. 918.)

Ppt. (Reichard, B. 1898, **31**. 2167.)

Could not be obtained. (Stavenhagen, *l.c.*)

Silver arsenite ammonia, $2\text{Ag}_2\text{O}$, As_2O_3 , 4NH_3 .

Insol. in H_2O or alcohol. (Girard.)

Sodium arsenites.

Correspond to potassium arsenites, but have not been obtained in crystalline form. All are very sol. in H_2O . (Pasteur, A. **68**. 308.)

Na_3AsO_3 . Very sol. in H_2O . (Stavenhagen, *l.c.*)

Insol. in ethyl acetate. (Naumann, B. 1904, **37**. 3602.)

Sodium arsenite bromide, $2\text{As}_2\text{O}_3$, NaBr .

Decomp. by warm H_2O . (Rüdorff, B. **21**. 3052.)

Sodium arsenite iodide, $2\text{As}_2\text{O}_3$, NaI .

Decomp. by hot H_2O . (Rüdorff.)

Strontium arsenite, $\text{Sr}_3(\text{AsO}_3)_2$.

Sol. in H_2O . (Stavenhagen, *l.c.*)

Sol. in H_2O , insol. in alcohol (identical with Stein). (Reichard, B. 1894, **27**. 1036.)

$\text{Sr}_2\text{As}_2\text{O}_5 + 2\text{H}_2\text{O}$. Quite easily sol. in H_2O . (Stein.)

Sl. sol. in H_2C , $\text{SrO}_2\text{H}_2 + \text{Aq.}$ or $\text{H}_3\text{AsO}_4 + \text{Aq.}$ (Dumas.)

Very sl. sol. in alcohol. (Stein.)

Easily sol. in H_2O and in acids. (Stavenhagen, J. pr. 1895, (2) **51**. 17.)

$\text{Sr}_3\text{As}_4\text{O}_9$. Moderately sol. in H_2O . (Reichard, B. 1894, **27**. 1036.)

Strontium arsenite iodide, SrI_2 , $3\text{As}_2\text{O}_3 + 12\text{H}_2\text{O}$.

As Ba comp. (Grühl, Dissert. 1897.)

Thallium arsenite, Tl_3AsO_3 .

Sl. sol. in H_2O and alcohol; easily sol. in acids, especially in dil. H_2SO_4 . (Stavenhagen, *l.c.*)

Tin (stannous) arsenite, $\text{Sn}_2(\text{AsO}_2)_2$.

Ppt.; decomp. by acids and alkali. (Reichard, B. 1898, **31**. 2169.)

+ $2\text{H}_2\text{O}$. Sl. sol. in H_2O . Easily sol. in dil. acids and alkalies. (Stavenhagen, *l.c.*)

Tin (stannic) arsenite, $\text{Sn}_2(\text{AsO}_3)_2 + 5\frac{1}{2}\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Stavenhagen, *l.c.*)

5SnO_2 , $2\text{As}_2\text{O}_3$. Ppt. Sol. in acids without decomp. (Reichard, B. 1894, **27**. 1025.)

$\text{Sn}_7\text{As}_2\text{O}_{17}$. Ppt. (Reichard, B. 1898, **31**. 2169.)

Uranium arsenite, UO_2 , As_2O_3 .

Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$; only sl. sol. $\text{KOH} + \text{Aq.}$ Sol. in acids. (Reichard, B. 1894, **27**. 1029.)

Zinc arsenite, ZnO , As_2O_3 .

Ppt. (Avery, J. Am. Chem. Soc. 1906, **28**. 1163.)

3ZnO , As_2O_3 . Sol. in acids without decomp. Easily sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Reichard, B. 1894, **27**. 1033.)

Arseniovanadic acid, As_2O_5 , $\text{V}_2\text{O}_5 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O , but solution easily decomposes; crystallizes from H_2O with $10\text{H}_2\text{O}$. Composition is vanadium dihydrogen arsenate $(\text{VO}_2)_2\text{H}_2\text{AsO}_4$. (Friedheim, B. **23**. 2600.)

+14, and +18 H_2O . (Ditte, C. R. **102**. 757.) Could not be obtained. (Friedheim.)

$3\text{As}_2\text{O}_5$, $2\text{V}_2\text{O}_5$. (Berzelius.) Correct formula is as above. (Friedheim.)

$3\text{H}_2\text{O}$, $7\text{As}_2\text{O}_5$, $6\text{V}_2\text{O}_5$. (Gibbs, Am. Ch. J. **7**. 209.) Could not be obtained. (Friedheim.)

$3\text{H}_2\text{O}$, $5\text{As}_2\text{O}_5$, $8\text{V}_2\text{O}_5 + 24\text{H}_2\text{O}$. (Gibbs.) Could not be obtained. (Friedheim.)

Arseniovanadates.

According to Friedheim (Z. anorg. 1892, **2**. 319) the arseniovanadates are double arsenates of VO_2 and NH_4 .

Ammonium arseniovanadate, $(\text{NH}_4)_2\text{O}$, As_2O_5 , $2\text{V}_2\text{O}_5 + 5\text{H}_2\text{O}$.

Efflorescent in dry air; sl. sol. in cold, decomp. by hot H_2O . Composition is ammonium divanadium arsenate $= (\text{VO}_2)_2(\text{NH}_4)\text{AsO}_4 + 2\frac{1}{2}\text{H}_2\text{O}$. (Friedheim, B. **23**. 2600.)

Sl. sol. in cold H_2O . Somewhat more easily sol. in hot H_2O with separation of V_2O_5 . (Schmitz-Dumont, Dissert. 1891.)

$2(\text{NH}_4)_2\text{O}$, $3\text{As}_2\text{O}_5$, $2\text{V}_2\text{O}_5 + 4\text{H}_2\text{O}$. Cannot be crystallized from H_2O . Composition is $(\text{NH}_4)_2\text{HAsO}_4 + 2(\text{VO}_2)_2\text{H}_2\text{AsO}_4$. (Friedheim.)

Decomp. under H_2O to $(\text{NH}_4)_2\text{O}$, $2\text{V}_2\text{O}_5$, $\text{As}_2\text{O}_5 + 5\text{H}_2\text{O}$. (Schmitz-Dumont, *l.c.*)

$5(\text{NH}_4)_2\text{O}$, $4\text{As}_2\text{O}_5$, $2\text{V}_2\text{O}_5 + 18\text{H}_2\text{O}$. Sol. in H_2O . (Ditte, C. R. **102**. 1019.) Does not exist. (Friedheim, B. **23**. 2605.)

arseniovanadate, 2CaO , $3\text{As}_2\text{O}_5$, $+21\text{H}_2\text{O} = \text{CaHAsO}_4 + 2(\text{VO}_2)_2\text{O}_4 + 8\text{H}_2\text{O}$.

crystallized in presence of vanadic but decomp. (Friedheim.)

cent. Sol. in H_2O . (Schmitz-l.c.)

eniovandate, CoO , As_2O_5 , V_2O_5 , $+ = \text{Co}(\text{VO}_2)_2\text{H}_2(\text{AsO}_4)_2 + 8\text{H}_2\text{O}$.

I_2O . (Friedheim.)

eniovandate, CuO , As_2O_5 , V_2O_5 , $+ = \text{Cu}(\text{VO}_2)_2\text{H}_2(\text{AsO}_4)_2 + 3\text{H}_2\text{O}$.

I_2O . (Friedheim.)

n arseniovanadate, MgO , As_2O_5 , $+10\text{H}_2\text{O} = (\text{VO}_2)_2\text{MgH}_2(\text{AsO}_4)_2 +$

I_2O . (Friedheim.)

ely sol. in H_2O . Solution decomp. g. (Schmitz-Dumont, l.c.)

As_2O_5 , $2\text{V}_2\text{O}_5$, $+23\text{H}_2\text{O} = \text{MgHAsO}_4$

$\text{I}_2\text{AsO}_4 + 9\text{H}_2\text{O}$. Sol. in H_2O . (Fried-
 I_2O but solution decomp. on evap-
Schmitz-Dumont, l.c.)

arseniovanadate, K_2O , As_2O_5 , $+5\text{H}_2\text{O} = (\text{VO}_2)_2\text{KAsO}_4 + 2\frac{1}{2}\text{H}_2\text{O}$.

I_2O . (Friedheim.)

n cold H_2O . Partially decomp. on
Schmitz-Dumont.)

arseniovanadate, 2SrO , $3\text{As}_2\text{O}_5$, $+20\text{H}_2\text{O} = \text{SrHAsO}_4 + 2(\text{VO}_2)_2\text{H}_2$
 $-7\frac{1}{2}\text{H}_2\text{O}$.

I_2O . (Friedheim.)

l. Easily sol. in H_2O . (Schmitz-

iovanadate, ZnO , As_2O_5 , V_2O_5 , $+ = \text{Zn}(\text{VO}_2)_2\text{H}_2(\text{AsO}_4)_2 + 5\frac{1}{2}\text{H}_2\text{O}$.

I_2O . (Friedheim.)

As_2O_5 , $2\text{V}_2\text{O}_5$, $+5\text{H}_2\text{O}$, and $+18\text{H}_2\text{O}$
 $+2(\text{VO}_2)_2\text{H}_2\text{AsO}_4$, and $+6\frac{1}{2}\text{H}_2\text{O}$.

l. (Friedheim.)

nadicotungstic acid.

arseniovanadicotungstate,
 $(\text{NH}_4)_2\text{O}$, $2\text{As}_2\text{O}_5$, $14\frac{1}{2}\text{V}_2\text{O}_5$, 29WO_3 , $+$

cold H_2O . Readily sol. in boiling
l. in alcohol, ether, benzene, CS_2 ,
etone, nitrobenzene, aniline and
ydride. (Rogers, J. Am. Chem.
25. 307.)

nadicovanadic acid.

arseniovanadicovanadate,
 $(\text{NH}_4)_2\text{O}$, $12\text{As}_2\text{O}_5$, 12VO_2 , $6\text{V}_2\text{O}_5$, $+$

cold, sol. in hot H_2O , from which
—
, $9\text{As}_2\text{O}_5$, 9VO_2 , $8\text{V}_2\text{O}_5$, $+11\text{H}_2\text{O}$.
l. (Gibbs, Am. Ch. J. 7. 209.)

Arseniovanadicovanadiotungstic acid.

Ammonium arseniovanadicovanadiotungstate,
 $17(\text{NH}_4)_2\text{O}$, $2\text{As}_2\text{O}_5$, $7\text{V}_2\text{O}_5$, $4\text{V}_2\text{O}_5$, 32WO_3 ,
 $+73\text{H}_2\text{O}$.

Sl. sol. in cold, readily sol. in boiling H_2O .
(Rogers, J. Am. Chem. Soc. 1903, 25. 310.)

Arseniovanadiotungstic acid.

Ammonium arseniovanadiotungstate,
 $18(\text{NH}_4)_2\text{O}$, $2\text{As}_2\text{O}_5$, $13\text{V}_2\text{O}_5$, 39WO_3 , $+$
 $88\text{H}_2\text{O}$.

Sol. in H_2O . Insol. in organic solvents.
(Rogers, J. Am. Chem. Soc. 1903, 25. 306.)

Arseniuretted hydrogen, AsH_3 .

See Arsenic hydride.

Arsenochromic acid.

Potassium arsenochromate, $\text{K}_4\text{Cr}_2\text{As}_2\text{O}_{16}$, $+$
 $12\text{H}_2\text{O}$.

Sol. in moderately conc. mineral acids.
(Tarugi, C. C. 1897, II. 724.)

$\text{K}_7\text{Cr}_2\text{As}_2\text{O}_{22} + 24\text{H}_2\text{O}$. Ppt. Sol. in dil.
warm acids. (Tarugi.)

Potassium hydrogen arsenochromate,
 $\text{K}_4\text{H}_6\text{Cr}_2\text{As}_2\text{O}_{16}$.

(Tarugi, C. C. 1897, II. 724.)

Arsenosoarseniotungstic acid.

Potassium arsenosoarseniotungstate, $10\text{K}_2\text{O}$,
 $4\text{As}_2\text{O}_5$, As_2O_3 , 21WO_3 , $+26\text{H}_2\text{O}$.

Precipitate. Sol. in a large amount of hot
 H_2O . (Gibbs, Am. Ch. J. 7. 313.)

Arsenosomolybdic acid.

Ammonium arsenosomolybdate, $3(\text{NH}_4)_2\text{O}$,
 $5\text{As}_2\text{O}_3$, 12MoO_3 , $+24\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Gibbs, Am. Ch. J. 7. 313.)

Ammonium barium arsenosomolybdate,
 $3(\text{NH}_4)_2\text{O}$, 2BaO , $5\text{As}_2\text{O}_3$, 10MoO_3 , $+$
 $50\text{H}_2\text{O}$.

Ppt. (Ephraim, Z. anorg. 1910, 66. 57.)

Ammonium cupric arsenosomolybdate,
 $(\text{NH}_4)_2\text{O}$, CuO , $2\text{As}_2\text{O}_3$, 4MoO_3 , $+2\text{H}_2\text{O}$,
and $2(\text{NH}_4)_2\text{O}$, CuO , $3\text{As}_2\text{O}_3$, 6MoO_3 , $+$
 $13\text{H}_2\text{O}$.

Ppts. (Ephraim, Z. anorg. 1910, 66. 58.)

Barium arsenosomolybdate, 3BaO , $2\text{As}_2\text{O}_3$,
 8MoO_3 , $+13\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (Gibbs.)

Copper arsenosomolybdate, 2CuO , $3\text{As}_2\text{O}_3$,
 6MoO_3 .

Sol. in H_2O . (Gibbs.)

Manganese arsenosomolybdate, 2MnO ,
 $3\text{As}_2\text{O}_3$, 6MoO_3 , $+6\text{H}_2\text{O}$, and $+15\text{H}_2\text{O}$.

Insol. in H_2O . (Gibbs.)

Potassium arsenosomolybdate, $3K_2O, As_2O_3, 5MoO_3 + 3H_2O$.

Easily sol. in H_2O . (Ephraim, Z. anorg. 1910, **66**. 54.)

$3K_2O, As_2O_3, 8MoO_3 + 18H_2O$. Easily sol. in H_2O . (Ephraim.)

Sodium arsenosomolybdate, $Na_2O, As_2O_3, 2MoO_3 + 6H_2O$.

Easily sol. in H_2O . (Ephraim, Z. anorg. 1910, **66**. 56.)

$2Na_2O, As_2O_3, 4MoO_3 + 13H_2O$. Ppt. (Ephraim.)

Zinc arsenosomolybdate, $2ZnO, 3As_2O_3, 6MoO_3 + 6H_2O$.

Sol. in H_2O . (Gibbs.)

Arsenosophosphotungstic acid.

Potassium arsenosophosphotungstate, $10K_2O, 14As_2O_3, 3P_2O_5, 32WO_3 + 28H_2O$.

Moderately sol. in cold, very easily in hot H_2O . (Gibbs.)

$7K_2O, 2As_2O_3, 4P_2O_5, 60WO_3 + 55H_2O$. Sol. in hot H_2O with decomp. (Gibbs.)

Potassium sodium arsenosophosphotungstate, $5K_2O, Na_2O, 2As_2O_3, 2P_2O_5, 12WO_3 + 15H_2O$.

(Gibbs, Am. Ch. J. 7. 313.)

Arsenosotungstic acid.

Ammonium arsenosotungstate, $7(NH_4)_2O, 2As_2O_3, 18WO_3 + 18H_2O$.

Sol. in H_2O . (Gibbs.)

Barium arsenosotungstate, $4BaO, As_2O_3, 9WO_3 + 21H_2O$.

Precipitate. Nearly insol. in H_2O . (Gibbs.)

Sodium arsenosotungstate, $9Na_2O, 8As_2O_3, 16WO_3 + 55H_2O$.

Very sol. in H_2O . (Gibbs, Am. Ch. J. 7. 313.)

Arsenyl bromide, $AsOBr$.

H_2O dissolves out As_2O_3 ; insol. in alcohol. (Sérullas.)

$+H_2O$. (Wallace, Phil. Mag. (4) **17**. 122.)

$As_2O_3Br_3 = 2AsBr_3, 3As_2O_3 + 12H_2O$.

Arsenyl bromide with MBr.

See Arsenite bromide, M.

Arsenyl chloride, $AsOCl$.

Sol. in H_2O with decomp.

$+H_2O$. (Wallace, Phil. Mag. (4) **16**. 358.)

As_2O_3Cl . (Wallace.)

Arsenyl chloride with MCl.

See Arsenite chloride, M.

Arsenyl potassium fluoride, $AsOF_3, KF + H_2O$.

(Marignac, A. **145**. 237.)

Arsenyl iodide, $As_2I_2O_{11} = 2AsOI, 3As_2O_3 + 12H_2O$.

Decomp. by H_2O . (Wallace, Phil. Mag. (4) **17**. 122.)

Sl. sol. in cold H_2O , less sol. in alcohol. (Plisson, J. Pharm. **14**. 46.)

Arsenyl iodide with MI.

See Arsenite iodide, M.

Arsenyl sulphiodide, $As_2I_2S_2O_8$.

Scarcely attacked by cold H_2O . Boiling H_2O extracts AsI_3 . Decomp. by hot HNO_3 or H_2SO_4 . Easily sol. in KOH , or $NH_4OH + Aq$. (Schneider, J. pr. (2) **36**. 513.)

Arsine.

See Arsenic hydride.

Atmospheric air.

See Air, atmospheric.

Auriamine, $Au(OH)_2NH_2$.

(Jacobsen, C. R. 1908, **146**. 1214.)

Diauriamine, $Au_2(OH)_4NH_2$.

(Jacobsen, C. R. 1908, **146**. 1214.)

Sesquiauriamine, NAu_3, NH_3 .

Decomp. by H_2O into NAu_3 . (Raschig, A. **235**. 341.)

Auric acid, HAu_2O_4 .

Sol. in HBr , or $HCl + Aq$. (Krüss, B. **19**. 2546.)

Ammonium aurate.

See Auroamidoimide.

Barium aurate, $BaAu_2O_4 + 5H_2O$.

Sl. sol. in H_2O . (Weigand, Zeit. angew. Ch. 1905, **19**. 139.)

$+6H_2O$. Sl. sol. in H_2O . Sol. in dil. H_2SO_4 and in dil. HNO_3 . Sol. in HCl . Decomp. by alcohol. (Meyer, C. R. 1907, **145**. 806.)

Calcium aurate (?).

Insol. in H_2O ; sol. in $CaCl_2 + Aq$. (Freymy, A. ch. (3) **31**. 485.)

$CaAu_2O_4 + 6H_2O$. As Ba salt. (Meyer, C. R. 1907, **145**. 806.)

Magnesium aurate (?).

Ppt. Insol. in H_2O ; sol. in $MgCl_2 + Aq$. (Pelletier.)

Potassium aurate, $KAuO_2 + 3H_2O$.

Very sol. in H_2O , and easily decomp. (Freymy, A. ch. (3) **31**. 483.)

Sol. in alcohol; the solution in alcohol does not decomp. below 50° . (Figuier, A. ch. (3) **11**. 364.)

Potassium aurate sulphite, $KAuO_2, 2K_2SO_3 + 5H_2O$.

Sol. in H_2O with decomp. Nearly insol. in alkaline solutions. (Freymy, A. ch. (3) **31**. 485.)

urate, $\text{Na}_2\text{Au}_2\text{O}_4 + 2\text{H}_2\text{O}$.

H_2O . Sol. in dil. H_2SO_4 , dil. HNO_3 , with decomp. Decomp. by alcohol. C. R. 1907, **145**. 806.)

aurate, $\text{SrAu}_2\text{O}_4 + 6\text{H}_2\text{O}$.

salt. (Meyer.)

le chloride, $\text{Au}(\text{NH})\text{Cl}$.

ig.)

nitrate, $\text{Au}_2\text{N}_2\text{H}_2\text{O}$, 2HNO_3 , or $\text{HNO}_3 + \frac{1}{2}\text{H}_2\text{O}$, or $\text{Au}_2\text{O}(\text{NH})_2$, O_3 .

liquescent. Decomp. by hot H_2O $(\text{NH})_2$. (Schottländer, J. B. 1884.

doimide, $\text{Au}(\text{HN})\text{NH}_2 + 3\text{H}_2\text{O}$.

nating gold.) Insol. in H_2O ; not by dil. acids; sol. in conc. acids, moderately dil. acids, when freshly pre-

Insol. in alkalies or alcohol. Sol. -Aq.

mine, $\text{Au}_3\text{N} + 5\text{H}_2\text{O}$.

comp. by boiling dil. acetic acid, H_2SO_4 . (Raschig, A. 1886, **235**.

hydric acid, $\text{HAu}(\text{CN})_4 + 1\frac{1}{2}\text{H}_2\text{O}$.

sol. in H_2O , alcohol, or ether.

Bromauricyanides.

Chlorauricyanides.

Iodauricyanides.

m auricyanide, $\text{NH}_4\text{Au}(\text{CN})_4$.

sol. in H_2O or alcohol. Insol. in

s auricyanide, $\text{Co}[\text{Au}(\text{CN})_4]_2 + 9\text{H}_2\text{O}$.

in cold, easily in hot H_2O . Sl. sol. l. (Lindbom.)

n auricyanide, $\text{KAu}(\text{CN})_4 + 1\frac{1}{2}\text{H}_2\text{O}$.

cent. Sl. sol. in cold, easily in hot sily sol. in alcohol.

ricyanide, AgAuCN_4 .

in H_2O or $\text{HNO}_3 + \text{Aq}$. Sol. in -Aq.

amine nitrate.

rimide nitrate.

mhydric acid.

mauric acid.

mic acid.

auric acid.

orhydric acid.

lorauric acid.

ric acid.

lorauric acid.

Aurocyanhydric acid.

Aurocyanides with MCN .

See Cyanide, aurous with MCN .

Azinosulphonic acid.

Ammonium azinosulphonate, $\text{N}_2\text{SO}_3\text{NH}_4$.

(Traube, B. 1914, **47**. 944.)

Barium azinosulphonate, $(\text{N}_2\text{SO}_3)_2\text{Ba}$.

(Traube, B. 1914, **47**. 944.)

Potassium azinosulphonate, $\text{N}_2\text{SO}_3\text{K}$.

Easily sol. in H_2O . Can be cryst. from boiling abs. alcohol. (Traube, B. 1914, **47**. 943.)

Sodium azinosulphonate, $\text{N}_2\text{SO}_3\text{Na}$.

(Traube, B. 1914, **47**. 944.)

Azoimide, HN_3 .

Miscible with H_2O and alcohol. (Curtius and Radershausen, J. pr. (2) **43**. 207.)

Stable in aq. solution; decomp. slowly by dil. boiling HCl . (Curtius, J. pr. 1898, (2) **58**. 265.)

For salts of HN_3 , see azoimide of metal under metal.

Azoimide, hydroxylamine, $\text{N}_3\text{H}_2\text{NH}_2\text{OH}$.

Sol. in H_2O . Gradually volatilizes at ord. temp. (Dennis, J. Am. Chem. Soc. 1907, **29**. 22.)

Azophosphoric acid.

See Pyrophosphamic acid.

Deutazophosphoric acid.

See Pyrophosphodiamic acid.

Barium, Ba.

Decomp. by H_2O and abs. alcohol. (Guntz, C. R. 1901, **133**. 874.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 827.)

Barium amalgam, BaHg_{13} .

Stable in contact with liquid amalgam up to 30° . Can be cryst. from Hg without decomp. if temp. does not exceed 30° . (Kerp, Z. anorg. 1900, **25**. 68.)

BaHg_{13} . Stable in contact with liquid amalgam from 30° – 100° . Can be cryst. from Hg without decomp. at any temp. within these limits. (Kerp.)

Barium amide, $\text{Ba}(\text{NH}_2)_2$.

B.-pt. 280° . (Mentrel, C. C. 1903, I. 276.)

Decomp. by H_2O . (Guntz and Mentrel, Bull. Soc. 1903, (3) **29**. 578.)

Barium potassium amide.

See Potassium ammonobarate.

Barium ammonia, $\text{Ba}(\text{NH}_3)_4$.

Takes fire in the air. Only sl. sol. in liquid NH_3 . Violently decomp. by H_2O . (Mentrel, C. R. 1902, **135**. 740.)

Barium arsenide, Ba₃As₂.

Decomp. by H₂O. (Lebeau, C. R. 1899, 129. 48.)

Barium azoimide, Ba(N₃)₂.

Very sl. hygroscopic; explosive.

12.5 pts. are sol. in 100 pts. H₂O at 0°

16.2 " " " " 100 " H₂O " 10.5°

16.7 " " " " 100 " H₂O " 15°

17.3 " " " " 100 " H₂O " 17°

0.0172 pts. are sol. in 100 pts. abs. alcohol at 16°.

Insol. in ether. (Curtius, J. pr. 1898, (2) 58. 290.)

See also Barium nitride.

Barium boride, BaB₆.

Sol. in fused oxidizing agents, not decomp. by H₂O; insol. in aq. acids; sl. sol. in conc. H₂SO₄, sol. in dil. and conc. HNO₃. (Moissan, C. R. 1897, 125, 634.)

Barium subbromide sodium bromide, BaBr, NaBr.

Decomp. by H₂O. (Guntz, C. R. 1903, 136. 750.)

Barium bromide, BaBr₂, and +2H₂O.

100 pts. H₂O dissolve—

at 0° 20° 40° 60° 80° 100°
98 104 114 123 135 149 pts. BaBr₂.

Sat. BaBr₂ + Aq contains at:

—20° —9° +7° 16° 19° 40°
45.7 46.5 48.5 48.8 49.3 50.9% BaBr₂

71° 76° 77° 104° 145° 160° 175°

55.1 55.5 55.6 56.6 60.5 59.4 60.3% BaBr₂
(Étard, A. ch. 1894, (7) 2. 540.)

Sp. gr. of BaBr₂ + Aq at 19.5° containing:

5 10 15 20 25 30% BaBr₂
1.045 1.092 1.114 1.201 1.262 1.329

35 40 45 50 55% BaBr₂.
1.405 1.485 1.580 1.685 1.800

(Kremers, Pogg. 99. 444, calculated by Gerlach, Z. anal. 8. 285.)

BaBr₂ + Aq containing 7.74% BaBr₂ has sp. gr. 20°/20° = 1.0716.

BaBr₂ + Aq containing 16.76% BaBr₂ has sp. gr. 20°/20° = 1.1674.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 279.)

Sat. BaBr₂ + Aq boils at 113°. (Kremers, Pogg. 99. 43.)

Solubility in BaI₂ + Aq at t°.

t°	Sat. solution contains	
	% BaBr ₂	% BaI ₂
—16	4.7	57.9
—16	5.0	59.0
+60	5.5	66.0
135	9.3	67.3
135	9.0	67.2
170	11.0	67.4
210	14.9	67.7

(Étard, A. ch. 1894, (7) 3. 287.)

Very sol. in absolute alcohol. (Hünefeld.)

100 pts. absolute methyl alcohol dissolve 50 pts. BaBr₂ at 22.5°.

100 pts. absolute ethyl alcohol dissolve 3 pts. BaBr₂ at 22.5°. (de Bruyn, Z. phys. Ch. 10. 783.)

Sat. solution in 87% alcohol contains 6% BaBr₂. (Richards, Z. anorg. 3. 455.)

100 pts. absolute methyl alcohol dissolve 45.8 pts. BaBr₂ + 2H₂O at 15°.

100 pts. 93.5% methyl alcohol dissolve 27.3 pts. BaBr₂ + 2H₂O at 15°.

100 pts. 50% methyl alcohol dissolve 4 pts. BaBr₂ + 2H₂O at 15°. (de Bruyn, Z. phys. Ch. 10. 787.)

100 g. BaBr₂ + CH₃OH contain 0.4 g. BaBr₂ at the critical temp. (Centnerszwer, Z. phys. Ch. 1910, 72. 437.)

At 15°, 1 pt. by weight is sol. in:

36 pts. methyl alcohol, sp. gr. 0.709

207 " ethyl " " " 0.8035

652 " propyl " " " 0.8085

(Rohland, Z. anorg. 1897, 15. 413.)

Nearly insol. in boiling amyl alcohol, 10 ccm. dissolving only an amt. equal to 1.3 mg BaO. (Browning, Sill. Am. J. 144. 459.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328; Eidmann, C. C. 1899, II. 1014.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3789.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Barium cadmium bromide, BaBr₂, CdBr₂ + 4H₂O.

Sol. in H₂C. (v. Hauer, W. A. B. 20. 40.)

Barium rhodium bromide.

See Bromorhodite, barium.

Barium bromide ammonia, BaBr₂·8NH₃.

Decomp. by H₂O. (Joannis, C. R. 1905, 140. 1244.)

Barium bromide hydrazine, BaBr₂·2N₂H₄.

Hygroscopic. Very sol. in H₂O. Insol. in alcohol. (Franzen, Z. anorg. 1908, 60. 291.)

Barium bromofluoride, BaBr₂·BaF₂.

Insol. in and undecomp. by boiling alcohol. Sol. in HBr and in HNO₃. Decomp. by H₂O, hot H₂SO₄, dil. HCl, dil. HNO₃, or dil. acetic acid. (Defacqz, C. R. 1904, 138. 199.)

Barium carbide, BaC₂.

Decomp. by H₂O. (Maquenne, C. R. 144. 360.)

Sp. gr. 3.75. Easily decomp. by H₂O and dil. acids. (Moissan, Bull. Soc. 1894, (3) 11. 1008.)

carbonyl, Ba(CO)₂.
in H₂O. (Guntz and Mentrel, Bull. 33, (3) 29. 586.)

subchloride, BaCl.
mp. by H₂O. (Guntz, C. R. 1903, l.)

subchloride sodium chloride, BaCl, Cl.
mp. by H₂O. (Guntz, C. R. 1903, l.)

chloride, BaCl₂, and +2H₂O.
anent in dry air.

1. H₂O at t° dissolve (a) pts. BaCl₂ and (b) pts. BaCl₂+2H₂O.

a	b	t°	a	b
34.86	43.50	74.89	59.94	65.51
43.84	55.63	105.48	59.58	77.89

(Gay-Lussac, A. ch. (2) 11. 309.)
H₂O at t° dissolve 32.62 + 0.2711t pts. BaCl₂.

ts. H₂O dissolve pts. BaCl₂ + 2H₂O at t°.

Pts. BaCl ₂ + 2H ₂ O	t°	Pts. BaCl ₂ + 2H ₂ O
39.66	62.50	48.0
42.22	75.00	63.0
43.7	87.00	65.0
51.0	100	72.0
65.0

(Brandes.)
2.67 pts. H₂O at 18.75°. (Abl.)
BaCl₂ is sol. in 2.86 pts. H₂O at 15.5°, and 1.67 boiling temp. (M. R. and P.)
H₂O at 15.5° dissolve 20 pts. BaCl₂, and 43.7°. (Ure's Dict.)

Solubility in 100 pts. H₂O at t°.

Pts. BaCl ₂	t°	Pts. BaCl ₂
31.1	77.5	51.9
33.9	95.65	57.7
41.2	102.5	58.9
47.7	105	59.7

(Nordenskiöld, Pogg. 136. 316.)

100 pts. H₂O dissolve pts. BaCl₂ at t°.

Pts. BaCl ₂	t°	Pts. BaCl ₂
33.2	50	43.7
38.1	58	45.9
40.0

(Gerardin, A. ch. (4) 5. 143.)
BaCl₂ + 2H₂O is sol. in 2.18 pts. H₂O, and the solution has sp. gr. = 1.2878. A. 109. 326.)
anhydrous BaCl₂ is sol. in 2.86 pts. H₂O at 15°. (Gerlach.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. BaCl ₂	t°	Pts. BaCl ₂	t°	Pts. BaCl ₂
0	30.9	36	39.7	71	49.7
1	31.2	37	40.0	72	50.0
2	31.5	38	40.2	73	50.3
3	31.7	39	40.5	74	50.6
4	31.9	40	40.7	75	50.9
5	32.2	41	41.0	76	51.2
6	32.4	42	41.3	77	51.5
7	32.6	43	41.6	78	51.8
8	32.8	44	41.9	79	52.1
9	33.1	45	42.2	80	52.4
10	33.3	46	42.5	81	52.7
11	33.5	47	42.7	82	53.0
12	33.8	48	43.0	83	53.3
13	34.0	49	43.3	84	53.6
14	34.2	50	43.6	85	54.0
15	34.5	51	43.9	86	54.3
16	34.7	52	44.2	87	54.6
17	35.0	53	44.4	88	55.0
18	35.2	54	44.7	89	55.3
19	35.5	55	45.0	90	55.6
20	35.7	56	45.3	91	55.9
21	36.0	57	45.6	92	56.2
22	36.2	58	45.9	93	56.6
23	36.5	59	46.2	94	56.9
24	36.7	60	46.4	95	57.2
25	37.0	61	46.7	96	57.6
26	37.2	62	47.0	97	57.9
27	37.5	63	47.3	98	58.2
28	37.7	64	47.6	99	58.5
29	38.0	65	47.9	100	58.8
30	38.2	66	48.2	101	59.2
31	38.5	67	48.5	102	59.5
32	38.7	68	48.8	103	59.8
33	39.0	69	49.1	104	60.2
34	39.2	70	49.4	104.1	60.3
35	39.5

(Mulder, calculated from his own and other observations. Scheik. Verhandel. 1864. 45.)

The saturated solution contains—
60.3 pts. BaCl₂ to 100 pts. H₂O, and boils at 104.1°. (Mulder.)
60.1 pts. BaCl₂ to 100 pts. H₂O, and boils at 104.4°. (Legrand.)
61.8 pts. BaCl₂ to 100 pts. H₂O, and boils at 104.5°. (Griffith.)
59.58 pts. BaCl₂ to 100 pts. H₂O, and boils at 105.48° (Gay-Lussac); at 106° (Kremers).
54.1 pts. BaCl₂ to 100 pts. H₂O, and forms crust at 104.4°; highest temperature observed, 104.9°. (Gerlach, Z. anal. 26. 426.)

Sat. BaCl₂ + Aq contains at:
100° 130° 144° 160° 180° 215°
36 37.3 37.5 38.9 40.7 43.1% BaCl₂
(Étard, A. ch. 1894, (7) 2. 535.)

Aq. solution contains 27.6% BaCl₂ at 30°. (Shreinemakers, C. C. 1910, I. 9.)

Solubility of $\text{BaCl}_2 + 2\text{H}_2\text{O}$ in H_2O equals 1.745 mol.-litre at 30° . (Masson, Chem. Soc. 1911, 99. 1136.)

$\text{BaCl}_2 + \text{Aq}$ sat. at 8° has sp. gr. 1.27. (Anthon.)
 $\text{BaCl}_2 + \text{Aq}$ sat. at 15° has sp. gr. 1.232. (Michel and Kraft.)

$\text{BaCl}_2 + \text{Aq}$ sat. at 18.1° has sp. gr. 1.235, and contains 44.31 pts. $\text{BaCl}_2 + 2\text{H}_2\text{O}$ to 100 pts. H_2O . (Karsten.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$ at 19.5° .

% BaCl_2	Sp. gr.	% BaCl_2	Sp. gr.
8.88	1.0760	27.53	1.2245
18.24	1.1521	35.44	1.2337

(Kremers, Pogg. 99. 444.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$ at 15° .

% BaCl_2	Sp. gr.	% BaCl_2	Sp. gr.
1	1.00917	14	1.13778
2	1.01834	15	1.14846
3	1.02750	16	1.15999
4	1.03667	17	1.17152
5	1.04584	18	1.18305
6	1.05569	19	1.19458
7	1.06554	20	1.20681
8	1.07538	21	1.21892
9	1.08523	22	1.23173
10	1.09508	23	1.24455
11	1.10576	24	1.25736
12	1.11643	25	1.27017
13	1.12711	

(Gerlach, Z. anal. 8. 283.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$ at 21.5° .

% $\text{BaCl}_2 + 2\text{H}_2\text{O}$	Sp. gr.	% $\text{BaCl}_2 + 2\text{H}_2\text{O}$	Sp. gr.
1	1.0073	16	1.1302
2	1.0147	17	1.1394
3	1.0222	18	1.1488
4	1.0298	19	1.1584
5	1.0374	20	1.1683
6	1.0452	21	1.1783
7	1.0530	22	1.1884
8	1.0610	23	1.1986
9	1.0692	24	1.2090
10	1.0776	25	1.2197
11	1.0861	26	1.2304
12	1.0947	27	1.2413
13	1.1034	28	1.2523
14	1.1122	29	1.2636
15	1.1211	30	1.2750

(Schiff, calculated by Gerlach, l.c.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$ at 18° .

% BaCl_2	Sp. gr.	% BaCl_2	Sp. gr.
5	1.0445	20	1.2047
10	1.0939	24	1.2559
15	1.1473	

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$ at 20° .

g. mols. BaCl_2 per l.	Sp. gr.
0.01	1.001878
0.025	1.00475
0.05	1.00929
0.075	1.01369
0.10	1.01766
0.25	1.0456
0.40	1.0726

(Jones and Pearce, Am. Ch. J. 1907, 38. 701.)

$\text{BaCl}_2 + \text{Aq}$ containing 6.94% BaCl_2 has sp. gr. $20^\circ/20^\circ = 1.0640$.

$\text{BaCl}_2 + \text{Aq}$ containing 11.38% BaCl_2 has sp. gr. $20^\circ/20^\circ = 1.1086$.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 279.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$ at 25° .

$\text{BaCl}_2 + \text{Aq}$	Sp. gr.
1-normal	1.0884
$\frac{1}{2}$ " "	1.0441
$\frac{1}{4}$ " "	1.0226
$\frac{1}{8}$ " "	1.0114

(Wagner, Z. phys. Ch. 1890, 5. 35.)

Sp. gr. of $\text{BaCl}_2 + \text{Aq}$.

t°	Concentration of $\text{BaCl}_2 + \text{Aq}$	Sp. gr.
25°	1 pt. BaCl_2 in 3.684 pts. H_2O	1.2194
22.8	1 " " " 52.597 " "	1.0145

(Hittorf, Z. phys. Ch. 1902, 39. 628.)

Temp. of Maximum Density.

Weight of BaCl_2 in 1000 grams H_2O	Temp. of maximum density	Molecular reduction of temp. of M. D.
0	3.982°	
6.73	3.207°	23.94
10.42	2.783°	23.88
20.83	1.572°	24.04
41.72	-0.843°	24.04

(De Coppet, C. R. 1897, 125. 533.)

$\text{BaCl}_2 + \text{Aq}$ containing 10% BaCl_2 boils at 100.6° . (Gerlach.)

$\text{BaCl}_2 + \text{Aq}$ containing 20% BaCl_2 boils at 101.9° . (Gerlach.)

B.-pt. of $\text{BaCl}_2 + \text{Aq}$ containing pts. BaCl_2 to 100 pts. H_2O . G = according to Gerlach (Z. anal. 26. 443); L = according to Legrand (A. ch. (2) 59. 452).

B.-pt.	G	L
100.5°	6.4	11.0
101.0	12.7	19.6
101.5	19.0	26.2
102.0	25.3	32.5
102.5	31.6	38.6
103.0	37.7	44.5
103.5	43.7	50.3
104.0	49.5	56.0
104.4	60.1
104.5	55.2

sol. in H_2O containing HCl than in H_2O , and scarcely sol. in conc. $HCl + Aq.$ (lus.)

ity of $BaCl_2$ in $HCl + Aq$ at 0° . $l_1 = no.$ $\frac{1}{2}$ mols. (in milligrammes) dissolved in 10 cc. of the liquid; $HCl = no.$ mols. (in milligrammes) contained in the same quantity of liquid.

l_1	HCl	Sum of mols.	Sp. gr.
5	0	29.45	1.250
	1.1	28.9	1.242
75	2.8	28.875	1.228
	5.0	28.4	1.210
	14.36	28.36	1.143
	18.775	28.975	1.118
7	22.75	29.42	1.099
4	32.0	34.74	1.079
9	50.5	50.79	1.088

(Engel, Bull. Soc. (2) 45. 653.)

n about 8000 pts. conc. $HCl + Aq.$ in about 20,000 pts. conc. $HCl + Aq.$ which HCl gas was passed. ically insol. in conc. $HCl + Aq$ conc. $\frac{1}{4}$ vol. ether. (Mar, Sill. Am. J. 143.)

Solubility in $HCl + Aq$ at 30° .

Composition of the solution	Solid phase
% by wt. $BaCl_2$	
27.6	$BaCl_2, 2H_2O$
12.97	"
3.85	"
0.46	"
0.00	"
0.00	$BaCl_2, 2H_2O + BaCl_2, H_2O$
0.00	$BaCl_2, H_2O$

(Schreinemakers, Z. phys. Ch. 1909, 68. 89.)

less sol. in $HNO_3 + Aq$ than in H_2O , $Ba(NO_3)_2$ is nearly insol. therein.

is sol. in about—
pts. H_2O .
pts. $NH_4OH + Aq$ (conc.).
pts. $NH_4OH + Aq$ (1 vol. conc.: 3 vols.)

pts. $HCl + Aq$ (1 vol. conc.: 4 vols.)

pts. $HC_2H_3O_2 + Aq$ (1 vol. commercial vol. H_2O).

pts. $NH_4Cl + Aq$ (1 pt. NH_4Cl : 10 pts.)

pts. $NH_4C_2H_3O_2 + Aq$ (dil. NH_4OH + realized by dil. $HC_2H_3O_2 + Aq$.)

pts. $NaC_2H_3O_2 + Aq$ (commercial $\frac{1}{2}$ neutralized by Na_2CO_3 , and dil. vol. H_2O).

pts. $Cu(C_2H_3O_2)_2 + Aq$. See Stolba (Z. 390).

pts. grape sugar (1 pt. grape sugar: H_2O). (Pearson, Zeit. Chem. 1869.)

$BaCl_2 + NH_4Cl$. Solubility of $BaCl_2$ in $NH_4Cl + Aq$ at 30° .

Composition of the solution		Solid phase
% NH_4Cl	% $BaCl_2$	
0	27.6	$BaCl_2, 2H_2O$
5.71	22.16	"
10.06	18.36	"
13.84	15.42	"
20.00	10.89	"
24.69	8.33	"
25.79	7.95	$BaCl_2, 2H_2O + NH_4Cl$
26.06	7.99	"
27.47	3.56	NH_4Cl
29.5	0	"

(Schreinemakers, Z. phys. Ch. 1909, 66. 688.)

See also under Ammonium chloride.

$BaCl_2 + Ba(OH)_2$. Solubility of $BaCl_2$ in $BaO + Aq$ at 30° .

Composition of the solution		Solid phase
% by wt. BaO	% by wt. $BaCl_2$	
0	27.6	$BaCl_2, 2H_2O$
1.78	27.42	"
1.79	27.31	$BaCl_2, 2H_2O + BaCl(OH), 2H_2O$
1.75	27.41	"
2.33	24.98	$BaCl(OH), 2H_2O$
2.50	24.20	"
3.27	21.46	"
4.67	19.18	"
4.86	18.97	$BaCl(OH), 2H_2O + BaO, 9H_2O$
4.29	18.83	"
4.64	18.77	"
4.65	18.10	"
4.62	18.04	$BaO, 9H_2O$
4.60	17.08	"
4.58	12.81	"
4.45	10.77	"
4.99	0	"

(Schreinemakers, Z. phys. Ch. 1909, 68. 88.)

Sol. in $CuCl_2, NH_4Cl + Aq$ at 30° . (Schreinemakers, Z. phys. Ch. 1909, 66. 688.)

The solubility data for the system $BaCl_2 + CuCl_2 + KCl + Aq$ have been determined at 40° and 60° . (Schreinemakers, C. C. 1915, I. 933.)

$BaCl_2 + HgCl_2$. Solubility of $BaCl_2 + HgCl_2$ in H_2O .

t°	Gms. per 100 g. solution		Solid phase
	$BaCl_2$	$HgCl_2$	
10.4°	23.58	50.54	$BaCl_2, 2H_2O + HgCl_2$
10.4	23.44	50.74	$BaCl_2, 3HgCl_2, 6H_2O$
10.4	22.58	51.23	
10.4	22.48	51.41	
10.4	22.10	51.66	
10.4	21.64	51.74	$BaCl_2, 2H_2O + HgCl_2$
25.0	23.02	54.83	

(Foote and Bristol, Am. Ch. J. 32. 248.)

Solubility of $\text{BaCl}_2 + \text{HgCl}_2$ in H_2O .

Temp. = 20°			Temp. = 0°		
% HgCl_2	% BaCl_2	Solid phase	% HgCl_2	% BaCl_2	Solid phase
0	27.77	$\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$	0	23.70	$\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$
2.90	27.56	"	14.25	24.0	"
7.09	27.47	"	36.20	24.89	"
12.98	26.99	"	46.12	24.07	$\text{BaCl}_2, 3\text{HgCl}_2 \cdot 6\text{H}_2\text{O} +$
22.61	26.89	"	46.05	24.03	" $\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$
34.57	26.10	"	46.07	24.05	"
46.50	25.22	"	46.59	23.28	$\text{BaCl}_2, 3\text{HgCl}_2 \cdot 6\text{H}_2\text{O}$
55.16	23.46	$\text{HgCl}_2 + \text{BaCl}_2 \cdot 2\text{H}_2\text{O}$	47.78	21.05	"
55.32	23.08	"	48.43	20.64	$\text{BaCl}_2, 3\text{HgCl}_2 \cdot 6\text{H}_2\text{O} + \text{HgCl}_2$
55.19	22.98	"	48.49	20.71	"
48.97	17.87	HgCl_2	44.33	18.50	HgCl_2
41.30	14.28	"	29.0	11.59	"
27.62	8.41	"	16.36	6.11	"
14.19	2.65	"	3.95	0	"
7.67	0	"			
			Temp. = 40°		
			56.57	22.98	$\text{BaCl}_2 \cdot 2\text{H}_2\text{O} + \text{HgCl}_2$

(Schreinemakers, Ch. Weekbl. 1911, 7. 202.)

$\text{BaCl}_2 + \text{KCl}$. Sol. in sat. $\text{KCl} + \text{Aq}$, at first without pptn. The KCl is pptd. after a time until a state of equilibrium is reached.

100 pts. H_2O at 16.6° dissolve 33.8–27.2 pts. KCl and 18.2–34.9 pts. BaCl_2 . (Kopp, A. 34. 267.)

100 g. sat. solution of $\text{BaCl}_2 + \text{KCl}$ contain 13.83 g. BaCl_2 and 18.97 g. KCl at 25°. (Foote, Am. Ch. J. 32. 253.)

$\text{BaCl}_2 + \text{Ba}(\text{NO}_3)_2$. BaCl_2 is sol. in sat. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$.

Solubility of $\text{BaCl}_2 + \text{Ba}(\text{NO}_3)_2$ in H_2O . Both salts present in solid phase.

t°	Gms. per 100 gms. solution		t°	Gms. per 100 gms. solution	
	BaCl_2	$\text{Ba}(\text{NO}_3)_2$		BaCl_2	$\text{Ba}(\text{NO}_3)_2$
0	22.5	4.3	100	31	II
20	24.5	6.0	140	32	20
40	26.5	7.5	180	33	26
60	28.5	9.5	210	32	32

(Etard, A. Ch. (7) 2. 535.)

Very slowly sol. in sat. $\text{NaNO}_3 + \text{Aq}$ with separation of $\text{Ba}(\text{NO}_3)_2$.

Rapidly sol. in sat. $\text{KNO}_3 + \text{Aq}$, forming $\text{Ba}(\text{NO}_3)_2$, which separates out. (Karsten.)

$\text{BaCl}_2 + \text{NaCl}$. BaCl_2 is sol. in $\text{NaCl} + \text{Aq}$ at first without separation of NaCl , which, however, finally separates.

100 pts. H_2O dissolve, when both salts are in excess—

	1	2	3	4	5	6
NaCl . . .	35.9	4.1		40.4	35.3	...
BaCl_2 . . .		34.5	35.0	..	19.4	60.3
		38.6			54.7	

1, 2, and 3 are at 17°. (Kopp, A. 34. 268.)

4, 5, and 6 are at b.-pt. (Mulder.)

Solubility of $\text{BaCl}_2 + \text{NaCl}$.

100 pts. H_2O dissolve pts. BaCl_2 and NaCl at t°.

t°	Pts. BaCl_2	Pts. NaCl	t°	Pts. BaCl_2	Pts. NaCl
III	4.1	33.9	60	9.7	33.5
20	4.1	33.8	70	11.7	33.0
30	5.0	33.7	80	13.9	33.0
40	6.3	33.6	90	15.9	33.6
50	7.9	33.5	100	17.9	33.6

(Precht and Wittgen, B. 14. 1667.)

Solubility of $\text{BaCl}_2 + \text{NaCl}$ in $\text{HCl} + \text{Aq}$ at 30°.

Solid phase, NaCl			Solid phase, $\text{BaCl}_2 + 2\text{H}_2\text{O}$		
Sp. gr. of sat. solution	G. mol. litre		Sp. gr. of sat. solution	G. mol. litre	
	HCl	NaCl		HCl	BaCl_2
1.2018	0.0000	5.400	1.3056	0.0000	1.745
1.1906	0.4575	4.932	1.2651	0.4709	1.468
1.1801	0.969	4.386	1.2147	1.107	1.122
1.1633	1.786	3.589	1.1789	1.622	0.861
1.1512	2.412	2.978	1.1419	2.234	0.592
1.1427	3.052	2.463	1.1068	3.041	0.307
1.1289	4.152	1.628	1.0880	3.953	0.124
1.1188	5.950	0.630	1.0895	3.059	0.020
1.1258	7.205	0.268	1.1024	6.234	0.00
			1.1609	10.25	0.00

(Masson, Chem. Soc. 1911, 99. 1136.)

ility of BaCl₂+NaCl in HCl+Aq at 30°.

l	% NaCl	% BaCl ₂	Solid phase
	23.85	3.8	NaCl, BaCl ₂ .2H ₂ O
4	18.07	2.27	"
2	9.55	0.82	"
0	4.65	0.29	"
6	1.54	0.00	"
6	0.47	0.00	"
1	0.12	0.00	NaCl+BaCl ₂ .H ₂ O

einemakers, Arch. Néer. Sc. ex. nat. (2) 15. 91.)

ol in liquid NH₃. (Franklin, Am. Ch. 38, 20. 827.)

lity in alcohol: 100 pts. alcohol of given sp. gr. dissolve pts. of the anhydrous, and crystallized lt.

Sp. gr.	Pts. BaCl ₂	Pts. BaCl ₂ +2H ₂ O
0.900	1.00	1.56
0.848	0.29	0.43
0.834	0.185	0.32
0.817	0.09	0.06

(Kirwan.)

sol. in abs. alcohol, or below 19° in al- of over 91%. Dil. alcohol dissolves less, than corresponds to the amount of H₂O nt. (Gerardin, A. ch. (4) 5. 142.)

ility in 100 pts. alcohol at t°. D=sp. gr. of alcohol; S=solubility.

9904	D=0.9848		D=0.9793		D=0.9726	
s	t°	s	t°	s	t°	s
29.1	14	25.0	11	19.6	15	15.6
32.0	32	29.1	15	20.4	23	17.0
33.5	39	30.9	20	21.7	33	19.1
37.4	50	33.2	35	24.6	50	22.0
39.8	63	37.6	45	26.8

.9573	D=0.9390		D=0.8967		D=0.8429	
s	t°	s	t°	s	t°	s
10	12	6.5	12	0.1	12	0.00
11.4	23	7.2	30	4.3	19	0.00
12.9	31	8.3	47	4.9	25	0.04
13.8	37	9.0	50	0.28
15.2	47	10.1	67	0.377

(Gerardin, A. ch. (4) 5. 142.)

ility in dil. alcohol of x% by weight at 15°.

cohol	0	10	20	30	40	60	80
BaCl ₂ .2H ₂ O	30.25	23.7	18.0	12.8	9.3	3.4	0.5

(Schiff, A. 118. 365.)

L in 6885-8108 pts. 99.3% alcohol at and in 1857 pts. at ebullition. (Frese-)

Solubility of BaCl₂ in alcohol+Aq.

t°	% alcohol	% BaCl ₂	Solid phase
30°	0	27.95	BaCl ₂ .2H ₂ O
"	32.67	10.63	"
"	50.16	5.68	"
"	66.72	2.23	"
"	92.53	0.05	"
"	94.83	0.07	BaCl ₂ .2H ₂ O+BaCl ₂ .H ₂ O
"	94.75	0.05	"
"	94.60	0.07	"
"	97.14	BaCl ₂ .H ₂ O
"	98.17	0.08	BaCl ₂ .H ₂ O+BaCl ₂
"	99.41	BaCl ₂
60°	0	31.57	BaCl ₂ .2H ₂ O
"	16.68	20.16	"
"	34.10	13.21	"
"	66.02	2.82	"
"	88.55	0.25	"
"	90.11	0.09	BaCl ₂ .2H ₂ O+BaCl ₂ .H ₂ O
"	90.39	"
"	93.95	BaCl ₂ .H ₂ O

(Schreinemakers and Massink, Chem. Weekbl. 1910, 7. 213.)

100 pts. absolute methyl alcohol dissolve 2.18 pts. BaCl₂ at 15.5°, and 7.3 pts. BaCl₂.2H₂O at 6°. (de Bruyn, Z. phys. Ch. 10. 783.)

At 15° C. 1 pt. by weight is sol. in:—

78 pts. methyl alcohol of sp. gr. 0.790
7,000 " ethyl " " " 0.8035
100,000 " propyl " " " 0.8085
(Rohland, Z. anorg. 1897, 15. 413.)

Absolutely insol. in boiling amyl alcohol. (Browning, Sill. Am. J. 144. 459.)

Absolutely insol. in acetic ether. (Cann, C. R. 102. 363.)

Very sl. sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

100 pts. by weight of glycerine dissolve 10 pts. BaCl₂ at 15.5°. (de Bruyn, Z. phys. Ch. 10. 783.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.) (Eidmann, C. C. 1899, II. 1014.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in anhydrous pyridine, 97% pyridine+Aq. and 95% pyridine+Aq. Sl. sol. in 93% pyridine+Aq. (Kahlenberg, J. Am. Chem. Soc. 1908, 30. 1107.)

+H₂O. Solution of monohydrate sat. at 6° contains 31.57% BaCl₂. (Schreinemakers, Chem. Weekbl. 1910, 7. 213.)

2.5 grams of the monohydrate are sol. in 100 cc. of methyl alcohol at 14°. (Kirschner, Z. phys. Ch. 1911, 76. 176.)

Exact solubility in methyl alcohol cannot be determined as BaCl₂+H₂O separates out from a sat. solution of the dihydrate. (Kirschner, Z. phys. ch. 1911, 76. 177.)

Barium cadmium chloride, BaCl₂,CdCl₂+4H₂O.

Easily sol. in H₂O. (v. Hauer.)

Solubility in H₂O at t°.

t°	100 pts. solution contain pts.			100 g. of solution contain g. salt	100 g. H ₂ O dissolve g. salt	100 mola. H ₂ O dissolve mola. of anhydrous salt
	Cl	Ba	Cd			
22.5	15.19	14.71	11.98	41.88	72.06	3.32
32.9	16.18	16.09	12.40	44.59	80.73	3.72
41.4	16.95	16.81	13.05	46.87	88.01	4.06
53.4	18.21	18.13	13.95	50.30	101.21	4.66
62.0	18.81	18.74	14.73	52.28	109.56	5.05
97.8	22.48	22.00	17.57	62.05	163.50	7.53
108.3	23.51	22.79	18.53	64.83	184.33	8.49
109.2	23.69	29.95	18.67	65.31	188.27	8.67

(Rimbach, B. 1897, 30. 3083.)

BaCl₂.2CdCl₂+5H₂O. Quite difficultly sol. in H₂O. (v. Hauer.)

Solubility in H₂O at t°.

t°	100 pts. by wt. of solution contain pts. by wt.			100 g. of solution contain g. salt	100 g. H ₂ O dissolve g. salt	100 mola. H ₂ O dissolve mola. of anhydrous salt
	Cl	Ba	Cd			
22.6	16.89	11.00	17.71	45.60	83.82	2.63
41.3	18.15	11.77	19.22	49.14	96.62	3.03
53.9	18.78	12.41	19.85	51.04	104.25	3.27
62.2	19.66	12.83	20.59	53.08	113.13	3.55
69.5	20.18	13.09	21.20	54.47	119.64	3.76
107.2	23.31	14.87	24.11	62.29	165.18	5.19
107.2	23.16	14.93	24.39	62.48	166.53	5.23

(Rimbach, B. 1897, 30. 3083.)

Barium mercuric chloride, basic, BaCl₂, HgO +6H₂O.

Decomp. by H₂O. (André, C. R. 104. 431.)

Barium mercuric chloride, BaCl₂, 2HgCl₂+2H₂O.

Efflorescent in dry air; sol. in H₂O. (v. Bonsdorff, Pogg. 17. 130.)

The salt BaCl₂, 2HgCl₂+2H₂O described by Bonsdorff does not form under the conditions which he gives. (Foote, Am. Ch. J. 1904, 32. 251.)

BaCl₂,3HgCl₂+6H₂O. Solubility determinations with mixtures of BaCl₂ and HgCl₂ show that these chlorides do not form a double salt at 25°, but that a transition temp. exists at about 17.2° below which the salt BaCl₂, 3HgCl₂+6H₂O forms. (Foote, Am. Ch. J. 1904, 32. 251.)

+8H₂O. Less sol. in H₂O than the Sr and Mg double salts. (Swan, Am. Ch. J. 1898, 20. 633.)

Barium rhodium chloride, 3BaCl₂, Rh₂Cl₄.

See Chlororhodite, barium.

Barium stannous chloride, BaCl₂, SnCl₂+4H₂O.

Sol. in H₂O. (Poggiale, C. R. 20. 1183.)

Barium stannic chloride.

See Chlorostannate, barium.

Barium uranium chloride, BaCl₂,UCl₄.

Decomp. by H₂O. (Aloy, Bull. Soc. 1899, (3) 21. 265.)

Barium zinc chloride, BaCl₂, ZnCl₂+4H₂O.

Deliquescent, and sol. in H₂O. (Warner, C. N. 27. 271.)

Pptd. from warm solution only. (Ephraim, Z. anorg. 1910, 67. 381.)

+2½H₂O. Pptd. from cold solution. (Ephraim.)

Barium chloride hydrazine, BaCl₂, 2N₂H₄.

Hydrosopic. (Franzen, Z. anorg. 1908, 60. 290.)

Barium chloride hydroxylamine, BaCl₂, 2NH₂OH.

Very sol. in H₂O. (Crismer, Bull. Soc. (3) 3. 118.)

Barium chloride sulphuric anhydride, BaCl₂, 2SO₃.

Decomp. by H₂O. (Schultz-Sellack, B. 4. 113.)

Barium chlorofluoride, BaClF.

Difficultly sol. in H₂O, but much more sol. than BaF₂. Decomp. by H₂O, so that when washed on filter, the filtrate contains more BaCl₂ than BaF₂. (Berzelius, Pogg. 1. 19.)

Insol. in and undecomp. by boiling alcohol; sol. in conc. HCl and HNO₃. Decomp. by hot H₂O, hot H₂SO₄, dil. acetic acid, dil. HCl or dil. HNO₃. (Defacqz, C. R. 1904, 138. 198.)

Barium cyanamide, BaCN₂.

Decomp. by H₂O. (Frank, C. C. 1902, 11. 774.)

subfluoride sodium fluoride, BaF,
p. by H₂O. (Guntz, C. R. 1903,
)
fluoride, BaF₂.
ly sol. in H₂O (Berzelius); less sol. in
CaF₂.
H₂O dissolves 1630 mg. BaF₂ at 18°. (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)
g. are contained in 1 l. of sat. solu-
3°. (Kohlrausch, Z. phys. Ch. 1908,
in molten MnCl₂, MnBr₂, MnI₂,
BaCl₂, MnBr₂+BaBr₂ and MnI₂+
Defacqz, A. ch. 1904, (8) 1. 350.)
sol. in HCl, HNO₃, or HF+Aq.
ssac and Thénard.)
in liquid HF. (Franklin, Z. anorg.
2.)
in ethyl acetate. (Naumann, B.
314.)
an aqueous solution of sodium cit-
piller.)
in (stannic) fluoride.
ostannate; barium.
tellurium fluoride, BaF₂, 2TeF₄.
p. by H₂O. (Högbom, Bull. Soc. (2)
titanium fluoride.
otitanate, barium.
itanyl fluoride, TiO₂F₂, BaF₂.
uoxypertitanate and fluoxytitanate,
ranyl fluoride.
oxyuranate, barium.
anadyl fluoride.
oxyvanadate, barium.
irconium fluoride, 3BaF₂, 2ZrF₄+
)
le precipitate. (Marignac.)
o Fluozirconate, barium.
luoiodide, BaF₂, BaI₂.
p. by H₂O, dil. HCl, dil. HNO₃, or
). Sol. in HI and HNO₃. Insol.
ndecomp. by boiling alcohol. De-
dil. acetic acid. (Defacqz, C. R.
1. 199.)
ydride, BaH.
p. by H₂O or HCl+Aq. (Winkler,
79.)
p. by H₂O. (Guntz, C. R. 1901,
)
ydrosulphide, BaS₂H₂.
sol. in H₂O. Insol. in alcohol.
) Sol. in H₂O, and the solution dis-
(Veley, Chem. Soc. 49. 369.)

Barium hydroxide, BaO₂H₂.
100 pts. cold H₂O dissolve 5 pts. BaO₂H₂.
boiling " 50 "
(Davy.)
100 pts. H₂O at 20° dissolve 3.45 pts. BaO.
(Bineau, C. R. 41. 509.)
100 pts. H₂O at 13° dissolve 2.86 pts. BaO.
" " 47° " 13.3 "
" " 70° " 17.9 "
(Osann.)
100 pts. H₂O dissolve pts. BaO at t°.

t°	Pts. BaO	t°	Pts. BaO	t°	Pts. BaO
0	1.5	30	5.0	60	18.76
5	1.75	35	6.17	65	24.67
10	2.22	40	7.36	70	31.9
15	2.89	45	9.12	75	56.85
20	3.48	50	11.75	80	90.77
25	4.19	55	14.71

(Rosenthal and Rühlmann, J. B. 1870. 314.)
100 pts. H₂O dissolve at 25° 55.08 millimols.
BaO₂H₂.
2 (Herz and Knoch, Z. anorg. 1904,
41. 315.)
Sp. gr. of BaO₂H₂+Aq.

%BaO	Sp. gr.	%BaO	Sp. gr.
30	1.6	1.8	1.02
19	1.3	0.9	1.01
2.6	1.03

(Dalton.)
Sp. gr. of BaO₂H₂+Aq at 18° containing
1.25% BaO₂H₂=1.0120; containing 2.5%=
1.0253. (Kohlrausch, W. Ann. 1879, 6. 41.)
Sp. gr. of BaO₂H₂+Aq at 80°.

Sp. gr.	%BaO ₂ H ₂ by volume	%BaO ₂ H ₂ by weight	Sp. gr.	%BaO ₂ H ₂ by volume	%BaO ₂ H ₂ by weight
1.514	58.22	38.45	1.219	24.53	20.12
1.500	56.31	37.54	1.200	23.00	19.17
1.479	54.14	36.60	1.195	22.15	18.53
1.458	49.38	33.87	1.174	19.83	16.89
1.450	48.90	33.72	1.152	17.78	15.43
1.413	45.99	32.55	1.129	16.01	14.18
1.400	45.00	32.14	1.125	15.80	14.04
1.390	44.22	31.81	1.114	14.56	13.07
1.375	42.40	30.84	1.100	13.06	11.87
1.368	41.45	30.30	1.076	10.58	9.83
1.350	38.60	28.59	1.062	9.16	8.62
1.338	37.30	27.88	1.049	7.55	7.20
1.312	35.02	26.69	1.040	6.51	6.26
1.301	34.02	26.13	1.031	5.18	5.02
1.278	31.48	24.67	1.022	4.78	4.67
1.249	28.14	22.52	1.015	3.90	3.84
1.236	26.41	21.36	1.009	3.37	3.34

(Haff, C. N. 1902, 86. 284.)
Insol. in liquid NH₃. (Franklin, Am. Ch.
J. 1898, 20. 827.)
More sol. in NaCl+Aq, KNO₃+Aq, or
NaNO₃+Aq than in H₂O. (Karsten.)
Not precipitated by alcohol.

Sol. with combination in absolute alcohol and anhydrous methyl alcohol. Insol. in ether.

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Solubility in acetone+Aq at 25°.

A=cc. acetone in 100 cc. acetone+Aq.
 $\frac{\text{BaO}_2\text{H}_2}{2}$ =millimols. BaO₂H₂ in 100 cc. of the solution.

S=sp. gr. of the solution.

A	$\frac{\text{BaO}_2\text{H}_2}{2}$	S
0	55.08	1.04790
10	31.84	1.01677
20	17.79	0.99268
30	9.10	0.97630
40	4.75	0.95605
50	1.54	0.93980
60	0.48	0.91790
70	0.08	0.89562

(Herz, Z. anorg. 1904, 41. 321.)

BaO₂H₂ is sol. in an aqueous solution of cane sugar (Hunton, Phil. Mag. (3) 11. 156); also in an aqueous sol. of mannite (Favre, A. ch. (3) 11. 76); sorbine (Pelouze); hot solution of quercite, separating on cooling (Des-saignes).

+3H₂O. Decomp. by H₂O free from carbonic acid. Sl. sol. in alcohol and ether. (Bauer, Z. anorg. 1905, 47, 416.)

Solubility in H₂O the same as that of the comp. with 8H₂O. Insol. in alcohol and ether. (Bauer, Zeit. angew. Ch. 1903, 17. 341.)

Nearly insol. in alcohol and ether. (Bauer, Zeit. angew. Ch. 1903, 16. 349.)

+8H₂O. Sol. in 20 pts. cold, and 3 pts. boiling H₂O (Graham); 17.5 pts. H₂O at 15.5°, and in all proportions of hot H₂O. (Hope.) Sol. in 19 pts. H₂O at 15°, and 2 pts. at 100°. (Wittstein.)

If BaO₂H₂+8H₂O is heated it dissolves in the crystal H₂O and the solution has the following bpts.

%BaO	49.05	50.05	52.43	53.72
B.-pt.	103°	104°	105°	106°

%BaO	55.35	57.49	58.74	61.44
B.-pt.	107°	108°	108.5°	109°

BaO₂H₂+3H₂O separates at 109°. (Bauer, Zeit. angew. Ch. 1903, 17. 345.)

B.-pt. of BaO₂H₂.8H₂O+Aq at 732 mm.

Bpt.	Time	%BaO
78° (mpt.)	0	48.45
78	4'	48.45
103	6' 30"	49.05
104	6' 45"	50.05
105	7' 30"	52.43
106	9' 25"	53.72
107	10' 45"	55.35

B.-pt. of BaO₂H₂.8H₂O+Aq, etc.—*Continued*

Bpt.	Time	%BaO
108	12'	57.49
108.5	—	58.74
109	13'	61.44
109	17' 40"	63.65
108	17' 50"	66.53
105	18'	67.51
100	18' 45"	68.17

(Bauer, Z. anorg. 1905, 47. 407.)

Solubility in Ba(NO₃)₂+Aq at 25°. Solution sat. with respect to both Ba(NO₃)₂ and BaO₂H₂.8H₂O.

Sp. gr. 25°/25°	g. BaO as Ba(OH) ₂ in 100 g. H ₂ O	g. Ba (NO ₃) ₂ in 100 g. H ₂ O
1.1448	5.02	11.48
1.1371	4.93	10.21
1.1288	4.83	8.66
1.1220	4.72	7.55
1.1133	4.72	7.01
1.1062	4.65	6.82
1.1044	4.61	6.55
1.1010	4.64	6.08
1.0975	4.60	5.66
1.0949	4.55	5.46
1.0937	4.54	5.32
1.0885	4.52	4.44
1.0864	4.53	4.41
1.0840	4.52	4.04
1.0790	4.48	3.47
1.0774	4.46	3.14
1.0731	4.40	2.79
1.0711	4.42	2.53
1.0651	4.35	1.88
1.0626
1.0640	4.35	1.45
1.0538	4.29	0.43
1.0512	4.29	0

(Parsons and Corson, J. Am. Chem. Soc. 1910, 32. 1385.)

Solubility of Ba(OH)₂+8H₂O (solid phase) in MCl+Aq (mol. per litre of solution) at 25°.

Solution of	(Cl')	(OH')
LiCl	0	0.555
"	0.75	0.745
"	1.42	0.937
"	2.30	1.336
KCl	0.86	0.645
"	1.75	0.660
"	3.40	0.676
NaCl	0	0.555
"	0.73	0.630
"	1.43	0.699
"	2.82	0.806
RbCl	1.25	0.648

(Herz, Z. anorg. 1910, 67. 366.)

solubility of BaO in NaOH+Aq at 30°.

BaO	%BaO	Solid phase
	4.99	BaO. 9H ₂ O
8	1.29	"
3	0.89	"
3	0.57	"
2	0.53	"
7	0.47	"
3	1.06	"
3	1.87	BaO. 9H ₂ O + BaO. 4H ₂ O
1	1.84	BaO. 4H ₂ O
2	1.75	"
3	1.58	"
1	1.34	BaO. 4H ₂ O + BaO. 2H ₂ O
2	0.82	BaO. 2H ₂ O
2	0.59	"
3	0.57	BaO. 2H ₂ O + NaOH. H ₂ O
	0	NaOH. H ₂ O

reinemakers, Z. phys. Ch. 1909, 68. 84.)

% alcohol dissolves less than 0.5% of rt. of BaO₂.H₂+8H₂O. (Beckmann, J. 883, (2) 27. 138.)

in subiodide sodium iodide, BaI, NaI. comp. by H₂O. (Guntz, C. R. 1903, 750.)

in iodide, BaI₂.
not deliquescent. Very sol. in H₂O and sol. 100 pts. of anhydrous salt dissolve:
19.5° 30° 40° 60° 90° 106°
48 44 43 41 37 35 pts. H₂O.
(Kremers, Pogg. 103. 66.)

gr. of BaI₂+Aq containing:
10 15 20 25 30%BaI₂
5 1.091 1.143 1.201 1.265 1.333
40 45 50 55 60%BaI₂
2 1.495 1.596 1.704 1.825 1.970
mers, Pogg. 111. 63, calculated by Gerlach, Z. anal. 8. 279.)

sily sol. in alcohol. (Henry.)
sol. in benzonitrile. (Naumann, B. 47. 1369.)
l. in acetone. (Naumann, B. 1904, 37. ; Eidmann, C. C. 1899, II. 1014.)
l. in methyl acetate. (Naumann, B. 42. 3789.)
2H₂O. At 15° C., 1 pt. by weight in sol. in:
22 pts. methyl alcohol sp. gr. 0.790
93 " " " " 0.8035
07 " " " " 0.8085
(Rohland, Z. anorg. 1897, 15. 413.)

7H₂O. (Thomson, B. 10. 1343.)

the composition of the hydrates formed BaI₂ at different dilutions is calculated determinations of the lowering of the produced by BaI₂ and of the conduc- and sp. gr. of BaI₂+Aq. (Jones, Am. 1. 1905, 34. 306.)

Barium iodide, basic, Ba(OH)I+9H₂O.

See Barium oxyiodide.

Barium bismuth iodide, BaI₂, 2BiI₃+18H₂O.

Deliquescent; decomp. by H₂O. (Linau, Pogg. 111. 240.)

Barium cadmium iodide, BaI₂, CdI₂+5H₂O.

Deliquescent. (Croft.)

Barium mercuric iodide, BaI₂, 2HgI₂.

Decomp. by much H₂O. (Boullay.)

BaI₂, HgI₂. Sol. in H₂O. (Boullay.)

Sp. gr. of sat. solution = 3.575-3.588. (Rohrbach, W. Ann. 20. 169.)

+5H₂O. (Duboin, C. R. 1906, 143. 314.)

2BaI₂, 3HgI₂+16H₂O. (Duboin, C. R. 1906, 142. 888.)

BaI₂, 5HgI₂+8H₂O. As the corresponding Ca salt. (Duboin, C. R. 1906, 142. 888.)

3BaI₂, 5HgI₂+21H₂O. Very deliquescent. (Duboin, C. R. 1906, 142. 889.)

Barium stannous iodide.

Very sol. in H₂O. (Boullay.)

Barium zinc iodide, BaI₂, 2ZnI₂.

Deliquescent, and sol. in H₂O. (Rammelsberg.)

+4H₂O. Very hygroscopic. (Ephraim, Z. anorg. 1910, 67. 385.)

Barium nitride, Ba₃N₂.

Decomp. H₂O violently, not alcohol. (Maquenne, A. ch. (6) 29. 219.)

BaN₆.

See Barium azoimide.

Barium oxide, BaO.

Sol. in H₂O with evolution of heat.

Easily sol. in dil. HNO₃, or HCl+Aq.

Solubility in NaOH+Aq. See Barium hydroxide.

Solubility in Na₂O, HCl, +H₂O at 30°. (Schreinemakers, Z. phys. Ch. 1909, 68. 98.)

Solubility in Na₂O, NaCl, BaCl₂+Aq at 30°. (Schreinemakers.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 827.)

Sol. with combination in absolute alcohol and anhydrous wood-spirit. Insol. in ether.

Easily sol. in absolute methyl alcohol.

1 l. absolute ethyl alcohol sat. with BaO at 9° contains 213.8 g. BaO. (Berthelot, Bull. Soc. 8. 389.)

Sol. in methyl alcohol. (Neuberg and Neimann, Biochem. Z. 1906, 1. 173.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

See also Barium hydroxide.

Barium peroxide, BaO₂.

Insol. in H₂O; decomp. by boiling H₂O.

Sol. in acids with formation of hydrogen dioxide.

Forms hydrate with $8\text{H}_2\text{O}$; also $10\text{H}_2\text{O}$ (Berthelot, A. ch. (5) 21. 157); also a compound $\text{BaO}_2 \cdot \text{H}_2\text{O}$, which is very unstable, sl. sol. in cold H_2O , and insol. in alcohol or ether. (Schöne, A. 192. 257.)

$+8\text{H}_2\text{O}$. 100 cc. pure H_2O dissolve 0.168 g. $\text{BaO}_2 \cdot 8\text{H}_2\text{O}$; if H_2O contains 0.3 g. $\text{Ba}(\text{OH})_2 \cdot 8\text{H}_2\text{O}$, only 0.102 g. $\text{BaO}_2 \cdot 8\text{H}_2\text{O}$ are dissolved; if 0.6 g. $\text{Ba}(\text{OH})_2 \cdot 8\text{H}_2\text{O}$ only 0.019 g. $\text{BaO}_2 \cdot 8\text{H}_2\text{O}$ are dissolved. (Schöne, A. 1878, 192. 266.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Barium oxybromide, $\text{Ba}(\text{OH})\text{Br} + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Beckmann, J. pr. (2) 27. 132.)

$\text{BaBr}_2 \cdot \text{BaO} + 5\text{H}_2\text{O}$. Sl. sol. in H_2O . (Tassilly, C. R. 1895, 120. 1340.)

Barium oxychloride, $\text{Ba}(\text{OH})\text{Cl} + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Beckmann, J. pr. (2) 26. 388, 474.)

Barium mercury oxychloride, $\text{BaCl}_2 \cdot \text{HgO} + 6\text{H}_2\text{O}$.

Decomp. by H_2O . (André, C. R. 104. 431.)

Barium oxyiodide, $\text{Ba}(\text{OH})\text{I} + 9\text{H}_2\text{O}$.

Decomp. by H_2O and alcohol. (Beckmann, B. 14. 2154.)

$\text{BaI}_2 \cdot \text{BaO} + 9\text{H}_2\text{O}$. Sl. sol. in H_2O . (Tassilly, C. R. 1895, 120. 1340.)

**Barium oxysulphides, $\text{Ba}_2\text{O}_4\text{S}_3 + 58\text{H}_2\text{O}$,
 $\text{Ba}_2\text{OS} + 10\text{H}_2\text{O}$, $\text{Ba}_4\text{OS}_2 + 28\text{H}_2\text{O}$.**

Very unstable; decomp. by recrystallization into BaS_2H_2 and BaO_2H_2 .

Barium phosphide, BaP_2 .

Decomp. by H_2O . (Dumas, A. ch. 32. 364.)

Ba_3P_2 . Crystallized. Sol. in dil. acids; insol. in conc. acids; decomp. by H_2O . Insol. in organic solvents at ord. temp. (Jaboin, C. R. 1899, 129. 765.)

Barium selenide, BaSe .

Sol. in H_2O with decomp.

Sl. sol. in H_2O . (Favre, C. R. 102. 1469.)

Barium silicide, Ba_2Si .

(Jüngst, C. C. 1905, I. 195.)

BaSi_2 . Slowly decomp. by H_2O , not by $\text{NH}_4\text{OH} + \text{Aq}$. Rapidly decomp. by conc. NaOH . Sol. in HNO_3 , H_2SO_4 , and H_3PO_4 with evolution of spontaneously inflammable gas. Sol. in HF and HCl . Sol. in acetic acid without evolution of gas. (Moissan, Traité ch. min. 1904, III. 680.)

Decomp. rapidly in both hot and cold H_2O . (Bradley, C. N. 1900, 82. 150.)

Barium sulphide, BaS .

Sol. in H_2O with decomp.

Crystallized. Decomp. by H_2O .

Attacked by cold conc. HNO_3 . (Mourlot, A. ch. 1899, (7) 17. 521.)

Cryst. modification is less readily acted on by air and other reagents than the amorphous modification; sol. in fuse oxidizing agents. (Mourlot, C. R. 1898, 126. 645.)

$+ \text{H}_2\text{O}$. (Neuberg and Neimann, Biochem. Z. 1906, 1. 174.)

$+ 6\text{H}_2\text{O}$. Slowly sol. in boiling H_2O , with decomp.; insol. in, but decomp. by boiling alcohol. (Schöne.)

Barium sulphide, $\text{Ba}_4\text{S}_7 + 25\text{H}_2\text{O}$ (?).

Sol. in H_2O . (Schöne, Pogg. 112. 215.)

Barium trisulphide, BaS_3 .

Sol. in large amount of boiling H_2O . (Schöne, Pogg. 112. 215.)

Barium tetrasulphide, $\text{BaS}_4 + \text{H}_2\text{O}$.

Easily sol. in H_2O , especially if hot; sol. in 2.42 pts. H_2O at 15° ; insol. in CS_2 or alcohol. (Schöne, Pogg. 112. 224.)

$+ 2\text{H}_2\text{O}$. (Velej, Chem. Soc. 49. 369.)

Barium pentasulphide, BaS_5 .

Known only in solution.

Barium mercuric sulphide, $\text{BaS} \cdot \text{HgS} + 5\text{H}_2\text{O}$.

Sol. in H_2O . (Wagner, J. pr. 98. 23.)

Barium nickel sulphide, $\text{BaS} \cdot 4\text{NiS}$.

Sol. in warm conc. HCl . (Bellucci, C. A. 1909, 293.)

Barium stannic sulphide.

See Sulphostannate, barium.

Barium uranyl sulphide, $6\text{BaS} \cdot \text{UO}_2\text{S} + x\text{H}_2\text{O}$ (?).

Decomp. by $\text{HCl} + \text{Aq}$. (Remelé, Pogg. 124. 159.)

Baryta.

See Barium oxide, BaO .

Beryllium, Be .

For beryllium and its salts, see Glucinum and the corresponding salts.

Bismuth, Bi .

Not attacked by H_2O . Very slowly attacked by $\text{HCl} + \text{Aq}$ (Troost). Very sl. sol. in conc. $\text{HCl} + \text{Aq}$ (Schützenberger, Willm). Not attacked by dil. $\text{HCl} + \text{Aq}$ (Naquet and Hanriot). Very slowly attacked by cold $\text{HCl} + \text{Aq}$ (Godeffroy). According to very careful experiments pure Bi is absolutely unattacked by hot or cold, dil. or conc. $\text{HCl} + \text{Aq}$ except in presence of oxygen. (Ditte and Metzner, A. ch. (6) 29. 397.)

Not attacked by dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Decomp.

by hot conc. H_2SO_4 . Easily sol. in dil. or conc. HNO_3 + Aq, or aqua regia.

Not attacked by pure HNO_3 + Aq of 1.52 to 1.42 sp. gr. at 20° ; violently attacked by a more dil. acid, but the acid becomes concentrated thereby. Conc. HNO_3 + Aq attacks only by heating or adding NO_2 . (Millon, A. ch. (3) 6. 95.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0091 g. Bi in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Bismuth arsenide, Bi_2As_3 .

(Descamp, C. R. 86. 1065.)

Bismuth dibromide, Bi_2Br_4 .

Not known in a pure state. (Weber, Pogg. 107. 599.)

Bismuth tribromide, BiBr_3 .

Very deliquescent. Decomp. by H_2O . Sol. in alcohol or ether.

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913, 84. 27.)

Bismuth hydrogen bromide, BiBr_3 , $2\text{HBr} + 4\text{H}_2\text{O}$.

Deliquescent.

Decomp. in the air. (Aloy, Bull. Soc. 1906, (3) 35. 398.)

Bismuth caesium bromide, 2BiBr_3 , 3CsBr .

Ppt. Insol. in HBr .

Sol. in HCl and in HNO_3 . (Hutchins, J. Am. Chem. Soc. 1907, 29. 33.)

Bismuth potassium bromide, BiBr_3 , 2KBr .

Decomp. by H_2O . (Aloy, Bull. Soc. 1906, (3) 35. 398.)

Bismuth bromide ammonia, BiBr_3 , 3NH_3 .

Sol. in HCl + Aq.

BiBr_3 , 2NH_3 (?).

2BiBr_3 , 5NH_3 . Not deliquescent; not decomp. by H_2O ; easily sol. in dil. acids. (Muir, Chem. Soc. 29. 144.)

Bismuth bromide potassium chloride,

$\text{K}_2\text{BiCl}_3\text{Br}_2 + 1\frac{1}{2}\text{H}_2\text{O}$.

Decomp. by H_2O . (Atkinson, Chem. Soc. 43. 289.)

Bismuth dichloride, Bi_2Cl_4 .

Very deliquescent. Decomp. by H_2O , dil. acids, or conc. NH_4Cl + Aq. (Weber, Pogg. 107. 596.)

Bismuth trichloride, BiCl_3 .

Deliquescent. Decomp. by H_2O . Sol. in dil. HCl + Aq, and alcohol. Not decomp. by H_2O in presence of citrates. (Spiller.)

0.08 g. sol. in 100 ccm. liquid H_2S . (Antony, C. C. 1905, I. 1692.)

Moderately sol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

1 g. BiCl_3 is sol. in 5.59 g. acetone at 18° . Sp. gr. of sat. solution $18^\circ/4^\circ = 0.9194$. (Naumann, B. 1904, 37. 4331.)

Sol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

1 pt. is sol. in 60.36 pts. ethyl acetate at 18° . Sp. gr at $18^\circ/40^\circ = 0.9106$. (Naumann, B. 1910, 43. 320.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Bismuth chloride, Bi_2Cl_6 (?).

Decomp. by H_2O . (Dehérain, C. R. 54. 724.)

Bismuth hydrogen chloride, 2BiCl_3 , $\text{HCl} + 3\text{H}_2\text{O}$.

Not deliquescent. Decomp. by H_2O . (Engel, C. R. 106. 1797.)

BiCl_3 , 2HCl . (Jacquelin, A. ch. (2) 62. 363.)

Bismuth caesium chloride, BiCl_3 , 3CsCl .

Decomp. by H_2O . Sl. sol. in cold dil. HCl + Aq, but easily sol. on warming. (Brigham, Am. Ch. J. 14. 181.)

2BiCl_3 , 3CsCl . As above. (Brigham.)

BiCl_3 , 6CsCl . Easily sol. in H_2O and dil. HCl + Aq. (Godeffroy, B. 8. 9.)

Does not exist. (Brigham.)

Bismuth hydrazine chloride, BiCl_3 , $3\text{N}_2\text{H}_4\text{HCl}$.

Sol. in acids, from which it is pptd. by H_2O . (Ferratini, C. A. 1912. 1613.)

Bismuth nitrosyl chloride, BiCl_3 , NOCl .

Very deliquescent. Decomp. by H_2O . (Sudborough, Chem. Soc. 59. 662.)

Bismuth potassium chloride, BiCl_3 , $\text{KCl} + \text{H}_2\text{O}$.

Decomp. by H_2O . Cannot be recryst. except from conc. BiCl_3 + HCl . Decomp. by HCl + Aq into BiCl_3 , $2\text{KCl} + 2\text{H}_2\text{O}$. (Brigham, Am. Ch. J. 14. 167.)

BiCl_3 , 2KCl . Decomp. by H_2O . (Arppe, Pogg. 64. 37.)

Deliquescent.

Sol. in H_2O with decomp. into the oxychloride when excess H_2O is used. (Aloy, Bull. Soc. 1906, (3) 35. 397.)

+ $2\text{H}_2\text{O}$. Decomp. by H_2O . (Jacquelin, J. pr. 14. 1.)

Sol. in moderately conc. HCl + Aq.

BiCl_3 , 3KCl . Decomp. by H_2O . (Arppe.) Does not exist. (Brigham.)

Bismuth rubidium chloride, BiCl_3 , $\text{RbCl} + \text{H}_2\text{O}$.

Decomp. by H_2O ; sol. in dil. $\text{HCl} + \text{Aq}$, from which BiCl_3 , 3RbCl crystallizes. (Brigham, Am. Ch. J. 14. 174.)

BiCl_3 , 3RbCl . Decomp. by H_2O ; sol. in dil. $\text{HCl} + \text{Aq}$ without decomp. (Brigham.)

BiCl_3 , 6RbCl . Decomp. by H_2O ; sol. in $\text{HCl} + \text{Aq}$ (Godeffroy, B. 8. 9); does not exist. (Brigham.)

10BiCl_3 , 23RbCl (?). As above. (Brigham.)

Bismuth sodium chloride, BiCl_3 , $2\text{NaCl} + \text{H}_2\text{O}$.

+ $3\text{H}_2\text{O}$. Decomp. by H_2O . (Arppe, Pogg. 64. 237.)

BiCl_3 , 3NaCl .

Bismuth thallous chloride, BiCl_3 , 3TlCl .

Ppt. (Ephraim, Z. anorg. 1909, 61. 254.)

BiCl_3 , 6TlCl . Ppt. (Ephraim.)

Bismuth chloride ammonia, 2BiCl_3 , NH_3 .

Stable. (Dehérain, C. R. 54. 724.)

BiCl_3 , 2NH_3 . (D.)

BiCl_3 , 3NH_3 . (D.)

Bismuth chloride nitric oxide, BiCl_3 , NO .

Very hygroscopic. (Thomas, C. R. 1895, 121. 129.)

Bismuth chloride nitrogen peroxide, BiCl_3 , NO_2 .

Decomp. by moist air, but stable in dry air. (Thomas, C. R. 1896, 122. 612.)

Bismuth chloride selenide.

See Bismuth selenochloride.

Bismuth trifluoride, BiF_3 .

Insol. in H_2O or alcohol. (Gott and Muir, Chem. Soc. 53. 138.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

Bismuth hydrogen fluoride, BiF_3 , 3HF .

Deliquescent. Decomp. by boiling H_2O . (Muir, Chem. Soc. 39. 21.)

Bismuth gold, Au_3Bi .

Insol. in equal pts. of HNO_3 and tartaric acids. (Roessler, Z. anorg. 1895, 9. 71.)

Bismuthous hydroxide, $\text{Bi}(\text{OH})_3$.

Sol. in strong acids. Insol. in solutions of alkalis, alkali carbonates, $(\text{NH}_4)_2\text{CO}_3$, or NH_4NO_3 ; or of amyl amine (Wurtz). When recently pptd. is sol. in $\text{NH}_4\text{Cl} + \text{Aq}$, but insol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$ (Brett, 1837). Not pptd. in presence of Na citrates (Spiller).

Solubility of freshly pptd. $\text{Bi}(\text{OH})_3$ in $\text{NaOH} + \text{Aq}$.

g. NaOH per l.	g. Bi dissolved per l. at 20°	g. Bi dissolved per l. at 100°
400	0.16	1.70
320	0.11	1.20
240	0.11	...
200	0.10	0.5
160	0.08	0.5
120	0.07	...
80	0.04	0.35
40	trace	0.2
20	0	0.15

(Moser, Z. anorg. 1909, 61. 386.)

Solubility of freshly pptd. $\text{Bi}(\text{OH})_3$ in $\text{KOH} + \text{Aq}$.

KOH per l. g.	g. Bi dissolved per l. at 20°	g. Bi dissolved per l. at 100°
560	0.14	1.65
448	0.11	1.20
336	0.11	...
280	0.10	0.5
224	0.08	0.5
168	0.06	...
112	0.03	0.3
56	trace	0.2
28	0	0.15

(Moser, Z. anorg. 1909, 61. 386.)

Bi_2O_3 , $2\text{H}_2\text{O}$.

Bi_2O_3 , H_2O . (Muir, Chem. Soc. 32. 131.)

See also Bismuth trioxide.

Bismuth tetrahydroxide, Bi_2O_4 , H_2O .

Bi_2O_4 , $2\text{H}_2\text{O}$. (Wernicke, Pogg. 141. 109.)

Bismuthic hydroxide (Bismuthic acid), Bi_2O_4 , H_2O .

Insol. in H_2O ; easily decomp. by acids. (Fremy, A. ch. (3) 12. 495.) Decomp. by H_2SO_4 ; not attacked by $\text{SO}_2 + \text{Aq}$; neither dissolved nor decomp. by dil. $\text{HNO}_3 + \text{Aq}$, but slowly converted into an allotropic modification (?). Partially decomp. by conc. HNO_3 . Slowly but wholly dissolved by hot conc. HNO_3 . Sl. sol. in conc. $\text{KOH} + \text{Aq}$. (Arppe.)

Sol. in about 100 pts. boiling $\text{KOH} + \text{Aq}$, so conc. that it solidifies on removing the lamp. (Muir, Chem. Soc. 51. 77.)

Bi_2O_4 , $2\text{H}_2\text{O}$. (Bödeker, A. 123. 61.)

Does not exist. (Hoffmann and Geuther.)

Bismuth iodide, BiI_3 .

Not attacked by cold H_2O , but by boiling, BiOI is formed. 100 pts. absolute alcohol dissolve $3\frac{1}{2}$ pts. salt at 20° . (Gott and Muir, Chem. Soc. 57. 138.)

Sol. in HNO_3 , and $\text{HI} + \text{Aq}$, from which it is repptd. by H_2O or alcohol. Sol. in $\text{KI} + \text{Aq}$ or $\text{KOH} + \text{Aq}$. (Rammelsberg.)

1. in liquid NH_3 . (Franklin, Am. Ch. 20. 827.)

absolute alcohol dissolve 3.5 g. BiI_3 . (Gott and Muir, Chem. Soc. 67. 138.)
in acetone. (Naumann, B. 1904, 87.)

pts. methylene iodide dissolve 0.15 pt. BiI_3 at 12° , and very little more at higher temperatures. (Retgers, Z. anorg. 3. 343.)
in methyl acetate. (Naumann, B. 2. 3790.)

1 hydrogen iodide, BiI_3 , $\text{HI} + 4\text{H}_2\text{O}$. (Pogg. 44. 248.)

1 cesium iodide, $3\text{CsI} \cdot 2\text{BiI}_3$.
sl. sol. in H_2O . (Wells, Am. J. Sci. 10. 3. 464.)

1 calcium iodide, 2BiI_3 , $\text{CaI}_2 + 18\text{H}_2\text{O}$.
luculent; decomp. by H_2O . (Linau, 11. 240.)

1 magnesium iodide, 2BiI_3 , $\text{MgI}_2 + 3\text{H}_2\text{O}$.
luculent; decomp. by H_2O . (Linau, 11. 240.)

1 potassium iodide, BiI_3 , 4KI .
(Arppe, Pogg. 44. 237.)
 3KI . (Astre, C. R. 110. 1137.)
 2KI . Sol. in acetic ether. (Astre.)
 H_2O . Sol. in small amt. H_2O without
decomp. by much H_2O .
 2KI , HI . (Arppe.)
 $3\text{KI} + 2\text{H}_2\text{O}$. (Astre.)
 $\text{KI} + \text{H}_2\text{O}$. Decomp. by H_2O . (Nicklès,
1. 1097.)
 KI . Sol. in acetic ether. (Astre.)

1 sodium iodide, BiI_3 , $\text{NaI} + \text{H}_2\text{O}$.
luculent; decomp. by H_2O . (Nicklès,
1. 1097.)
 $3\text{NaI} + 12\text{H}_2\text{O}$. As above. (Linau,
11. 240.)

1 zinc iodide, 2BiI_3 , $\text{ZnI}_2 + 12\text{H}_2\text{O}$.
deliquescent. (Linau, Pogg. 111. 240.)

1 iodide ammonia, BiI_3 , 3NH_3 .
mp. by H_2O . (Rammelsberg.)

1 iodide zinc bromide.
in H_2O . (Linau, Pogg. 111. 240.)

1 nitride.
white. (Fischer, B. 1910, 43. 1471.)
Ppt. Decomp. by H_2O or dil. acids.
in, J. Am. Chem. Soc. 1905, 27. 847.)

1 dioxide, Bi_2O_3 .
in conc. $\text{HNO}_3 + \text{Aq}$. Decomp. by
acids, and boiling $\text{KOH} + \text{Aq}$.
mp. by H_2O . (Tanatar, Z. anorg.
7. 488.)

Bismuth trioxide, Bi_2O_3 .

Insol. in H_2O . Sol. in conc. acids.

Solubility of Bi_2O_3 in $\text{HNO}_3 + \text{Aq}$ at 20° .

In 100 g. of the liquid phase		Solid phase
g. Bi_2O_3	g. N_2O_5	
0.321	0.963	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot 2\text{H}_2\text{O}$
0.337	0.982	"
3.54	4.68	"
6.37	7.17	"
13.67	12.50	"
14.85	13.31	"
18.74	15.90	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot \text{H}_2\text{O}$
23.50	19.21	"
23.50	19.29	"
27.15	20.96	"
28.11	21.64	"
29.50	22.53	"
30.19	22.90	"
31.48	23.70	"
32.93	24.83	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot \text{H}_2\text{O} +$
32.80	24.86	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O}$.
32.67	24.70	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O}$.
32.59	24.60	"
32.24	24.68	"
30.74	25.13	"
29.83	25.30	"
24.16	28.25	"
16.62	35.40	"
12.17	43.37	"
11.66	46.62	"
11.19	49.38	"
11.19	50.20	"
15.20	54.66	"
20.76	53.75	"
27.85	51.02	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O} +$
8.58	68.28	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 3\text{H}_2\text{O}$.
4.05	74.90	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 3\text{H}_2\text{O}$.

(Rutten, Z. anorg. 1902, 30. 386.)

Solubility of Bi_2O_3 in $\text{HNO}_3 + \text{Aq}$ at t° .

t°	% Bi_2O_3	% N_2O_5	Solid phase
9°	20.8	17.1	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot \text{H}_2\text{O}$
	24.02	19.1	"
	31.09	23.8	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot \text{H}_2\text{O} +$
	31.2	23.9	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O}$
30°	34.2	26.5	"
	28.2	29.6	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O}$
	16.1	47.7	"
65°	5.55	7.44	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot \text{H}_2\text{O}$
	27.62	22.46	"
	40.80	31.60	$\text{Bi}_2\text{O}_3 \cdot \text{N}_2\text{O}_5 \cdot \text{H}_2\text{O} + \text{Bi}_2\text{O}_3 \cdot$
			$3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O}$
	37.82	35.80	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O}$
	35.73	47.02	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 10\text{H}_2\text{O} +$
	4.59	77.90	$\text{Bi}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 \cdot 3\text{H}_2\text{O}$

(Rutten.)

Solubility of Bi ₂ O ₃ in HNO ₃ +Aq at t°.			
t°	% Bi ₂ O ₃	% N ₂ O ₅	Solid phase
72°	37.23	47.76	Bi ₂ O ₃ .3N ₂ O ₅ .4H ₂ O
75°	36.74	47.91	"
80°	39.75	45.16	"
9°	31.2	23.9	Bi ₂ O ₃ .3N ₂ O ₅ .10H ₂ O +
20°	32.8	24.8	Bi ₂ O ₃ .N ₂ O ₅ .H ₂ O
30°	34.2	26.4	"
50°	36.9	28.9	"
64°	40.6	31.1	"
65°	40.8	31.6	"
75.5°	45.4	34.6	"
72°	45.9	35.6	"
11.5°	25.36	52.57	} Bi ₂ O ₃ .3N ₂ O ₅ .10H ₂ O + Bi ₂ O ₃ .3N ₂ O ₅ .3H ₂ O
20°	27.85	51.02	
50°	32.22	49.29	
65°	35.73	47.02	

(Rutten.)

Solubility in NaOH+Aq at 25°.	
Conc. of NaOH Mol./l.	g. Bi ₂ O ₃ in 100 cc. of solution. Mean result.
1.0	0.0013 ± 0.0002
2.0	0.0026 ± 0.0002
3.0	0.0049 ± 0.0005

(Knox, Chem. Soc. 1909, 95. 1767.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

Min. *Bismite*. Easily sol. in HNO₃+Aq.
See also Bismuthous hydroxide.

Bismuth tetroxide, Bi₂O₄.

Sol. in conc. HCl+Aq, with evolution of Cl; in oxygen acids with evolution of O. Less easily sol. in conc. H₂SO₄ than in HNO₃, or HCl+Aq.

Bismuth oxide, Bi₄O₇ (?).

(Hoffmann and Geuther.)

Bismuth pentoxide, Bi₂O₅.

Sol. in dil. acids. Combines with H₂O to form bismuthic hydroxide, which see. (Hasebroek, B. 20. 213.)

Bismuth oxybromide, etc.

See Bismuthyl bromide, etc.

Bismuth palladium, PdBi₂.

Insol. in equal pts. HNO₃ and tartaric acids. (Roessler, Z. anorg. 1895, 9. 70.)

Bismuth platinum, PtBi₂.

Insol. equal pts. HNO₃ and tartaric acids. (Roessler, Z. anorg. 1895, 9. 69.)

Bismuth phosphide, BiP.
(Cavazzi.)

Bismuth triselenide, Bi₂Se₃.

Insol. in H₂O, alkalies, or alkali sulphides +Aq; sl. attacked by HCl+Aq; oxidized by HNO₃+Aq. (Schneider, Pogg. 94. 628.)
Min. *Frenzelite*.

Bismuth potassium selenide.

See Selenobismuthite, potassium.

Bismuth selenochloride, BiSeCl.

Not attacked by H₂O; very sl. sol. in HCl+Aq; easily and completely sol. with decomp. in HNO₃+Aq. (Schneider.)

Bismuth disulphide, Bi₂S₃+2H₂O (?).

Insol. in H₂O. Decomp. by HCl+Aq.

Bismuth trisulphide, Bi₂S₃.

Insol. in H₂O.

1 l. H₂O dissolves 0.35 x 10⁻⁴ moles Bi₂S₃ at 18°. (Weigel, Z. phys. Ch. 1907, 58. 294.)

Easily sol. in moderately dil. HNO₃+Aq, and conc. HCl+Aq, with separation of S. Insol. in alkalies, alkali sulphides, Na₂S₂O₃, or KCN+Aq; insol. in NH₄Cl, or NH₄NO₃+Aq (Brett). Insol. in potassium thiocarbonate+Aq. (Rosenbladt, Z. anal. 26. 15.)

Insol. in alkali hydroxides or alkali hydro-sulphides.

Insol. in 2N-(NH₄)₂S+Aq.

0.0090 g. Bi₂S₃ is sol. in 100 cc. N-Na₂S₃+Aq at 25°. (Knox, Chem. Soc. 1909, 95. 1764.)

Somewhat sol. in Na₂S+Aq. 75 cc. of Na₂S+Aq (sp. gr. 1.06) dissolve an amt. of Bi₂S₃ corresponding to 0.031 g. Bi₂O₃. (Stillman, J. Am. Chem. Soc. 1896, 18. 683.)

Solubility in Na₂S+NaOH+Aq at 25°.

Conc. of Na ₂ S Mol./l.	Conc. of NaOH Mol./l.	g. Bi ₂ S ₃ in 100 cc. of solution
0.5	1.0	0.0185
1.0	1.0	0.0838

(Knox, Chem. Soc. 1909, 95. 1763.)

Bismuth sulphide pptd. from acid solution is not dissolved by subsequent treatment with K₂S+Aq. (Stone, J. Am. Chem. Soc. 1896, 18. 1091.)

Sol. in K₂S+Aq. (Ditte, C. R. 1895, 120. 187.)

Solubility in K₂S+KOH+Aq at 25°.

Conc. of K ₂ S Mol./l.	Conc. of KOH Mol./l.	g. Bi ₂ S ₃ in 100 cc. of solution
0.5	1.0	0.0240
1.0	1.0	0.1230
1.25	1.25	0.2354

(Knox, Chem. Soc. 1909, 95. 1763.)

ty in alkali sulphides + Aq at 25°.

Conc. of alkali sulphide Mol./l.	g. Bi_2S_3 in 100 cc. of solution
0.5	0.0040
1.0	0.0238
1.5	0.1023
0.5	0.0042
1.0	0.0337
1.25	0.0639

ix, Chem. Soc. 1909, **95**. 1762.)

o. by FeCl_3 + Aq. (Cammerer, C. C. 525.)

1 KCN + Aq. (Hoffmann, A. 1884,

bismuthinite. Easily sol. in HNO_3 +

suprous sulphide, Bi_2S_3 , Cu_2S .

1 H_2O . Sol. with decomp. in HNO_3 ,
Schneider, J. pr. (2) **40**. 564.)

amplectonite.

potassium sulphide, Bi_2S_3 , K_2S .

der, Pogg. **136**. 460.)

in ethyl acetate. (Naumann, B. 314.)

$\text{K}_2\text{S} + 4\text{H}_2\text{O}$. Decomp. by H_2O .
in K_2S + Aq. Efflorescent in dry
e, C. R. 1895, **120**. 186.)

Sulphobismuthite, potassium.

silver sulphide, Bi_2S_3 , Ag_2S .

1 cold HCl , or HNO_3 . Sol. in warm
h separation of S, in boiling HCl
ation of H_2S .

enargyrite, Matildite.

ler, J. pr. 1890, (2) **41**. 414.)

sodium sulphide, Bi_2S_3 , Na_2S .

ler.)

ulphide telluride, Bi_2S_3 , $2\text{Bi}_2\text{Te}_3$.

tetradymite. Sol. in HNO_3 with sepa-

Bi_2Te_3 .

seite. As above.

ulphobromide, BiSBr_2 .

nd Eagles, Chem. Soc. 1895, **67**. 91.)

ulphochloride, BiSCl_2 .

n H_2O or dil. HCl + Aq. Sol. in
l, or HNO_3 + Aq. Decomp. by
Aq. (Schneider, Pogg. **93**. 464.)

ulphoiodide, BiSI .

acked by boiling H_2O , and dil. acids.
by hot conc. HCl + Aq, and HNO_3 +
l + Aq dissolves out I_2 . (Schneider,
114.)

Bismuth telluride, Bi_2Te_3 .

Min. *Tetradymite*. Sol. in HNO_3 + Aq.

See also *Bismuth sulphide telluride*.

Bismuthic acid, HBiO_3 .

See *Bismuthic hydroxide*.

Potassium bismuthate, KBiO_3 .

Sol. in H_2O . (Arppe.)

$\text{KH}(\text{BiO}_3)_2$. Insol. in H_2O .

Not decomp. by boiling H_2O . (André, C. R. **113**. 860.)

No salts of HBiO_3 can exist. (Muir and Carnegie, Chem. Soc. **51**. 77.)

Bismuthicotungstic acid.

Ammonium bismuthicotungstate, $3(\text{NH}_4)_2\text{O}$,
 $2\text{Bi}_2\text{O}_3$, 11WO_3 + $10\text{H}_2\text{O}$.

A yellow oil which dries to a yellow glass.
(E. F. Smith, J. Am. Chem. Soc. 1903, **25**. 1232.)

Potassium bismuthicotungstate, $3\text{K}_2\text{O}$,
 $2\text{Bi}_2\text{O}_3$, 11WO_3 + $15\text{H}_2\text{O}$.

A yellow oil which dried to a pale yellow
glass. (E. F. Smith, J. Am. Chem. Soc. 1903,
25. 1233.)

Strontium bismuthicotungstate, 3SrO ,
 $2\text{Bi}_2\text{O}_3$, 11WO_3 + $11\text{H}_2\text{O}$.

A yellow wax, insol. in pure H_2O , but sol.
in H_2O containing a few drops HNO_3 . (E.
F. Smith, J. Am. Chem. Soc. 1903, **25**. 1233.)

Bismuthyl bromide, BiOBr .

Insol. in H_2O ; sol. in moderately conc.
 HBr + Aq.

Insol. in H_2O . (Herz, Z. anorg. 1903, **36**. 348.)

$\text{Bi}_5\text{O}_7\text{Br}_6$. Insol. in H_2O ; easily sol. in
conc. HCl , or HNO_3 + Aq; less sol. in dil.
 HNO_3 + Aq.

$\text{Bi}_{11}\text{O}_{13}\text{Br}_7$. As the preceding comp.
(Muir.)

Bismuthyl chloride, BiOCl .

Insol. in H_2O or dil. acids. Sol. in conc.
 HCl , or HNO_3 + Aq.

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 827.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329.)

+ H_2O . (Heintz, Pogg. **63**. 55.)

+ $3\text{H}_2\text{O}$. (Phillips, Br. Arch. (1) **39**. 41.)

$\text{Bi}_7\text{O}_9\text{Cl}_3$. (Arppe.)

BiO_2Cl_2 . Insol. in H_2O ; sol. in hot HCl , or
 HNO_3 + Aq. (Muir.)

Bismuthyl fluoride, BiOF .

Insol. in H_2O ; sol. in HCl , HBr , or HI + Aq.
(Gott and Muir, Chem. Soc. **33**. 139.)

BiOF , 2HF . Insol. in H_2O .

Bismuthyl iodide, BiOI.

Not decomp. by H_2O or alkaline solutions.
Sol. in $HCl + Aq.$ Decomp. by $HNO_3 + Aq.$
(Schneider, J. pr. 79. 424.)

Insol. in KCl , or $KI + Aq.$
 $3BiOI, 7Bi_2O_3$. Sol. in dil. HCl ; decomp.
by HNO_3 ; insol. in boiling H_2O and alkali.
(Blyth, C. N. 1896, 74. 200.)

$BiI_3, 5Bi_2O_3$. Ppt. Sl. sol. in $HC_2H_3O_2 + Aq.$
Not decomp. by H_2O . (Fletcher and
Cooper, Pharm. J. (3) 13. 254.)

$4BiI_3, 5Bi_2O_3$. Easily sol. in $HCl + Aq.$
Decomp. by $HNO_3 + Aq.$ Sl. attacked by
 H_2SO_4 ; somewhat sol. in $H_2C_4H_4O_6$, and
 $KHC_4H_4O_6 + Aq.$

Sol. in $(NH_4)_2S$, and $KOH + Aq.$ (Storer's
Dict.)

Bismuthyl sulphide, Bi_2O_3S .

(Hermann, J. pr. 75. 452.)

Bi_2O_3S . Insol. in H_2O . (Scherpenberg,
C. C. 1889, II. 641.)

Bi_4O_5S .

Min. Karelinite.

Boracic acid.

See Boric acid.

Borax.

See Tetraborate, sodium.

Boric acid, anhydrous, B_2O_3 .

See Boron trioxide.

Metaboric acid, HBO_2 .

Sol. in H_2O .

Sl. sol. in hot glacial acetic acid. (Holt,
Chem. Soc. 1911, 100. (2) 720.)

Orthoboric acid, H_3BO_3 .

Sol. in 33 pts. H_2O at 10° .

" 25 " " 20° .

" 3 " " 100° .

(Berzelius.)

Sol. in 20 pts. H_2O at 18.75° . (Abl.)

100 pts. H_2O at 100° dissolve 2 pts. (Ure's Dict.)

1 pt. crystallized acid dissolves in—

25.66 pts. H_2O at 19° .

14.88 " " 25° .

12.66 " " 37.5° .

10.16 " " 50° .

6.12 " " 62.5° .

4.73 " " 75° .

3.55 " " 87.5° .

2.97 " " 100° .

Or, 100 pts. H_2O dissolve at—

19° 3.9 pts. H_3BO_3 .

25° 6.8 " "

37.5° 7.8 " "

50° 9.8 " "

62.5° 16.0 " "

75° 21.0 " "

87.5° 28.0 " "

100° 34.0 " "

Or, sat. aqueous solution contains at—

19° 3.75% H_3BO_3 .

25° 6.27 " "

37.5° 7.32 " "

50° 8.96 " "

62.5° 14.04 " "

75° 17.44 " "

87.5° 21.95 " "

100° 25.17 " "

(Brandes and Firnhaber, Arch. Pharm. 7

1 litre H_2O dissolves at—

0° 19.47 g. H_3BO_3 .

12° 29.20 " "

20° 39.92 " "

40° 69.91 " "

62° 114.16 " "

80° 168.15 " "

102° 291.16 " "

(Ditte, C. R. 85. 1069.)

1 l. H_2O dissolves 0.901 mol. H_3BO_3 at
(Herz, Z. anorg. 1910, 66. 359.)

1 l. H_2O dissolves 0.898 mol. H_3BO_3 at
Sp. gr. of the solution = 1.0168. (Müller
phys. Ch. 1907, 57. 529.)

1 l. H_2O dissolves 0.887 mol. H_3BO_3 at
and 1.025 mol. at 30° . (Ageno and
Ist. Ven. (VIII) 14. II, 331.)

Solubility in H_2O at t° .

t°	g. H_3BO_3 in 100 g. of the soluti
0	2.59
12.2	3.69
21	4.90
31	6.44
40	8.02
50	10.35
60	12.90
69.5	15.58
80	19.11
90	23.30
99.5	28.10
108	36.7
115	45.0
120	52.4

(Nasini and Ageno, Z. phys. Ch. 1906
483.)

Solubility curve for orthoboric aci
 H_2O at various temp. up to 120° . (N
and Ageno, Gazz. ch. it. 1911, 41. (1) 1

Sp. gr. of $H_3BO_3 + Aq$ sat. at 8° = 1.014. (Anth
24. 241.)

Sp. gr. of $H_3BO_3 + Aq$ sat. at 15° = 1.0243. (Stol
pr. 90. 457.)

Sp. gr. of $H_3BO_3 + Aq$ at 15° .

% H_3BO_3	Sp. gr.	% H_3BO_3	Sp. g
1	1.0034	4	1.01
2	1.0069	Sat. sol.	1.01
3	1.0106

(Gerlach, Z. anal. 28. 473.)

gr. of $\text{H}_3\text{BO}_3 + \text{Aq}$ at 18° .

H_2O	0.776	1.92	2.88	3.612
.	1.0029	1.0073	1.0109	1.0131

(Bock, W. Ann. 1887, **30**. 638.)

atile with steam.

re sol. in dil. $\text{HCl} + \text{Aq}$ than in H_2O .
in warm conc. H_2SO_4 , HCl , or $\text{HNO}_3 +$

solubility in $\text{HCl} + \text{Aq}$ at 25° .

Millimols HCl in 10 ccm. of the solution	Millimols H_3BO_3 in 10 ccm. of the sat. solution
...	9.01
7.0	7.69
13.7	6.66

(Herz, Z. anorg. 1910, **66**. 359.)

solubility of H_3BO_3 in $\text{HCl} + \text{Aq}$ at 16° .

Normality of HCl	Normality of H_3BO_3
0.	0.907
0.130	0.895
0.260	0.870
0.390	0.842
1.30	0.645
2.16	0.542
4.32	0.308
6.00	0.338
7.08	0.327
8.74	0.327
9.51	0.338

(Herz, Z. anorg. 1902, **33**. 354.)

Solubility in $\text{HF} + \text{Aq}$ at 26° .

of	(2) Titer after saturation with H_3BO_3 at 26°	(3) Titer after addition of mannitol	(3)-(2) equals free boric acid
n.	1.61	2.36	0.75
n.	1.25(1.40?)	2.21	0.96(0.81?)

values 0.75 and 0.81 represent the
ty of H_3BO_3 in the concentrations of
ric acid resulting from the original
tration of $\text{HF} + \text{Aq}$.

Abegg, Z. anorg. 1903, **35**. 145.)

solubility of H_3BO_3 in acids + Aq at 26° .

	Normality of the acid	Normality of H_3BO_3
4	0.548	0.746
	2.74	0.518
	5.48	0.312
	8.75	0.092
3	0.241	0.818
	1.206	0.676
	1.607	0.593
	2.411	0.567
	5.96	0.268
	7.38	0.238

Herz, Z. anorg. 1903, **34**. 205.)

Solubility in $\text{KOH} + \text{Aq}$.

See Borates, potassium.

Solubility in $\text{NaOH} + \text{Aq}$.

See Borates, sodium.

Solubility in $\text{LiCl} + \text{Aq}$ at 25° .

Millimols LiCl in 10 ccm. of the solution	Millimols H_3BO_3 in 10 ccm. of the sat. solution
...	9.01
7.1	8.13
10.3	7.65
22.3	6.42
37.2	5.02

(Herz, Z. anorg. 1910, **66**. 359.)

Solubility in $\text{KCl} + \text{Aq}$ at 25° .

Millimols KCl in 10 ccm. of the solution	Millimols H_3BO_3 in 10 ccm. of the sat. solution
...	9.01
1.9	9.20
7.9	9.44
15.6	9.80
30.6	10.75

(Herz.)

Solubility in $\text{RbCl} + \text{Aq}$ at 25° .

Millimols RbCl in 10 ccm. of the solution	Millimols H_3BO_3 in 10 ccm. of the sat. solution
...	9.01
14.0	9.66
25.3	10.60

(Herz.)

Solubility in $\text{NaCl} + \text{Aq}$ at 25° .

Millimols NaCl in 10 ccm. of the solution	Millimols H_3BO_3 in 10 ccm. of the sat. solution
...	9.01
8.2	8.49
15.2	8.25
29.4	8.20

(Herz.)

Solubility in H_2O is increased by presence
of KCl , KNO_3 , K_2SO_4 , NaNO_3 , and Na_2SO_4 .

In general the solubility in H_2O is increased
by the presence of both electrolytes and non-
electrolytes. (Bogdan, C. C. 1903, II. 2.)

Sol. in borax + Aq . (McLauchlan, Z. anorg.
1903, **37**. 371.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch.
J. 1898, **20**. 827.)

Unattacked and undissolved by liquid NO_2 .
(Frankland, Chem. Soc. 1901, **79**. 1362.)

Sol. in 6 pts. alcohol (Wittstein), 5 pts.
boiling alcohol (Wenzel). Only traces dis-
solve in anhydrous ether. (Schiff.) Sol. in
100 pts. ether. (Hager's Comm.) Sol. in
several essential oils.

1 l. H_2O sat. with amyl alcohol dissolves
0.8952 mol. H_3BO_3 at 25° . (Auerbach, Z.
anorg. 1903, **37**. 357.)

Solubility of H_3BO_3 in amyl alcohol + Aq at t° .
M = millimols H_3BO_3 in 1 l. of H_2O .
A = millimols H_3BO_3 in 1 l. of alcohol.

t°	M	A
15°	607.2	176.4
	589.3	177.4
	589.0	177.1
	586.0	173.4
	427.4	127.6
	425.8	127.0
	289.1	84.9
	894.0	264.0
	372.0	110.0
	371.8	110.8
25°	301.2	85.7
	180.8	54.0
	49.15	15.45
	51.04	15.45
	26.02	8.05
35°	146.3	44.27

(Müller, Z. phys. Ch. 1907, 57. 514.)

Sp. gr. of amyl alcohol + Aq sat. with H_3BO_3 .

g. water in 1 l. of alcohol + Aq	$d_{25^\circ/4^\circ}$
32.481	0.82229
35.465	0.82324
37.339	0.82321
42.479	0.82392
45.175	0.82447
45.636	0.82456
47.883	0.82454
51.461	0.82527
52.043	0.82585
59.270	0.82699
63.179	0.82739
64.254	0.82779
66.403	0.82701
66.624	0.82670
68.253	0.82856
69.211	0.82884
75.610	0.82999(?)

(Müller.)

Solubility of H_3BO_3 in amyl alcohol and NaCl + Aq at 25°.

Water phase		Amyl alcohol phase			
NaCl normality	mol. H_3BO_3	Sp. gr. 25°/4°	1 l. contains		
			mol. H_2O	mol. amyl alcohol	mol. H_3BO_3
0.00	0.880	0.8296	4.10	8.39	0.2640
0.945	0.866	0.8277	3.55	8.49	0.2638
1.490	0.850	0.8268	3.27	8.54	0.2689
1.865	0.844	0.8259	3.03	8.56	0.2724
2.355	0.833	0.8254	2.86	8.59	0.2850
2.845	0.827	0.8247	2.62	8.62	0.2877
3.06	0.810	0.8241	2.39	8.66	0.2891
3.48	0.810	0.8240	2.32	8.69	0.3006
3.57	0.807	0.8236	2.15	8.70	0.3066
4.01	0.801	0.8233	1.99	8.72	0.3162
4.28	0.798	0.8229	1.78	8.75	0.3210

(Müller)

Solubility in hydroxy-compounds + Aq at 25°.

Organic substance added	Mol. of organic substance in 100 mol. of the mixture	Mol. of boric acid sol. in 1 l. of solution	Sp. gr. of the pure mixture	Sp. gr. of the mixture sat. with boric acid
Lactic acid	2.321	1.07	1.0252	1.0444
	6.819	1.61	1.0722	1.0986
	18.77	1.86	1.1405	1.1635
	36.33	2.08	1.2023	1.2254
Glycerine	24.64	1.208	1.1574	1.1707
	46.75	2.132	...	1.2260
	67.71	2.96	1.2370	1.2526
	90.58	3.78	1.2531	1.2710

Solubility in hydroxy-compounds etc.—Continued

Substance added	Mol. of organic substance in 100 mol. of the mixture	Mol. of boric acid sol. in 1 l. of solution	Sp. gr. of the pure mixture	Sp. gr. of the mixture sat. with boric acid
mannitol	0.790	1.007		1.0425
	0.810	1.015	1.0244	1.0433
	0.945	1.029	1.0288	
	1.585	1.136	1.0475	
sorbitol	0.065	0.8876	0.9995	1.0686
	0.130	0.9078	1.0018	1.0212
	0.260	0.9360	1.0060	1.0260

(Müller.)

Solubility of H_2BO_3 in alcohols + Aq at 25°.
M = Mol. of alcohol in 100 mol. of alcohol + Aq.
 H_2BO_3 = Mol. of H_2BO_3 in 1 l. of the solution.
 d_1 = Sp. gr. of alcohol + Aq.
 d_2 = Sp. gr. of alcohol + Aq sat. with H_2BO_3 .

Alcohol added	M	H_2BO_3	d_1	d_2
Ethanol	11.74	0.895		
	28.64	1.012		
	36.02	1.098		
	43.95	1.161		
	52.31	1.307		
	100	2.900	0.7924	0.8904
Methanol	8.996	0.829		
	22.28	0.800		
	44.46	0.729		
	55.62	0.700		
	79.89	0.893		
	88.10	1.105		
	99.26	1.527	0.7860	0.8353
Vinyl alcohol	23.66	0.6437	0.9043	0.9193
	53.63	0.4569	0.8231	0.8570
	83.65	0.5776	0.8133	0.8466
	100	0.961	0.8010	0.8297
Propyl alcohol	0.70	0.884	0.9923	1.0124
	2.15	0.857	0.9853	0.0038
	2.18	0.857	0.9855	0.0046
	71.4	0.323	0.8173	0.8351
	77.1	0.347	0.8133	0.8220
	85.6	0.4212	0.8081	0.8195
	100	0.6927	0.7984	0.8172
Butyl alcohol	0.448	0.883	0.9943	1.0132
	0.520	0.880	0.9936	1.0125
	0.525 ¹	0.880	0.9931	1.0123
	67.26 ²	0.2584	0.8232	0.829
	75.54	0.2722	0.8183	0.8253
	83.40	0.3190	0.8142	0.8223
	100	0.5703	0.8068	0.8220

¹ Water sat. with alcohol.² Alcohol sat. with water.

(Müller.)

Easily sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

Solubility in acetone+Aq at 20°. A = ccm. acetone in 100 ccm. acetone+Aq. H₂BO₃ = millimols H₂BO₃ in 100 ccm. of the solution.

A	H ₂ BO ₃
0	79.15
20	81.71
30	83.35
40	82.74
50	81.61
60	76.40
70	67.62
80	55.05
100	8.06

(Herz, Z. anorg. 1904, 41. 319.)

100 g. pure anhydrous ether dissolve 0.00775 g. H₂BO₃.

100 g. ether sat. with H₂O dissolve 0.2391 g. H₂BO₃.

(J. A. Rose, Dissert. 1902.)

Sol. in 10 pts. glycerine. (Häger.)

100 pts. glycerine (sp. gr. 1.26 at 15.5°) dissolve pts. H₂BO₃ at t°.

t°	Pts. H ₂ BO ₃	t°	Pts. H ₂ BO ₃	t°	Pts. H ₂ BO ₃
0	20	40	38	80	61
10	24	50	44	90	67
20	28	60	50	100	72
30	33	70	56

(Hooper, Ph. J. Trans. (3) 13. 258.)

Solubility of H₂BO₃ in glycerine+Aq at 25°.

G = g. glycerine in 100 g. glycerine+Aq. H₂BO₃ = Millimols H₂BO₃ in 100 cc. of the solution.

G	H ₂ BO ₃	Sp. gr.
0	90.1	1.0170
7.15	90.1	1.0379
20.44	90.6	1.0629
31.55	92.9	1.0897
40.95	97.0	1.1130
48.7	103.0	1.1328
69.2	140.2	1.1871
100	390.3	1.2719

(Herz, Z. anorg. 1905, 45. 268.)

Solubility of H₂BO₃ in organic acids+Aq at 26°.

Acid	Normality of the acid	Normality of H ₂ BO ₃
Acetic	0.570	0.887
	2.85	0.538
	5.70	0.268

Solubility of H₂BO₃, etc.—Continued.

Acid	Normality of the acid	Normality of H ₂ BO ₃
Tartaric	0.955	0.890
	1.909	0.923
	2.51	0.962
	3.316	1.07

(Herz, Z. anorg. 1903, 34. 206.)

The solubility of H₂BO₃ in H₂O is increased by the presence of racemic acid.

Millimols racemic acid in 10 ccm. of the solvent	Millimols boric acid in 10 ccm. of the solution
0	9.01
6.3	9.86
12.6	10.46
24.7	11.65

(Herz, Z. anorg. 1911, 70. 71.)

Solubility of H₂BO₃ in H₂O is increased by the presence of tartaric acid.

Millimols tartaric acid in 10 ccm. of the solvent	Millimols boric acid in 10 ccm. of the solution
0	9.01
7.5	10.00
15	10.70
30	12.07

(Herz, Z. anorg. 1911, 70. 71.)

Solubility in oxalic acid+Aq at 25°.

Millimols oxalic acid in 10 ccm. of the solution	Millimols H ₂ BO ₃ in 10 ccm. of the sat. solution
...	9.01
2.97	9.95
5.95	10.80
13.77	11.98

(Herz, Z. anorg. 1910, 66. 93.)

Solubility in H₂O is increased by the presence of urea, acetone or propyl alcohol. (Bogdan, C. C. 1903, II. 2.)

Readily sol. in hot glacial acetic acid. (Holt Chem. Soc. 1911, 100 (2). 720.)

Sol. in 250 pts. benzene. (Häger.)

Solubility of H₂BO₃ in mannite+Aq at t°.

Solid phase, H ₂ BO ₃					
t°	Mg.-mole. in 1 l.		t°	Mg.-mols in 1 l.	
	Mannite	H ₂ BO ₃		Mannite	H ₂ BO ₃
25°	0	0.887	30°	0	1.025
"	0.1	0.951	"	0.1	1.056
"	0.3	1.015	"	0.2	1.086
"	0.4	1.039	"	0.3	1.118
"	0.5	1.071	"	0.4	1.157
"	0.6	1.102	"	0.5	0.193
"	0.7	1.142	"	0.6	1.219
"	0.8	1.173	"	0.7	1.258
"	1.043	1.244			
"	1.409	1.404			
"	1.781	1.521			

Solid phase, mannite		
t°	Mg. mols. in 1 l.	
	Mannite	H ₃ BO ₃
25°	1.075	0
"	1.1424	0.2646
"	1.259	0.463
"	1.265	0.559
"	1.354	0.794
"	1.409	0.927
"	1.536	1.243
"	1.781	1.521

(Ageno and Valla, Ist. Ven. (VIII) 14. 331.)

Distribution between H₂O and amyl alcohol at 25°.

w = concentration of H₃BO₃ in H₂O layer expressed in millimols.

a = concentration of H₃BO₃ in alcohol layer expressed in millimols.

w	a
265.8	76.6
196.5	59.5
159.6	47.5
126	37.1
87.9	33.2
75.2	22.7
64.6	19.76

(Abegg, Z. anorg. 1903, 35. 130.)

Partition of H₃BO₃ between water and mixtures of amyl alcohol and CS₂.

W = Millimols H₃BO₃ in 10 ccm. of the aqueous layer.

G = Millimols H₃BO₃ in 10 ccm. of the amyl alcohol—CS₂ layer.

Composition of the solvent mixture	G	W	W, G
75% by vol. amyl alcohol + 25% by vol. CS ₂	0.145	0.624	4.31
	0.275	1.198	4.36
	0.429	1.844	4.30
	0.589	2.565	4.45
50% by vol. amyl alcohol + 50% by vol. CS ₂	0.145	0.756	5.47
	0.259	1.353	5.21
	0.364	1.946	5.34
	0.555	2.889	5.22
25% by vol. amyl alcohol + 75% by vol. CS ₂	0.085	0.699	8.24
	0.175	1.467	8.40
	0.264	2.165	8.12
	0.384	3.129	8.14

(Herz, Z. Elektrochem. 1910, 16. 870.)

Distribution between HF + Aq and amyl alcohol at 25°.

c = HF concentration (millimols).

a = H₃BO₃ concentration in alcohol layer (expressed in millimols).

w = H₃BO₃ concentration in water layer (expressed in millimols).

c	a	w
500	14.3	71.2
"	19.2	99.2
"	25.3	144.2
"	114.3	979.0
250	30.1	144.5
"	37.0	194.8
"	56.8	321.5
"	108.0	652.0
125	39.0	170.5
"	47.2	214.0
"	52.8	240.5
"	96.0	442.0
62.5	30.4	111.2
"	39.4	151.8
"	65 (68?)	272.8
"	90.0	362.2

(Abegg, Z. anorg. 1903, 35. 131.)

See also Boron trioxide.

Pyroboric (tetraboric) acid, H₂B₄O₇.

Sol. in H₂O.

Sp. gr. of solutions of boric acid, calculated as H₂B₄O₇, containing—

6.3 1.27 1.91 2.54% H₂B₄O₇
1.0034 1.0069 1.0106 1.0147 sp. gr.

Sat. solution at 15° has sp. gr. 1.015. (Gerlach, Z. anal. 28. 473.)

Insol. in hot glacial acetic acid. (Holt, Chem. Soc. 1911, 100. (2) 720.)

Borates.

No borate is quite insol. in H₂O; the alkali borates are very sol. The less sol. borates are easily decomp. by H₂O; the easily sol. salts are also decomp., but less quickly. The less sol. borates are easily sol. in H₃BO₃, HNO₃, etc. They are more sol. in H₂O containing tartaric acid or potassium tartrate than in pure H₂O. (Souberain.) The normal borates of the alkaline-earths are sol. to no inconsiderable extent in H₂O, and more readily in hot, than in cold H₂O. (Berzelius, Pogg. 34. 568.)

All borates are insol., or sl. sol. in alcohol.

Aluminum borate, 2Al₂O₃, B₂O₃.

Min. *Jeremciwite*.

+3H₂O. Ppt. (Rose, Pogg. 91. 452.)

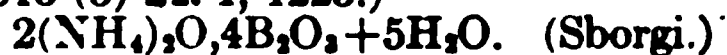
3Al₂O₃, B₂O₃. *Crystallized*. Insol. in HNO₃ + Aq. (Ebelmen, A. ch. (3) 33. 62.)

3Al₂O₃, 2B₂O₃ + 7H₂O. Ppt. (Rose, l. c.)

Ammonium borate.

The system (NH₄)₂O, B₂O₃, H₂O at 60° has

been studied by Sborgi. (Real. Ac. Linc. 1915 (5) 24. I, 1225.)



Ammonium diborate.

Difficultly sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Ammonium tetraborate, $(\text{NH}_4)_2\text{B}_4\text{O}_7 + 4\text{H}_2\text{O}$,
or perhaps $\text{NH}_4\text{H}(\text{BO}_2)_2 + 1\frac{1}{2}\text{H}_2\text{O}$.

Sol. in 12 pts. cold H_2O ; decomp. by heat. (Rammelsberg, Pogg. 90. 21.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

+ H_2O . (Arfvedson.)

Ammonium octoborate, $(\text{NH}_4)_2\text{B}_8\text{O}_{13} + 6\text{H}_2\text{O}$.

Sol. in 8 pts. cold, decomp. by boiling H_2O . (Rammelsberg, Pogg. 90. 21.)

+ $4\text{H}_2\text{O}$.

Min. *Lordellerite*. Sol. in H_2O with decomp.

Ammonium dekaborate, $(\text{NH}_4)_2\text{B}_{10}\text{O}_{16} + 6\text{H}_2\text{O}$.

Permanent. Sol. in H_2O . (Rammelsberg.)
+ $8\text{H}_2\text{O}$. (Atterberg, Bull. Soc. (2) 22. 350.)

Ammonium dodekaborate, $(\text{NH}_4)_2\text{B}_{12}\text{O}_{18} + 9\text{H}_2\text{O}$.

Sol. in hot H_2O . (Bechi, Sill. Am. J. (2) 17. 129.)

Ammonium perborate, NH_4BO_3 .

See Perborate, ammonium.

Ammonium calcium borate, $(\text{NH}_4)_2\text{CaB}_4\text{O}_{11}$
= $\text{CaB}_4\text{O}_7 + 4(\text{NH}_4)_2\text{O}$.

(Ditte, C. R. 96. 1663.)

Ammonium magnesium borate.

Sol. in H_2O , decomp. by boiling. (Rammelsberg, Pogg. 49. 451.)

Ammonium zinc borate, $4(\text{NH}_4)_2\text{B}_4\text{O}_7$,
 $\text{Zn}(\text{BO}_2)_2 + 5\text{H}_2\text{O}$.

(Ditte, C. R. 96. 1663.)

Barium borate, $\text{Ba}(\text{BO}_2)_2$.

Ppt. (Ouvrard, C. R. 1906, 142. 283.)

+ $2\text{H}_2\text{O}$. (Atterberg.)

+ $4\text{H}_2\text{O}$. (Benedikt, B. 7. 703.)

Sol. in 3,300 pts. 45% alcohol.

" 7,800 " 50 "

" 25,000 " 60 "

" 55,000 " 75 "

(Berg, Z. anal. 16. 25.)

+ $10\text{H}_2\text{O}$. Sl. sol. in cold, more readily in hot H_2O , especially in presence of ammonium salts. (Berzelius, Pogg. 34. 568.) Sol. in sodium citrate + Aq. (Spiller.) Insol. in wood spirit. (Ebelmen.)

$2\text{BaO} \cdot \text{B}_2\text{O}_3$. Decomp. by H_2O forming BaO , B_2O_3 + $4\text{H}_2\text{O}$. (Ouvrard, C. R. 1906, 142. 283.)

$3\text{BaO} \cdot \text{B}_2\text{O}_3$. Easily sol. in mineral acids. Sl. attacked by dil. acetic acid. (Ouvrard, C. R. 1901, 132. 258.)

$\text{BaB}_4\text{O}_{10}$. Slowly sol. in warm dilute HNO_3 + Aq. (Ditte, C. R. 77. 892.)

+ $5\text{H}_2\text{O}$. Sol. in 100 pts. cold, and more freely in hot H_2O . When freshly pptd. sol. in cold NH_4Cl + Aq (Wackenroder, A. 41. 315); NH_4NO_3 + Aq (Brett, Phil. Mag. (3) 10. 96); and BaCl_2 + Aq (Rose).

$\text{BaB}_6\text{O}_{10} + 13\text{H}_2\text{O}$. (Laurent, A. ch. (2) 67. 215.)

$\text{Ba}_2\text{B}_2\text{O}_5$. (Bloxam, Chem. Soc. 14. 143.)
 $5\text{BaO} \cdot 2\text{B}_2\text{O}_3$.

$\text{Ba}_3\text{B}_{10}\text{O}_{18} + 6\text{H}_2\text{O}$. Sol. in 100 pts. cold H_2O . Easily sol. in ammonium nitrate, or chloride, or barium chloride + Aq. (Rose, Pogg. 87. 1.)

$\text{Ba}_2\text{B}_6\text{O}_{11}$. Easily sol. in warm dilute acids.
+ $6\text{H}_2\text{O}$.

+ $7\text{H}_2\text{O}$.

+ $15\text{H}_2\text{O}$. (Laurent, A. ch. (2) 67. 215.)

Barium borate bromide, $3\text{BaO} \cdot 5\text{B}_2\text{O}_3 \cdot \text{BaBr}_2$.
(Ouvrard, C. R. 1906, 142. 283.)

Barium borate chloride, $3\text{BaO} \cdot 5\text{B}_2\text{O}_3 \cdot \text{BaCl}_2$.

Unaffected by H_2O . Sol. in acids. (Ouvrard, C. R. 1906, 142. 283.)

Bismuth borate, $\text{BiBO}_3 + 2\text{H}_2\text{O}$.

Ppt. Sl. sol. in H_2O . Decomp. by H_2S . Not decomp. by KOH + Aq. (Vanino, J. pr. 1906, (2) 74. 152.)

Cadmium borate, $\text{Cd}_2(\text{BO}_2)_2$.

Insol. in H_2O , easily sol. in dil. acids. (Ouvrard, C. R. 1900, 130. 174.)

$\text{Cd}(\text{BO}_2)_2$. Difficultly sol. in H_2O (Stromeyer); insol. in H_2O , sol. in HCl + Aq (Odling); easily sol. in warm NH_4Cl + Aq (Rose).

(Guertler, Z. anorg. 1904, 40. 242.)

$3\text{CdO} \cdot 2\text{B}_2\text{O}_3 + 3\text{H}_2\text{O}$. Ppt. Sl. sol. in H_2O . (Rose, Pogg. 88. 299.)

$\text{CdO} \cdot 2\text{B}_2\text{O}_3 + 2\text{H}_2\text{O}$. (Ditte, A. ch. 1883, (5) 30. 255.)

$\text{CdO} \cdot 4\text{B}_2\text{O}_3 + 10\text{H}_2\text{O}$. Sol. in H_2O ; decomp. on heating. (Ditte, A. ch. 1883, (5) 30. 255.)

Cadmium borate bromide, $6\text{CdO} \cdot 8\text{B}_2\text{O}_3 \cdot \text{CdBr}_2$.

Insol. in H_2O and fuming HCl or HBr + Aq. (Rousseau and Allaire, C. R. 1894, 119. 72.)

Cadmium borate chloride, $6\text{CdO} \cdot 8\text{B}_2\text{O}_3 \cdot \text{CdCl}_2$.

(Rousseau and Allaire, C. R. 1894, 118. 1256.)

Cadmium borate iodide, $6\text{CdO} \cdot 8\text{B}_2\text{O}_3 \cdot \text{CdI}_2$.

(Allaire, C. R. 1898, 127. 557.)

Cæsium borate, $\text{Cs}_2\text{B}_4\text{O}_{10}$.

Very sol. in H_2O , less in alcohol. (Reischle, Z. anorg. 4. 116.)

borate, $\text{Ca}(\text{BO}_2)_2$.

l. in H_2O ; insol. in alkali chlorides, or conc. acetic acid + Aq; sol. in cold or solutions of ammonium salts, especially ammonium nitrate, in CaCl_2 + Aq, and also sol. in dilute mineral acids at 50° . C. R. 80. 490, 561.)

O_7 .

O_7 ; two modifications of which one is unstable. (van't Hoff and Meyer-A. 1906, 351. 101.)

O_7 . When warmed in H_2O it goes to $\text{CaB}_2\text{O}_4 + 4\text{H}_2\text{O}$. (van't Hoff and offer.)

in H_2O without decomp.; 1 l. solution of 2 g. salt. (Ditte, C. R. 96. 1663.)

O_7 . Decomp. by H_2O . (Blount, C. N. 99. 276.)

sol. in methyl acetate. (Naumann, B. 1. 3790.)

O_7 . (Ditte, C. R. 96. 1663.)

O_7 . Min. *Bechilite*.

O_7 . Min. *Borocalcite*. Sol. in acids. $\text{O}_7 + 4\text{H}_2\text{O}$.

O_7 . Unstable. On standing in the solution in which it is formed it changes into $\text{O}_7 + 4\text{H}_2\text{O}$.

O_7 . Unstable. Goes over into $\text{O}_7 + 8\text{H}_2\text{O}$. (van't Hoff and Meyer-A. 1906, 351. 104.)

$\text{O}_{11} + 12\text{H}_2\text{O}$. (Ditte, C. R. 96. 1663.)

O_{11} . Insol. in H_2O , sol. in dil. acids. (Ditte, C. R. 1905, 141. 353.)

O_{11} . (Ditte, C. R. 77. 785.)

O_{11} . Min. *Pandermite*, *Priceite*. See $\text{B}_2\text{O}_3 + 9\text{H}_2\text{O}$.

O_{11} . Min. *Colemanite*.

the Ca is in form of colemanite, the sample contains in 100 g., 4.8 g. H_2BO_3 and 0.1 g. CaO . (van't Hoff, B. A. B. 1907, 653.)

O_{11} .

O_{11} . (van't Hoff and Meyerhoffer, A. 1. 101.)

O_{11} . Easily sol. in dil. acids. (Ditte, C. R. 1901, 132. 258.)

O_{11} . $5\text{B}_2\text{O}_3 + 9\text{H}_2\text{O}$. (van't Hoff, B. A. B. 1907, 568.)

O_{11} . $5\text{B}_2\text{O}_3 + 9\text{H}_2\text{O}$. True composition of *Priceite*. (van't Hoff, B. A. B. 1906, II.)

iron (ferrous) borate silicate,

$\text{FeB}_2\text{Si}_2\text{O}_{10}$.

Homilite. Easily sol. in HCl + Aq.

magnesium borate, CaO , MgO , $\text{O}_3 + 6\text{H}_2\text{O}$.

Hydroboracite. Somewhat sol. in H_2O . Sol. in warm HCl + Aq or HNO_3 + Aq. (Ditte, C. R. 77. 1257.)

sodium borate, 2CaO , Na_2O , $5\text{B}_2\text{O}_3$, H_2O .

(van't Hoff, B. A. B. 1907, 303.)

O_{11} , $\text{Na}_2\text{B}_4\text{O}_{10} + 15$, or $24\text{H}_2\text{O}$.

Min. *Natroborocalcite*, *Ulexite*. Decomp. by boiling with H_2O . Sol. in acids.

$\text{Ca}_2\text{Na}_4\text{B}_{10}\text{O}_{22} + 15\text{H}_2\text{O}$. Min. *Franklandite*. Sl. sol. in H_2O ; easily sol. in HCl , and HNO_3 + Aq.

Calcium borate bromide, 3CaO , $3\text{B}_2\text{O}_3$, CaBr_2 .

Sl. attacked by H_2O . Very sol. in dilute acetic acid. (Ouvrard, C. R. 1905, 141. 1023.)

3CaO , $5\text{B}_2\text{O}_3$, CaBr_2 . Hardly attacked by cold H_2O or very dil. acetic acid. Sol. in strong acids, even when dilute. (Ouvrard, C. R. 1905, 141. 1023.)

Calcium borate chloride, $\text{Ca}_2\text{B}_2\text{O}_6$, CaCl_2 .

Decomp. quickly by moist air or H_2O , slowly by absolute alcohol. (Chatelier, C. R. 99. 276.)

3CaO , $3\text{B}_2\text{O}_3$, CaCl_2 . (Ouvrard, C. R. 1905, 141. 353.)

3CaO , $5\text{B}_2\text{O}_3$, CaCl_2 . Sl. attacked by cold H_2O and dil. acetic acid + Aq. Strong acids dissolve even when very dilute. (Ouvrard, C. R. 1905, 141. 352.)

Calcium borate silicate, 2CaO , B_2O_3 , 2SiO_2 , $+ \text{H}_2\text{O}$.

Min. *Datolite*. Sol. in HCl + Aq with separation of gelatinous silica.

$+ 2\text{H}_2\text{O}$. Min. *Botryolite*.

CaO , B_2O_3 , SiO_2 . Min. *Danburite*. Very sl. attacked by HCl + Aq before ignition.

Chromous borate.

Precipitate. Sol. in free acids; insol. in borax + Aq. (Moberg.)

Chromic borate, $7\text{Cr}_2\text{O}_3$, $4\text{B}_2\text{O}_3$.

Insol. in H_2O ; sol. in excess of borax + Aq. (Hebberling, C. C. 1870. 122.)

Chromic magnesium borate, $3\text{Cr}_2\text{O}_3$, 6MgO , $2\text{B}_2\text{O}_3$.

Not attacked by acids. (Ebelmen, A. ch. (3) 33. 52.)

$2\text{Cr}_2\text{O}_3$, 9MgO , $3\text{B}_2\text{O}_3$. (Mallard, C. R. 105. 1260.)

Cobaltous borate, 3CoO , $2\text{B}_2\text{O}_3 + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Rose, Pogg. 88. 299.)

3CoO , B_2O_3 . (Mallard, C. R. 105. 1260.)

2CoO , B_2O_3 . (Ouvrard, C. R. 1900, 130. 337.)

Cobaltous borate bromide, 6CoO , $8\text{B}_2\text{O}_3$, CoBr_2 .

(Rousseau and Allaire, C. R. 1894, 119. 73.)

Cobaltous borate chloride, 6CoO , $8\text{B}_2\text{O}_3$, CoCl_2 .

(Rousseau and Allaire, C. R. 1894, 118. 1257.)

Cobaltous borate iodide, 6CoO , $8\text{B}_2\text{O}_3$, CoI_2 .

(Allaire, C. R. 1898, 127. 557.)

Cuprous borate, $3\text{Cu}_2\text{O}$, $2\text{B}_2\text{O}_3$.

(Guertler, Z. anorg. 1904, 38. 459.)

Cupric borate.

Composition depends on temperature and concentration of solutions. Boiling H_2O dissolves out all the boric acid. Sol. in acids; slowly sol. in hot conc. $NH_4Cl + Aq.$

$Cu(BO_2)_2$. Insol. in cold dil. acids, even HF . Slowly sol. in hot conc. HCl . Not attacked by alkalis or alkali carbonates + $Aq.$ (Guertler, Z. anorg. 1904, **38**. 456.)

Insol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

Cupric borate ammonia, $CuB_4O_7, 4NH_3 + 6H_2O$.

Efflorescent. Can be recrystallized from a little $NH_4OH + Aq.$ (Pasternack, A. **151**. 227.)

Didymium borate, $DiBO_3$.

Insol. in H_2O acidulated with $HCl + Aq.$ (Cleve, Bull. Soc. (2) **43**. 363.)

$Di_2(B_4O_7)_3$. Insol. in H_2O ; sol. in acids. (Frerichs and Smith, A. **191**. 355.)

Glucinum borate, basic, $5GfO, B_2O_3$.

Insol. in H_2O ; sol. in acids. (Krüss and Moraht, B. **23**. 735.)

Iron (ferrous) borate.

Ppt. H_2O dissolves out all the boric acid. (Tünnerman.)

Iron (ferric) borate, $Fe_2(BO_2)_6 + 3H_2O$.

Ppt. Insol. in H_2O .

Min. *Lagonite*. Sol. in acids.

$2Fe_2O_3, 3B_2O_3$. (Mallard, C. R. **105**. 1260.)

$6Fe_2O_3, B_2O_3 + 6H_2O$. Ppt. (Rose, Pogg. **89**. 473.)

$9Fe_2O_3, B_2O_3 + 9H_2O$. Ppt. (Rose.)

Iron (ferric) magnesium borate, $3Fe_2O_3, 6MgO, 2B_2O_3$.

Insol. in H_2O . Sol. in conc. $HCl + Aq.$ (Ebelmen, A. ch. (3) **33**. 53.)

$2Fe_2O_3, 9MgO, 3B_2O_3$. (Mallard, C. R. **105**. 1260.)

Iron (ferroferric) magnesium borate, $3MgO, FeO, Fe_2O_3, B_2O_3$.

Min. *Ludwigite*. Slowly sol. in $HCl + Aq.$ when finely powdered.

Iron (ferrous) borate bromide, $6FeO, 8B_2O_3, FeBr_2$.

Slowly sol. in hot $HNO_3 + Aq.$ (Rousseau and Allaire, C. R. **116**. 1445.)

Iron (ferrous) borate chloride, $6FeO, 8B_2O_3, FeCl_2$.

Slowly sol. in hot $HNO_3 + Aq.$ (Rousseau and Allaire, C. R. **116**. 1195.)

Lanthanum borate, $2La_2O_3, B_2O_3$.

(Nordenskjöld, Pogg. **114**. 618.)

$La_2(B_4O_7)_3$. Ppt. (Smith.)

Formula is $La_2B_6O_{13} + xH_2O$. (Cleve, B. **11**. 910.)

Lead borate, basic,

$2PbO, B_2O_3 + 2H_2O$. Ppt.

$4PbO, 3B_2O_3 + 4H_2O$. Ppt.

+ $5H_2O$. Ppt.

$6PbO, 5B_2O_3 + 6H_2O$. Ppt.

$8PbO, 3B_2O_3 + 8H_2O$. Ppt.

$9PbO, 5B_2O_3 + 9H_2O$. Ppt. (Rose, Pogg. **87**. 470.)

Lead borate, $Pb(BO_2)_2 + H_2O$.

Insol. in H_2O . Easily sol. in dil. HNO_3 , or boiling $HC_2H_3O_2 + Aq.$ Decomp. by H_2SO_4 , HCl , also by boiling KOH , or $NaOH + Aq.$ Insol. in alcohol. (Herapath, Phil. Mag. (3) **34**. 375.)

Sol. in $NH_4Cl + Aq$; sol. in sat. $NaCl + Aq.$ $2PbO, 3B_2O_3 + 4H_2O$. (Herapath.)

$PbB_4O_7 + 4H_2O$. Slightly sol. in pure H_2O , but insol. in solutions of Na salts as $Na_2B_4O_7 + Aq.$ (Soubeiran.)

Lead borate chloride, $Pb(BO_2)_2, PbCl_2 + H_2O$.

Insol. in cold, very slowly decomp. by hot H_2O into its constituents. Easily sol. in dil. hot $HNO_3 + Aq$; insol. in alcohol. (Herapath, Phil. Mag. (3) **34**. 375.)

Lead borate nitrate, $Pb(BO_2)_2, Pb(NO_3)_2 + H_2O$.

Insol. in alcohol. (Herapath.)

Lithium borate, $LiBO_2$.

Solubility in H_2O .

100 g. H_2O dissolve g. $LiBO_2$ at t° .

t°	g. $LiBO_2$	t°	g. $LiBO_2$
0	0.7	30	4.9
10	1.4	40	11.12
20	2.6	45	20.

(Le Chatelier, C. R. 1897, **124**. 1094.)

Insol. in acetone. (Eidmann, C. C. **1899**, II. 1014; Naumann, B. 1904, **37**. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, **43**. 314.)

+ $8H_2O$. (Le Chatelier, Bull. Soc. 1899, (3) **21**. 35.)

+ $16H_2O$. Effloresces in the air; slowly sol. in cold H_2O , rapidly in hot H_2O . (Le Chatelier, C. R. 1897, **124**. 1092.)

$Li_2H_4(BO_3)_2 + 14H_2O$. (Reischle, Z. anorg. **4**. 166.)

$Li_2B_4O_7$. Deliquescent; easily sol. in H_2O . (Arfvedson, A. ch. **10**. 82.)

Sol. in acetone. (Eidmann, C. C. **1899**, II. 1014.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

+ $5H_2O$. Insol. in alcohol. (Filsinger, Arch. Ph. (3) **8**. 198.)

$3B_2O_3 + 6H_2O$. Very sol. in H_2O ; alcohol. (Filsinger.)

B_2O_3 . Insol. in H_2O . (Le Chatelier, 1899, (3) 21. 35.)

O. Sol. in H_2O ; insol. in alcohol.

"lithium borate" is less sol. than the te. (Gmelin.)

$5B_2O_3 + 10H_2O$. (Dukelski, C. A. 9.)

m borate, $Mg(BO_2)_2$.

C. R. 77. 893.)

Min. *Pinnoite*.

(Laurent, A. ch. (2) 67. 215.)

Insol. in cold or hot H_2O ; easily $[Cl + Aq]$. Decomp. by conc. HCl to H_3BO_3 and $MgCl_2$. (Wöhler.)

$+ 8H_2O$. (Popp, A. Suppl. 8. 1.)

$3B_2O_3 + 8H_2O$. Very slowly sol. in rose, A. 84. 221.)

75 pts. cold H_2O . (Rammelsberg, 445.)

B_2O_3 . Insol. in H_2O , but sol. in $-Aq$. (Guertler, Z. anorg. 1904, 40.)

Very sl. sol. in $\frac{1}{10}$ N $HCl + Aq$. (Hoff, B. A. B. 1907, 658.)

tscharite.

B_2O_3 . Insol. in H_2O ; easily sol. in Ebelmen, A. 80. 208.)

sol. in cold, but somewhat decomp. $g H_2O$. (Rammelsberg.)

Somewhat sol. in cold H_2O . Pogg. 28. 525.)

$2B_2O_3$. Sol. in warm H_2SO_4 or Aq . (Ditte, C. R. 77. 893.)

$3B_2O_3 + 18H_2O = Mg(BO_2)_2, 10HBO_2$. (Rammelsberg, Pogg. 49. 445.)

$4B_2O_3$. Sol. in hot dil. acids; insol. acid. (Ditte, C. R. 77. 893.)

$2B_2O_3 + 1\frac{1}{2}$, and $3H_2O$. Min. e. Difficultly sol. in $HCl + Aq$.

B_2O_3 . (Mallard, C. R. 105. 260.)

m manganous borate, $3Mg_2B_2O_5, 12B_2O_3 + 7H_2O$.

tsusserite. Sol. in $HCl + Aq$.

m potassium borate, $KMg_2B_{11}O_{19} + 3H_2O$.

Kaliborite. Insol. in H_2O . (Feit, 189, 13. 1188.)

$2K_2O, 11B_2O_3 + 20H_2O$. (van't Lichtenstein, B. A. B. 1904, 936.)

m sodium borate, $Mg_2B_6O_{11}, B_4O_7 + 30H_2O$.

scent. About as sol. in cold H_2O as solution separates out a Mg borate ring, which redissolves on cooling. by boiling H_2O . (Rammelsberg.)

m strontium borate, $3MgO, 3SrO, 3B_2O_3$.

sol. in dil. acids. (Ditte, C. R. 77.

Magnesium borate bromide, $2Mg_2B_2O_5, MgBr_2$ or $6MgO, 8B_2O_3, MgBr_2$.

(Rousseau and Allaire, C. R. 1894, 119, 71.)

Magnesium borate chloride, $2Mg_2B_2O_5, MgCl_2$.

Min. *Boracite*. Insol. in H_2O ; slowly sol. in acids. (Kraut.)

Stassfurthite. Easily sol. in warm acids. (Bischof.)

Magnesium borate iodide, $6MgO, 8B_2O_3, MgI_2$.

(Allaire, C. R. 1898, 127. 556.)

Magnesium borate phosphate, $Mg(BO_2)_2, 2MgHPO_4 + 7H_2O$.

Min. *Lunenburgite*.

Magnesium borate sulphate, $2Mg_2B_2O_5, 3MgSO_4 + 12H_2O$.

Min. *Magnesium sulphoborite*.

Sol. in mineral acids when ground. (Nau-pert, B. 1893, 26. 874.)

Manganous borate, MnB_4O_7 (?)

Insol. in H_2O (Berzelius); very sl. sol. in H_2O (Thomas, Am. Ch. J. 4. 358); decomp. by warm, slowly by cold H_2O . Sol. in $MgSO_4 + Aq$ (Berzelius).

$+ 3H_2O$. (Endemann and Paisley, Zeit. angew. Ch. 1903, 16. 176.)

$+ 5H_2O$. Ppt. (Endemann and Paisley.)

Very hygroscopic. (Endemann, Am. Ch. J. 1903, 29. 72.)

$3MnO, B_2O_3$. (Mallard, C. R. 105. 1260.)

Not attacked by H_2O . Very sol. in acids. (Ouvrard, C. R. 1900, 130. 336.)

$3MnO, 2B_2O_3$. (Mallard.)

$MnH_4(BO_2)_2$. Very sl. sol. in H_2O .

Solubility in 2% $Na_2SO_4 + Aq$. At 18.5° , 0.77 g. $MnH_4(BO_2)_2$ are dissolved per litre; at 40° , 0.65 g.; at 60° , 0.36 g.; at 80° , 0.12 g.

Solubility in 2% $NaCl + Aq$. 1 l. solution dissolves 1.31 g. salt at 18.2° ; 0.6 g. at 59° ; and 0.29 g. at 80° .

Solubility in 2% $CaCl_2 + Aq$. 1 l. $CaCl_2 + Aq$ dissolves 2.91 g. salt at 17.6° ; 2.44 g. at 43.0° ; 2.25 g. at 61° ; and 1.35 g. at 80° . (Hartley and Ramage, Chem. Soc. 63. 129.)

Manganous borate bromide, $6MnO, 8B_2O_3, MnBr_2$.

(Rousseau and Allaire, C. R. 1894, 119. 73.)

Manganous borate chloride, $6MnO, 8B_2O_3, MnCl_2$.

(Rousseau and Allaire, C. R. 1894, 118. 1257.)

Molybdenum borate, $MoO_3, 2B_2O_3$ (?)

Insol. in H_2O ; sol. in $H_3BO_3 + Aq$. (Berzelius.)

Molybdenum borate, $\text{Mo}_2\text{O}_3, \text{B}_2\text{O}_3$.

Precipitate. Insol. in H_2O ; sl. sol. in a solution of boric acid. (Berzelius.)

See **Boromolybdic Acid**.

Nickel borate, $\text{Ni}(\text{BO}_2)_2 + 2\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in acids. Easily sol. in warm $\text{NH}_4\text{Cl} + \text{Aq}$. (Rose, Pogg. 88. 299.)

$2\text{NiO}, \text{B}_2\text{O}_3 + x\text{H}_2\text{O}$. Easily sol. in acids. (Rose.)

$3\text{NiO}, 2\text{B}_2\text{O}_3 + 5\text{H}_2\text{O}$. Easily sol. in acids. (Rose.)

$3\text{NiO}, \text{B}_2\text{O}_3$. Not attacked by H_2O ; sol. in acids. (Ouvrard, C. R. 1900, 130. 337.)

Nickel borate bromide, $6\text{NiO}, 8\text{B}_2\text{O}_3, \text{NiBr}_2$.
(Rousseau, C. R. 1894, 119. 73.)**Nickel borate chloride, $6\text{NiO}, 8\text{B}_2\text{O}_3, \text{NiCl}_2$.**
(Rousseau, C. R. 1894, 118. 1257.)**Potassium borates.**

Solubility of B_2O_3 in $\text{K}_2\text{O} + \text{Aq}$ at 30° .

Solution contains		Solid phase
% by wt. K_2O	% by wt. B_2O_3	
47.50	$\text{KOH}, 2\text{H}_2\text{O}$
46.45	0.72	"
46.36	0.91	$\text{K}_2\text{O}, \text{B}_2\text{O}_3, 2.5\text{H}_2\text{O}$
40.51	1.25	"
36.82	1.80	"
36.72	1.85	"
32.74	3.51	"
29.63	6.98	"
26.89	12.12	"
24.84	17.63	"
23.30	18.19	$\text{K}_2\text{O}, 2\text{B}_2\text{O}_3, 4\text{H}_2\text{O}$
16.21	13.10	"
11.78	9.82	"
9.18	8.00	"
6.22	9.13	"
7.79	13.20	"
7.73	13.37	$\text{K}_2\text{O}, 2\text{B}_2\text{O}_3, 4\text{H}_2\text{O} + \text{K}_2\text{O}, 5\text{B}_2\text{O}_3, 8\text{H}_2\text{O}$
7.81	13.28	"
7.67	13.19	"
7.71	13.21	$\text{K}_2\text{O}, 5\text{B}_2\text{O}_3, 8\text{H}_2\text{O}$
7.63	13.28	"
3.42	7.59	"
1.80	4.15	"
0.80	3.05	"
0.51	3.19	"
0.33	4.58	$\text{K}_2\text{O}, 5\text{B}_2\text{O}_3, 8\text{H}_2\text{O} + \text{B}(\text{OH})_3$
0.38	4.51	"
0.31	4.46	"
0.28	4.36	$\text{B}(\text{OH})_3$
.....	3.54	"

At 30° only the three potassium borates $\text{K}_2\text{O}, \text{B}_2\text{O}_3 + 2.5\text{H}_2\text{O}$; $\text{K}_2\text{O}, 2\text{B}_2\text{O}_3 + 4\text{H}_2\text{O}$ and $\text{K}_2\text{O}, 5\text{B}_2\text{O}_3 + 8\text{H}_2\text{O}$ exist in stable form.

(Dukelski, Z. anorg. 1906, 50. 42.)

Potassium metaborate, KBO_2 .

Sol. in small amount of H_2O . (Berzelius, Pogg. 34. 568.)

$+1\frac{1}{4}\text{H}_2\text{O}$. Only stabile hydrate. (Dukelski, Z. anorg. 1906, 50. 42.)

$+1\frac{1}{2}\text{H}_2\text{O}$. (Atterberg, Bull. Soc. (2) 22. 350.)

Potassium tetraborate, $\text{K}_2\text{B}_4\text{O}_7$.

Very sol. in H_2O .

$+4\text{H}_2\text{O}$. (Atterberg, Bull. Soc. (2) 22. 350.)

Only stabile hydrate. (Dukelski, l. c.)

$+5\text{H}_2\text{O}$. Very sol. in H_2O ; more sol. than $\text{K}_2\text{B}_6\text{O}_{10}$ or $\text{K}_2\text{B}_{12}\text{O}_{19}$.

$+6\text{H}_2\text{O}$. (Atterberg, l. c.)

Potassium hexaborate, $\text{K}_2\text{B}_6\text{O}_{10} + 5$, and $8\text{H}_2\text{O}$.

Easily sol. in H_2O .

Does not exist. (Dukelski, l. c.)

Potassium dekaborate, $\text{K}_2\text{B}_{10}\text{O}_{16} + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Rammelsberg.)

Only hydrate. (Dukelski.)

Potassium dodekaborate, $\text{K}_2\text{B}_{12}\text{O}_{18} + 10\text{H}_2\text{O}$.

Sl. sol. in cold, very sol. in hot H_2O . (Laurent, A. ch. 67. 215.)

$= \text{K}_2\text{B}_{10}\text{O}_{16}$. (Rammelsberg.)

Does not exist. (Dukelski.)

Potassium borate fluoride, KBO_2, KF .

Sol. in H_2O . (Schiff and Sestini, A. 228. 72.)

$\text{KBO}_2, 2\text{KF}$. Sol. in little, decomp. by much H_2O . Insol. in H_2O . (Schiff and Sestini, A. 228. 72.)

Rubidium borate, $\text{Rb}_2\text{B}_4\text{O}_7$.

Anhydrous. (Reischle, Z. anorg. 4. 166.)

$+6\text{H}_2\text{O}$. Not deliquescent or efflorescent.

Sol. in H_2O . (Reissig, A. 127. 33.)

Samarium borate, SmBO_3 .

Insol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$. (Cleve, Bull. Soc. (2) 43. 1670.)

Scandium borate, ScBO_3 .

Sol. in dil. acids. (Crookes, Phil. Trans. 1910, 210. A. 364.)

Silver borate, AgBO_2 .

Sl. sol. in H_2O . By washing with H_2O the boric acid is dissolved out. (Rose, Pharm. Centralbl. 1853. 205.)

Sol. with decomp. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ (Herschel); sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$ if pptd. cold.

1 l. H_2O dissolves ca. 6×10^{-2} gram-atoms at 25° . (Abegg and Cox, Z. phys. Ch. 1903, 46. 11.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

$3\text{Ag}_2\text{O}, 4\text{B}_2\text{O}_3$. (Rose, l. c.)

1 borates.

ibility of B₂O₃ in Na₂O+Aq at 30°.

olution contains		Solid phase
wt. ()	% by wt. B ₂ O ₃	
0	NaOH, H ₂ O
5	2.71	"
7	5.10	"
5	5.55	Na ₂ O, B ₂ O ₃ , 4H ₂ O
4	3.73	"
9	2.51	"
1	2.38	"
8	2.44	"
3	2.75	"
8	2.98	"
0	3.82	"
1	13.69	"
8	4.63	Na ₂ O, B ₂ O ₃ , 4H ₂ O + Na ₂ O, B ₂ O ₃ , 8H ₂ O
8	4.69	Na ₂ O, B ₂ O ₃ , 8H ₂ O
1	4.97	"
2	6.21	"
5	8.18	"
9	9.12	"
5	10.49	Na ₂ O, 2B ₂ O ₃ , 10H ₂ O
1	6.94	"
0	4.76	"
8	2.41	"
8	5.16	"
4	7.36	"
2	7.79	"
0	9.48	"
8	17.20	Na ₂ O, 2B ₂ O ₃ , 10H ₂ O + Na ₂ O, 5B ₂ O ₃ , 10H ₂ O
9	15.84	Na ₂ O, 5B ₂ O ₃ , 10H ₂ O
7	13.30	"
6	12.14	"
9	11.84	Na ₂ O, 5B ₂ O ₃ , 10H ₂ O + B(OH) ₃
6	11.78	B(OH) ₃
1	11.18	"
4	6.11	"
.	3.54	"

°, only the four sodium borates Na₂O, 4H₂O; Na₂O, B₂O₃+8H₂O; Na₂O, +10H₂O; and Na₂O, 5B₂O₃+10H₂O; stable phases. (Dukelski, Z. anorg. 1906, 50. 46.)

1 metaborate, NaBO₂.

ndrous. Easily sol. in H₂O, with evolu- heat.

O. Easily sol. in H₂O. (Benedikt.)

I₂O. Easily sol. in H₂O. (Benedikt, 03.)

I₂O. Easily sol. in H₂O. (Berzelius.)

I₂O. Sl. efflorescent. Sol. in hot, less cold H₂O. Melts at 57° in its crystal (Dukelski, Z. anorg. 50. 42.)

2H₂O. (Atterberg, Z. anorg. 1906, 48.)

2H₂O. (Atterberg.)

+8H₂O. (Atterberg.)

+4H₂O and +8H₂O are the only hydrates formed. (Dukelski.)

System Na₂O, B₂O₃, H₂O at 60° investi- gated by Sborgi. (Real. Ac. Linc. 1915, (5) 24. I, 443.)

Sodium tetraborate, Na₂B₄O₇ (Borax).

100 g. H₂O dissolve at:

5° 10° 21.5° 30° 37.5°
1.3 1.6 2.8 3.9 5.6 g. anhydrous salt.

45° 50° 54° 55° 56° 57°
8.1 10.5 13.3 14.2 15.0 16.0 g. anhydrous salt.

(Horn and van Wagener, Am. Ch. J. 1903, 30. 347.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Sol. in amyl alcohol in the presence of meta- arsenious acid and excess of H₂BO₃. (Auer- bach, Z. anorg. 1903, 37. 358.)

+4H₂O.

+5H₂O.

100 g. H₂O dissolve at:

65° 70° 80° 90° 100°
22.0 24.4 31.4 40.8 52.3 g. anhydrous salt.

(Horn and van Wagener, Am. Ch. J. 1903, 30. 347.)

+6H₂O. Grows opaque in the air. (Bechi, Sill. Am. J. (2) 17. 129.)

+10H₂O. Only stabile hydrate. (Dukel- ski, Z. anorg. 50. 30.) Efflorescent on surface in dry air. Not efflorescent when free from Na₂CO₃. (Sims.)

Sol. in 12 pts. cold, and 2 pts. hot H₂O. Sat. cold Na₂B₄O₇+Aq contains 9.23%, and sat. hot Na₂B₄O₇+ Aq contains 33.33% Na₂B₄O₇. (Gmelin.)

Sol. in 20 pts. cold, and 6 pts. boiling H₂O. (Wal- lerius.)

Sol. in 15 pts. H₂O at 18.75°. (Abl.)

100 pts. H₂O at 15.5° dissolve 5 pts.; at 65°, 40 pts.; at 100°, 166 pts. Na₂B₄O₇+10H₂O. (Ure's Dictionary.)

100 pts. sat. Na₂B₄O₇+Aq at 105.5° contain 52.5 pts. Na₂B₄O₇, or 100 pts. H₂O dissolve 110.54 pts. Na₂B₄O₇, or 1 pt. Na₂B₄O₇ is sol. in 0.9047 pt. H₂O at 105.5°. (Griffith, Quar. J. Sci. 18. 90.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. Na ₂ B ₄ O ₇	Pts. Na ₂ B ₄ O ₇ +10H ₂ O	t°	Pts. Na ₂ B ₄ O ₇	Pts. Na ₂ B ₄ O ₇ +10H ₂ O
0	1.49	2.83	60	18.09	40.43
10	2.42	4.65	70	24.22	57.85
20	4.05	7.88	80	31.17	76.19
30	6.00	11.90	90	40.14	116.66
40	8.79	17.90	100	55.16	201.43
50	12.93	27.41

(Poggiale, A. ch. (3) 8. 46.)

100 pts. H₂O dissolve 1.4 pts. Na₂B₄O₇ at 0°, and 55.3 pts. at 100°. (Mulder.)

Na₂B₄O₇+Aq sat. at 15° has sp. gr.= 1.0199, and contains 3.926 pts. Na₂B₄O₇ to 100 pts. H₂O. (Michel and Krafft, A. ch. (3) 41. 471.)

$\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$ sat. at 17° has sp. gr. = 1.0208. (Stolba, J. pr. 97. 503.)

Sp. gr. of $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$ at 15° .

$\frac{\% \text{Na}_2\text{B}_4\text{O}_7}{\text{Na}_2\text{B}_4\text{O}_7 + 10\text{H}_2\text{O}}$	Sp. gr.	$\frac{\% \text{Na}_2\text{B}_4\text{O}_7}{\text{Na}_2\text{B}_4\text{O}_7 + 10\text{H}_2\text{O}}$	Sp. gr.
1	1.0049	4	1.0199
2	1.0099	5	1.0249
3	1.0149	6	1.0299

(Gerlach, Z. anal. 28. 473.)

Sp. gr. of $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$ sat. at $15^\circ = 1.032$. (Gerlach.)

Sat. $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$ boils at 105.5° , and contains 110.5 pts. $\text{Na}_2\text{B}_4\text{O}_7$ to 100 pts. H_2O . (Griffith.)

Sat. $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$ forms a crust at 103° , and contains 60.14 pts. $\text{Na}_2\text{B}_4\text{O}_7$ to 100 pts. H_2O ; highest temp. observed, 104.3° . (Gerlach, Z. anal. 26. 427.)

B.-pt. of $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq}$ containing pts. $\text{Na}_2\text{B}_4\text{O}_7$ to 100 pts. H_2O .

B.-pt.	Pts. $\text{Na}_2\text{B}_4\text{O}_7$	B.-pt.	Pts. $\text{Na}_2\text{B}_4\text{O}_7$
100.5°	8.64	103.0°	61.2
101.0	17.2	103.5	75.4
101.5	26.5	104.0	90.8
102.0	37.5	104.5	109.0
102.5	48.5	104.6	112.3

(Gerlach, Z. anal. 26. 452.)

M.-pt. of $\text{Na}_2\text{B}_4\text{O}_7 + 10\text{H}_2\text{O}$ is 75.5° . (Tilden, Chem. Soc. 45. 407.)

Insol. in alcohol.

100 g. alcohol (0.941 sp. gr.) dissolve 2.48 g. at 15.5° (U. S. P.).

Sol. in alcoholic solution of $\text{NaC}_2\text{H}_3\text{O}_2$. (Stromeyer.)

Sol. in 14.7 pts. glycerine of 1.225 sp. gr. (Vogel.)

Sol. in 1 pt. glycerine. (Schultze, Arch. Pharm. (3) 6. 149.)

100 g. glycerine dissolve 60.3 g. at 15.5° (U. S. P.)

Min. Tincal.

Sodium borate, $\text{Na}_2\text{B}_4\text{O}_7 + 10\text{H}_2\text{O}$.

Sol. in 5-6 pts. cold H_2O . (Bolley, A. 68. 122.) Perhaps sodium hydrogen tetraborate $\text{NaHB}_4\text{O}_7 + 4\frac{1}{2}\text{H}_2\text{O}$.

$\text{Na}_2\text{B}_{10}\text{O}_{16} + 10\text{H}_2\text{O}$. Decomp. by H_2O . (Atterberg, Z. anorg. 48. 370.)

Stabile. (Dukelski, Z. anorg. 50. 42.)

+11 H_2O . (Laurent, C. R. 29. 5.)

Sodium borate fluoride, $\text{NaBO}_2 \cdot 3\text{NaF} + 4\text{H}_2\text{O}$.

Sol. in H_2O .

Basarow (B. 7. 112) considers this salt to be a mixture.

$\text{Na}_2\text{B}_4\text{O}_7$, $12\text{NaF} + 22\text{H}_2\text{O}$. Can be arated into its constituents by H_2O . (Zelius, Berz. J. B. 23. 96.)

Strontium borate, $\text{Sr}(\text{BO}_3)_2$.

(Ditte, C. R. 77. 788.)

Easily hydrated by H_2O forming SrO , 1 +2 H_2O . Very sol. in dil. acetic acid. (Ouvrard, C. R. 1906, 142. 282.)

Insol. in acetone. (Naumann, B. 1904 4329.)

+2 H_2O . (Ouvrard, l. c.)

+4 H_2O . (Ouvrard, l. c.)

+5 H_2O . 1 l. H_2O dissolves 2.3 g. at (Ditte, A. ch. 1883 (5) 30. 253.)

SrB_4O_7 . Insol. in H_2O ; sol. in dil. a (Guertler, Z. anorg, 1904, 40. 243.)

+4 H_2O . Sol. in 130 pts. boiling H_2O . pts. H_2O at 100° dissolve 7.7 pts. (Ure's D.

Easily sol. in cold NH_4 salts + Aq; sol. in HNO_3 + Aq.

2 SrO , B_2O_3 . Easily decomp. by H_2O forming B_2O_3 , SrO , 4 H_2O . Very sol. in a (Ouvrard, C. R. 1906, 142. 282.)

3 SrO , B_2O_3 . Less easily attacked by than Ca comp. Very sol. in mineral a Sl. attacked by dil. acetic acid. (Ouv C. R. 1901, 132. 258.)

$\text{SrB}_6\text{O}_{10}$. Very sl. sol. in H_2O ; sol. in a (Laurent.)

$\text{SrB}_6\text{O}_{11} + 7\text{H}_2\text{O}$. Ppt. (Laurent.)

+12 H_2O . (Ditte.)

$\text{Sr}_2\text{B}_4\text{O}_9$. Sol. in cold mineral acids acetic acid. (Ditte, C. R. 77. 785.)

2 SrO , 3 B_2O_3 . Easily sol. in acids. (I l. c.)

Strontium borate bromide, 3 SrO , 51 SrBr_2 .

As the chloride. (Ouvrard, C. R. 142. 283.)

Strontium borate chloride, 3 SrO , 51 SrCl_2 .

Sl. attacked by cold H_2O , not attacked dilute acetic acid. (Ouvrard, C. R. 1906, 282.)

Thallous borate, $\text{TlBO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Decomp. in the air. (Buchtala, J. pr. (2) 88. 784.)

$\text{Tl}_2\text{B}_2\text{O}_7$. (Buchtala.)

$\text{Tl}_2\text{B}_4\text{O}_7$. Ppt. Sol. in boiling H_2O ; in cold dil. H_2SO_4 + Aq. (Crookes.)

+2 H_2O . (Buchtala, J. pr. 1913 (2 774.)

$\text{Tl}_2\text{B}_6\text{O}_{10} + 3\text{H}_2\text{O}$. (Buchtala.)

$\text{Tl}_2\text{B}_8\text{O}_{13} + 4\text{H}_2\text{O}$. (Buchtala.)

$\text{Tl}_2\text{B}_{10}\text{O}_{16} + 8\text{H}_2\text{O}$. (Buchtala.)

$\text{Tl}_2\text{B}_{12}\text{O}_{19} + 7\text{H}_2\text{O}$. (Buchtala.)

Thorium borate (?).

Precipitate. Insol. in H_2O and H_2E Aq. (Berzelius.)

Tin (stannous) borate (?).

Ppt. (Wenzel.)

idyl borate.

l. in H_2O ; sol. in $H_3BO_3 + Aq.$ (Ber-

um borate, $YbBO_3$.

l. in conc. HCl ; sol. in HF . (Cleve, Z. 1902, **32**. 148.)

n borate.

ipitate. (Berlin, Pogg. **43**. 105.)

orate, $3ZnO, 2B_2O_3$.

llard, C. R. **105**. 1260.)

omp. by H_2O ; very sol. in dil. acids. rd, C. R. 1900, **130**. 336.)

, $2B_2O_3 + 4H_2O$. Sol. in H_2O with p. (Ditte, A. ch. 1883, (5) **30**. 256.)

, $4B_2O_3 + H_2O$. Ppt. (Holdermann, Pharm. 1904, **242**. 567.)

, $4B_2O_3 + 10H_2O$. (Ditte, A. ch. 1883, 256.)

, $4B_2O_3 + 9H_2O$. Sl. sol. in H_3BO_3 (Rose, Pogg. **88**. 299.)

, B_2O_3 . Insol. in mineral acids. (le ier, C. R. **113**. 1034.)

orate ammonia, $ZnB_4O_7, 4NH_3 + 6H_2O$.

ly sol. in $NH_4OH, HC_2H_3O_2, H_2SO_4$, and $HNO_3 + Aq.$ (Büchner, A. **151**.

orate bromide, $6ZnO, 8B_2O_3, ZnBr_2$.

isseau and Allaire, C. R. **116**. 1446.)

orate chloride, $6ZnO, 8B_2O_3, ZnCl_2$.

l. in HCl . (Rousseau, C. R. 1894, **118**.

orate iodide, $6ZnO, 8B_2O_3, ZnI_2$.

ire, C. R. 1898, **127**. 556.)

um borate, (?).

l. in H_2O .

ic acid.

Perboric Acid.

phosphoric acid.

Phosphoboric acid.

tungstic acid.

Borotungstic acid.

acid sulphur trioxide.

Borosulphuric acid.

de, $B_2(NH)_3$.

omp. by H_2O ; insol. in all indifferent s; sol. in liquid $NH_3 + S$ to form a dark lution. (Stock, B. 1901, **34**. 3044.)

de hydrochloride, $B_2(NH)_3, 3HCl$.

omp. by H_2O ; insol. in all ordinary or- olvents. (Stock, B. 1901, **34**. 3045.)

Borofluorhydric acid, $HBFl_4$.

See Fluoboric acid.

Borofluorides.

See Fluoborides.

Boromolybdic acid.

Sol. in H_2O . Decomp. by alcohol. (Ber- zelius.)

Boron, B.

(a) *Amorphous*. Somewhat sol. in pure H_2O , when not ignited. Salts and acids sep- arate it out of aqueous solution. Upon evap- oration of H_2O solution a crust is formed, which is only partially sol. in H_2O . (Ber- zelius, Pogg. **2**. 113.) Decomp. by hot H_2SO_4 and cold moderately conc. $HNO_3 + Aq.$ Strongly ignited amorphous B is much less easily attacked by reagents than freshly pptd., and is insol. in H_2O . (Berzelius.) Insol. in caustic alkalies + Aq ; also in alcohol and ether.

Above boron was very impure. (Moissan, C. R. **114**. 392.)

Pure B is not attacked by acids, but has a strong reducing action on $KMnO_4 + Aq, FeCl_3 + Aq$, etc. (Moissan, C. R. **114**. 617.)

Does not melt at 1500° . Readily sol. in conc. acids, as H_2SO_4, HNO_3, H_3PO_4 ; very sl. sol. in hydracids; decomp. H_2O at red heat. (Moissan, A. ch. 1895, (7) **6**. 313-14.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 827.)

(b) *Crystallized*. 1. Insol. in H_2O, HCl , or $KOH + Aq$. Very slightly and slowly attacked by boiling conc. H_2SO_4 . Gradually sol. in hot conc. HNO_3 . Formula is Al_2B_{14} . (Hampe, A. **183**. 75.)

2. Very slightly attacked by conc. HCl or H_2SO_4 ; slowly but completely sol. in conc. HNO_3 ; insol. in $KOH + Aq$. Formula is $C_2Al_2B_{18}$. (Hampe.)

$C_2Al_2B_{14}$. *Crystalline*. Insol. in a solution of CrO_3 in H_2SO_4 . Insol. in hot conc. HCl and H_2SO_4 . Sol. in hot conc. HNO_3 . (Biltz, B. 1910, **43**. 303.)

Boron tribromide, BBr_3 .

Sol. in H_2O or alcohol with decomp. (Nicklès, C. R. **60**. 800.)

Boron phosphorus bromide, BBr_3, PBr_3 .

Decomp. by H_2O .

Sol. in CS_2 , and $CHCl_3$. Decomp. by al- cohol, ether, etc. (Tarible, C. R. **116**. 1521.)

BBr_3, PBr_3 . Sl. sol. in cold, easily in hot CS_2 . (Tarible.)

Boron bromide ammonia, $BBr_3, 4NH_3$.

Decomp. by H_2O and alkalies. (Besson, C. R. **114**. 542.)

Boron bromide phosphine, BBr_3, PH_3 .

Violently decomp. by H_2O . (Besson, C. R. **113**. 78.)

Boron bromide phosphorus trichloride,
 $2\text{BBr}_3, \text{PCl}_3$.

Decomp. by H_2O . Sol. in BBr_3 , PCl_3 , CS_2 , and CHCl_3 . Insol. in petroleum ether. (Tarible, C. R. 1901, 132. 84.)

Boron bromide phosphorus pentachloride,
 $2\text{BBr}_3, \text{PCl}_5$.

Sol. in BBr_3 and CS_2 ; decomp. by H_2O ; insol. in light petroleum. (Tarible, C. R. 1901, 132. 85.)

Boron bromide phosphorus diiodide, 2BBr_3 ,
 P_2I_4 .

Sol. in BBr_3 , CS_2 , CHCl_3 ; insol. in light petroleum; decomp. by H_2O . (Tarible, C. R. 1901, 132. 205.)

Boron bromide phosphoryl chloride, BBr_3 ,
 POCl_3 .

Very easily decomp. (Oddo and Tealdi, Gazz. ch. it. 1903, 33. (2) 431.)

Boron bromoiodide, BBr_2I .

Decomp. violently by H_2O . (Besson, C. R. 112. 100.)

BBrI_2 . (Besson, C. R. 112. 100.)

Boron bromosulphide, $\text{B}_2\text{S}_3, \text{BBr}_3$.

Decomp. by H_2O . (Stock, B. 1901, 34. 3040.)

Boron carbide, B_4C .

Very stable; insol. in HF and in HNO_3 ; sol. in KOH at red heat. (Moissan, Bull. Soc. 1894, (3) 11. 1101.)

Insol. in acids; sol. in fused alkali. (Moissan, C. R. 1894, 118. 559.)

BC or B_2C_2 . Insol. in all the usual solvents. (Müllhäuser, Z. anorg. 5. 92.)

Boron trichloride, BCl_3 .

Rapidly absorbed by H_2O and alcohol with decomposition.

Boron nitrosyl chloride, $\text{BCl}_3, \text{NOCl}$.

Decomp. violently by H_2O . (Geuther, J. pr. (2) 8. 854.)

Boron phosphoryl chloride, $\text{BCl}_3, \text{POCl}_3$.

Decomp. immediately by H_2O . (Gustavson, Zeit. Chem. 1870. 521.)

Boron chloride ammonia, $2\text{BCl}_3, 3\text{NH}_3$.

Decomp. by H_2O . (Berzelius, Pogg. 2. 147.)

Boron chloride phosphine, $\text{BCl}_3, \text{PH}_3$.

Decomp. by H_2O . (Besson, C. R. 110. 516.)

Boron chlorosulphide, $\text{B}_2\text{S}_3, \text{BCl}_3$.

Decomp. by H_2O . (Stock, B. 1901, 34. 3040.)

Boron trifluoride, BF_3 .

H_2O absorbs 700 vols. BF_3 gas to form a

liquid of 1.77 sp. gr. On boiling, $\frac{1}{3}$ of the BF_3 is given off, and a residue boiling at 165–200° with composition $\text{BF}_3 + 2\text{H}_2\text{O}$ or $\text{HBO}_2 + 3\text{HF}$, is left. (J. Davy, A. ch. 86. 178.)

1 ccm. H_2O absorbs at 0° and 762 mm. pressure 1.057 ccm. BF_3 .

1 vol. conc. H_2SO_4 of 1.85 sp. gr. absorbs 50 vols. BF_3 .

Absorbed by alcohol with decomp.

Cold oil of turpentine absorbs 6.8% of BF_3 .

Boron fluoride ammonia, BF_3, NH_3 , BF_3 ,
 2NH_3 , and $\text{BF}_3, 3\text{NH}_3$.

Decomp. by H_2O .

Boron fluoride cyanhydric acid, BF_3, HCN .

Very unstable. (Patein, C. R. 113. 85.)

Boron fluoride phosphine, $2\text{BF}_3, \text{PH}_3$.

Very unstable at ordinary temp. Decomp. by H_2O . (Besson, C. R. 110. 80.)

Boron hydride, BH_3 .

Not obtained free from H . Sl. sol. in H_2O . (Jones, Chem. Soc. 35. 41.)

See *Cyclotriborene*.

B_4H_{10} . B.-pt. 16–17° at 760 mm.

Very unstable. Takes fire spontaneously in the air.

Decomp. by H_2O , dil. HCl , and oxidized by conc. HNO_3 with explosive violence.

Absorbed by $\text{NaOH} + \text{Aq}$.

Decomp. by alcohol. Sol. in dry benzene. (Stock, B. 1912, 45. 3562.)

B_6H_{12} . B.-pt. 100° at atmospheric pressure.

Decomp. by H_2O . With aqueous alkalis, hydrogen is evolved. (Stock, B. 1912, 45. 3565.)

B_2H_6 . Insol. in HCl . Sol. in aqua regia and $\text{Br}_2 + \text{Aq}$. (Winkler, B. 1890, 23. 778.)

$\text{B}_{10}\text{H}_{14}$. M.-pt. 99.5°; not attacked by cold or boiling H_2O . Sol. in dil. $\text{NaOH} + \text{Aq}$. Sol. in alcohol, ether, benzene, and CS_2 . (Stock, B. 1913, 46. 3360.)

Boron iodide, BI_3 .

Very hygroscopic, and instantly decomp. by H_2O or alcohol. Very sol. in CS_2 , CCl_4 , C_6H_6 ; less sol. in PCl_3 , AsCl_3 , and a great many organic liquids. (Moissan, C. R. 112. 717.)

Boron iodide ammonia, $\text{BI}_3, 5\text{NH}_3$.

Decomp. by H_2O . (Besson, C. R. 114. 542.)

Boron iodophosphide, BI_2P .

Very hygroscopic; decomp. by H_2O . Not attacked by cold conc. H_2SO_4 , even if fuming, but on heating decomposition takes place. Very sl. sol. in CS_2 . Insol. in benzene, PCl_3 , or CCl_4 . (Moissan, C. R. 113. 624.)

BIP . Less hygroscopic than BI_2P , but otherwise the properties are similar. (Moissan.)

nitride, BN.

in H_2O , conc. HNO_3 , conc. HCl + conc. solutions of alkalies.

mp. by hot conc. H_2SO_4 or HF . (r, A. 74. 70.)

trioxide, B_2O_3 .

luculent. Sol. in H_2O with a large increase in temp. (Ditte, C. R. 85. 1069.)

dissolves—

at 18.75° in 47.01 pts. H_2O .

“ 25° “ 27.75 “ “

“ 37.5° “ 18.73 “ “

“ 50° “ 15.13 “ “

“ 62.5° “ 9.29 “ “

“ 75° “ 7.28 “ “

“ 87.5° “ 5.58 “ “

“ 100° “ 4.74 “ “

pts. H_2O dissolve—

at 18.75° 2.13 pts. B_2O_3 .

“ 25° 3.60 “ “

“ 37.5° 4.24 “ “

“ 50° 6.61 “ “

“ 62.5° 10.76 “ “

“ 75° 13.73 “ “

“ 87.5° 17.92 “ “

“ 100° 21.09 “ “

es and Firnhaber, Arch. Pharm. 7. 50.)

e H_2O dissolves—

at 0° 11.00 g. B_2O_3 .

“ 12° 16.50 “ “

“ 20° 22.49 “ “

“ 40° 39.50 “ “

“ 62° 64.50 “ “

“ 80° 95.00 “ “

“ 102° 164.50 “ “

(Ditte, C. R. 85. 1069.)

H_2O solution boils at 100° . (Brandes and Firnhaber.)

H_2O solution boils at 103.3° . (Griffith, J. Sci. 18. 90.)

in acetic acid, hot conc. HCl + Aq, and H_2SO_4 . From the three latter it separates on cooling or dilution with H_2O .

Solubility in Na_2O + Aq at 30° .

Borates, sodium.

Solubility in K_2O + Aq at 30° .

Borates, potassium.

in hot glacial acetic acid. (Holt, Soc. 1911, 100. (2) 720.)

in alcohol. (Graham.)

in alcohol. (Berzelius, Ebelmen.)

in oils.

Iso Boric acid.

trioxide potassium fluoride, $B_2O_3, 2KF$.

usually sol. in H_2O . Decomp. by much H_2O . Insol. in alcohol. (Schiff and Sestini, 82.)

Oxychloride, $BOCl$.

Lawson, Zeit. Chem. 1870. 521.)

slowly decomp. by H_2O . (Counsell, pr. (2) 18. 399.)

Oxychlorides of either the above formulæ do not exist; the true formula for boron oxychloride is B_2O_3, Cl_2 . (Lorenz, A. 247. 226.)

Boron phosphide, BP.

Insol. in H_2O . Sol. in conc. boiling alkalies + Aq with decomp. Decomp. by HNO_3 + Aq. (Besson, C. R. 113. 78.)

Insol. in PCl_3 , $AsCl_3$, $SbCl_3$, CCl_4 , and in fact in all known solvents.

Not attacked by boiling H_2O , conc. HCl , or HI + Aq. Sol. in conc. HNO_3 with decomp. on heating. Not attacked by cold H_2SO_4 . (Moissan, C. R. 113. 726.)

B_2P_3 . Not attacked by boiling conc. HNO_3 + Aq. Insol. in all solvents. (Moissan.)

Boron phosphoiodide.

See Boron iodophosphide.

Boron selenide, B_2Se_3 .

Violently decomp. by H_2O . (Sabatier, C. R. 112. 1000.)

Boron trisulphide, B_2S_3 .

Decomp. with violence with H_2O . Combines with alcohol and ether. (Freymy, A. ch. (3) 38. 312.)

Insol. in most solvents, but sl. sol. in PCl_3 without decomp.; more sol. in SCl_3 , but does not crystallize from the solution. (Moissan, C. R. 115. 203.)

Boron trisulphide ammonia, $B_2S_3, 6NH_3$.

Ppt. (Stock, B. 1901, 34. 3042.)

Boron pentasulphide, B_2S_5 .

Decomp. by H_2O and alcohol. (Moissan, C. R. 115. 271.)

Borosulphuric acid, $BOHSO_4 + SO_3$.

Decomp. by H_2O . (Schultz-Sellac, B. 4. 12.)

$B(HSO_4)_3$. Very deliquescent. Easily sol. in fuming H_2SO_4 . (D'Arcy, Chem. Soc. 55. 155.)

$SO_3(O.BO)_2$. Hygroscopic. Deliquescent. Sol. in H_2O with decomp. Decomp. by cold alcohols. (Pictet, Bull. Soc. 1908, (4) 3. 1121.)

$(SO_3)_2B_2O_3$. Hygroscopic. Deliquescent. Sol. in H_2O with decomp. Decomp. by cold alcohols. (Pictet, Bull. Soc. 1908, (4) 3. 1121.)

Borononotungstic acid, $H_4B_2W_2O_{22} + 22H_2O = 9WO_3, B_2O_3, 2H_2O + 22H_2O$.

Sol. in less than $1/2$ pt. H_2O , and as easily sol. in alcohol and ether. Sp. gr. of aqueous solution is somewhat under 3. (Klein, A. ch. (5) 28. 370.)

Aluminum borononotungstate, $Al_4(B_2W_2O_{22})_2 + 65H_2O$.

Extremely sol. in H_2O . (Klein.)

Ammonium —, $(NH_4)_4B_2W_2O_{22} + 18H_2O$.

Quickly effloresces. (Klein.)

Barium boronotungstate, $\text{Ba}_2\text{B}_2\text{W}_8\text{O}_{32} + 19\text{H}_2\text{O}$.

Sol. in 4 pts. cold, and less than $\frac{1}{2}$ pt. hot H_2O . (Klein.)

Cadmium —, $\text{Cd}_2\text{B}_2\text{W}_8\text{O}_{32} + 18\text{H}_2\text{O}$.

Deliquescent.

100 pts. of salt dissolve in less than 8 pts. H_2O at 19° . Sp. gr. of solution is 3.28. (Klein.)

Sp. gr. of sat. solution at $15.6^\circ/4^\circ = 3.2887$; at $16.2^\circ/4^\circ = 3.2868$. (Kahlbaum, Z. anorg. 1902, 29. 229.)

Calcium —, $\text{Ca}_2\text{B}_2\text{W}_8\text{O}_{32} + 15\text{H}_2\text{O}$.

Sol. in $\frac{1}{10}$ pt. H_2O . Solution has sp. gr. = 3.10. (Klein.)

Cerium —, $\text{Ce}_4(\text{B}_2\text{W}_8\text{O}_{32})_3 + 57\text{H}_2\text{O}$.

Very sol. in H_2O ; sp. gr. of solution is over 3.

Chromium —, $\text{Cr}_4(\text{B}_2\text{W}_8\text{O}_{32})_3 + 74\text{H}_2\text{O}$.

Very sol. in H_2O ; sp. gr. of solution is 2.80. (Klein.)

Cobalt —, $\text{Co}_2\text{B}_2\text{W}_8\text{O}_{32} + 18\text{H}_2\text{O}$.

Very sol. in H_2O ; sp. gr. of solution sat. at $19^\circ = 3.36$. (Klein.)

100 pts. H_2O dissolve 306.8 pts. anhydrous salt at 16.2° ; 288 pts. at 18.5° ; 299.7 pts. at 19.6° ; 286 pts. at 21.8° .

Sp. gr. of solution sat. at $19.2^\circ/4^\circ = 3.1369$. (Kahlbaum, Z. anorg. 1902, 29. 218.)

Copper —, $\text{Cu}_2\text{B}_2\text{W}_8\text{O}_{32} + 19\text{H}_2\text{O}$.

25 pts. H_2O dissolve 100 pts. salt. Sp. gr. of solution = 2.6. (Klein.)

Lead —, $\text{Pb}_2\text{B}_2\text{W}_8\text{O}_{32} + 11\text{H}_2\text{O}$.

Sl. sol. in cold, easily sol. in hot H_2O . (Klein.)

Lithium —, (?).

Very sol. in H_2O . Sp. gr. of solution is about 3.

Magnesium —, $\text{Mg}_2\text{B}_2\text{W}_8\text{O}_{32} + 22\text{H}_2\text{O}$.

Very sol. in H_2O . (Klein.)

Manganous —, $\text{Mn}_2\text{B}_2\text{W}_8\text{O}_{32} + 17\text{H}_2\text{O}$.

100 pts. dissolve in 13 pts. H_2O . Sp. gr. of solution at $19^\circ = 3.15$. (Klein.)

Mercurous —, $3\text{Hg}_2\text{O}$, B_2O_3 , $9\text{WO}_3 + 14\text{H}_2\text{O}$ (?).

Precipitate.

Insol. in H_2O . (Klein.)

Sol. in 20,000 pts. dil. cold, and 1000 pts. boiling $\text{HNO}_3 + \text{Aq}$ of 1.42 sp. gr.

Nickel —, $\text{Ni}_2\text{B}_2\text{W}_8\text{O}_{32} + 18\text{H}_2\text{O}$.

Very sol. in H_2O ; sp. gr. of sat. solution at $19^\circ = 3.32$.

100 pts. H_2O dissolve 261.6 pts. at 21.2° .

Sp. gr. $15.75^\circ/4^\circ$ of solution = 2.2959. (Kahlbaum, Z. anorg. 1902, 29. 218.)

Potassium boronotungstate, $\text{K}_4\text{B}_2\text{W}_8\text{O}_{32} + 13\text{H}_2\text{O}$.

5 pts. salt dissolve in 8 pts. H_2O at 19° form a solution of 1.38 sp. gr. The solution sat. at 100° has sp. gr. of over 2. (Klein.)

Silver —, $\text{Ag}_4\text{B}_2\text{W}_8\text{O}_{32} + 14\text{H}_2\text{O}$.

Very sl. sol. in H_2O .

Sodium —, $\text{Na}_2\text{H}_2\text{B}_2\text{W}_8\text{O}_{32} + 23\text{H}_2\text{O}$.

Very sol. in H_2O . Solution sat. at 19° contains 84 pts. salt to 16 pts. H_2O . (Klein.)

$\text{Na}_4\text{B}_2\text{W}_8\text{O}_{32} + 12\text{H}_2\text{O}$. Sol. in less than 1 pt. H_2O .

Thallium —, $\text{Tl}_2\text{B}_2\text{W}_8\text{O}_{32} + 5\text{H}_2\text{O}$.

Sl. sol. in hot H_2O and nearly insol. in H_2O . (Klein.)

Uranyl —, $(\text{UO}_2)_2(\text{B}_2\text{W}_8\text{O}_{32})_2 + 30\text{H}_2\text{O}$.

Very sol. in H_2O . (Klein.)

Sp. gr. of solution = 3.1.

Zinc —, $\text{Zn}_2\text{B}_2\text{W}_8\text{O}_{32} + 2\text{H}_2\text{O}$.

Very sol. in H_2O . Sp. gr. of solution = 3. (Klein.)

Borodecitungstic acid.

Barium borodecitungstate, $\text{Ba}_2\text{B}_2\text{W}_{10}\text{O}_{40} + 20\text{H}_2\text{O}$.

Sol. in H_2O . (Klein, C. R. 99. 35.)

Boroduodecitungstic acid, $\text{H}_2\text{B}_2\text{W}_{12}\text{O}_{48} + 4\text{H}_2\text{O}$, B_2O_3 , 12WO_3 .

Known only in solution, which decomposes into boronotungstic acid and tungstic acid when evaporated to a certain concentration. (Klein, C. R. 99. 35.)

Barium potassium boroduodecitungstate, 3BaO , K_2O , B_2O_3 , $12\text{WO}_3 + 28\text{H}_2\text{O}$.

Potassium —, $\text{K}_2\text{B}_2\text{W}_{12}\text{O}_{48} + 21\text{H}_2\text{O}$.

Sol. in H_2O . (Klein.)

$2\text{K}_2\text{O}$, 12WO_3 , $\text{B}_2\text{O}_3 + 18\text{H}_2\text{O}$. Sol. in 1 pt. H_2O . (Klein.)

Boroquatuordecitungstic acid, $\text{H}_{12}\text{B}_2\text{W}_{14}\text{O}_{60} + 6\text{H}_2\text{O}$, B_2O_3 , 14WO_3 .

Has not been obtained in the free state. (Klein, A. ch. (5) 28. 353.)

Barium boroquatuordecitungstate,

$\text{Ba}_3\text{B}_2\text{W}_{14}\text{O}_{60} = 3\text{BaO}$, B_2O_3 , $14\text{WO}_3 + 5\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Klein.)

Barium sodium —, $3\frac{1}{2}\text{BaO}$, $1\frac{1}{2}\text{Na}_2\text{O}$, $5\text{H}_2\text{O}$, B_2O_3 , $14\text{WO}_3 + 29\text{H}_2\text{O}$.

Potassium —, $3\text{K}_2\text{O}$, H_2O , B_2O_3 , $14\text{WO}_3 + 22\text{H}_2\text{O}$.

Sol. in H_2O . (Klein.)

Silver —, $\text{Ag}_6\text{H}_2\text{B}_2\text{W}_{14}\text{O}_{60} + 7\text{H}_2\text{O}$.

Nearly insol. in cold H_2O . (Klein.)

Sodium —, $\text{Na}_4\text{H}_4\text{B}_2\text{W}_{14}\text{O}_{60} + 29\text{H}_2\text{O}$.

Sol. in H_2O . (Klein.)

Strontium boroquatordécitungsstate, $\text{SrO}, 1\frac{1}{2}\text{Na}_2\text{O}, \text{B}_2\text{O}_3, 14\text{WO}_3 + 29\text{H}_2\text{O}$.
p. by H_2O . (Klein.)

Trigintitungsstic acid.

Boroundevigintitungsstate, $4\text{BaO}, \text{B}_2\text{O}_3, 19\text{WO}_3 + 30\text{H}_2\text{O}$.

p. by H_2O . (Ebenhusen, 1905.)

Tuoretvigintitungsstic acid, $\text{B}_2\text{O}_3, 24\text{WO}_3 + 66\text{H}_2\text{O}$.

Insol. Somewhat more sol. in H_2O .
 $\text{B}_2\text{O}_3, 28\text{WO}_3 + 62\text{H}_2\text{O}$. Also more sol. in H_2O . (Copaux, C. R. 1908, 147. 975.)

Boroquatruoretvigintitungsstate, $\text{B}_2\text{O}_3, 24\text{WO}_3 + 54\text{H}_2\text{O}$.

p. by H_2O . (Copaux, C. R. 1908, 147. 217.)

$\text{B}_2\text{O}_3, 24\text{WO}_3 + 58\text{H}_2\text{O}$. (Copaux, C. R. 1908, 147. 217.)

—, $5\text{CdO}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 54\text{H}_2\text{O}$.

p. by H_2O . (Copaux, l. c.)

—, $5\text{CaO}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 44\text{H}_2\text{O}$.
Insol. in H_2O . (Copaux, l. c.)

—, $15\text{Li}_2\text{O}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 38\text{H}_2\text{O}$.
Insol. in H_2O . (Copaux, l. c.)

m —, $5\text{MgO}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 44\text{H}_2\text{O}$.

Insol. in H_2O . (Copaux, l. c.)

s —, $9\text{Hg}_2\text{O}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 54\text{H}_2\text{O}$.

Insol. in H_2O . (Copaux, l. c.)

l —, $5\text{K}_2\text{O}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 44\text{H}_2\text{O}$.

Insol. in H_2O . (Copaux, l. c.)

—, $5\text{Na}_2\text{O}, \text{B}_2\text{O}_3, 24\text{WO}_3 + 54\text{H}_2\text{O}$.
Insol. in H_2O . (Copaux, l. c.)

Quetvigintitungsstic acid.

Boroquinquetvigintitungsstate, $\text{B}_2\text{O}_3, 25\text{WO}_3 + 34\text{H}_2\text{O}$. (Ebenhusen, 1905.)

Tetrigintatungsstic acid, $\text{B}_2\text{O}_3, 24\text{WO}_3 + 62\text{H}_2\text{O}$.

Insol. in boiling aqueous solution. (Copaux, C. R. 1908, 147. 975.)

Boroduodetrigintatungsstate, $6\text{K}_2\text{O}, \text{B}_2\text{O}_3, 28\text{WO}_3 + 42\text{H}_2\text{O}$.

p. by boiling alkalis. (Copaux, C. R. 1908, 147. 217.)

Antimonadic acid.

Insol. in H_2O . Easily decomp. (Guyard, C. R. 1908, 147. 354.)

Metabromantimonic acid, $\text{HSbBr}_3 + 3\text{H}_2\text{O}$.

Very hygroscopic. Loses Br_2 in the air. Decomp. by H_2O with separation of antimonic acid. (Weinland, B. 1903, 36. 256.)

Ammonium metabromantimonate, $\text{NH}_4\text{SbBr}_3 + \text{H}_2\text{O}$.

Loses Br_2 in the air. Decomp. by H_2O . (Weinland, l. c.)

Iron (ferric) metabromantimonate, $\text{Fe}(\text{SbBr}_3)_3 + 14\text{H}_2\text{O}$.

Very hygroscopic. Decomp. by H_2O . (Weinland, l. c.)

Lithium metabromantimonate, $\text{LiSbBr}_3 + 4\text{H}_2\text{O}$.

Very hygroscopic. Loses Br_2 in the air. Decomp. by H_2O . (Weinland, l. c.)

Nickel metabromantimonate, $\text{Ni}(\text{SbBr}_3)_2 + 12\text{H}_2\text{O}$.

Hygroscopic. Decomp. by H_2O . (Weinland, l. c.)

Potassium metabromantimonate, $\text{KSbBr}_3 + \text{H}_2\text{O}$.

Loses Br_2 in the air. Decomp. by H_2O . (Weinland, l. c.)

Bromarsenious acid.

See Arsenyl bromide.

Bromauric acid, $\text{HAuBr}_4 + 3\text{H}_2\text{O}$.

(Lengfeld, Am. Ch. J. 1901, 26. 329.)
+ $5\text{H}_2\text{O}$. Very sol. in H_2O . (Thomsen, J. pr. (2) 13. 337.)

+ $6\text{H}_2\text{O}$. Sol. in ether and CHCl_3 without decomp. (Lengfeld, Am. Ch. J. 1901, 26. 329.)

Ammonium bromaurate, NH_4AuBr_4 .

Ppt. (Gutbier, Z. anorg. 1914, 85. 358.)

Barium bromaurate.

Not deliquescent. Sol. in H_2O . (v. Bonsdorff, Pogg. 17. 261.)

Cæsium bromaurate, CsAuBr_4 .

Sl. sol. in H_2O or alcohol. Insol. in ether. (Wells and Wheeler, Sill. Am. J. 144. 157.)
Ppt. (Gutbier, Z. anorg. 1914, 85. 360.)

Cerium bromaurate, $\text{CeAuBr}_3 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Jolin, Bull. Soc. (2) 21. 533.)

Didymium bromaurate, $\text{DiAuBr}_3 + 9\text{H}_2\text{O}$.

Very deliquescent. Sol. in H_2O . (Cleve.)

Lanthanum bromaurate, $\text{LaAuBr}_3 + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve.)

Magnesium bromaurate.

Deliquescent in moist air. (v. Bonsdorff.)

Manganese bromaurate.

Deliquescent. (v. Bonsdorff.)

Potassium bromaurate, KAuBr_4 .
Sl. sol. in H_2O . More sol. in cold alcohol than in H_2O . (v. Bonsdorff.)
+2 H_2O . Sol. in 5.12 pts. H_2O at 15°, 1.56 pts. at 40°, and 0.48 pt. at 67°. Decomp. by ether. Sl. sol. in $\text{KBr} + \text{Aq}$. (Schottländer, A. 217. 314.)
+5 H_2O . Efflorescent. (v. Bonsdorff.)

Rubidium bromaurate, RbAuBr_4 .
As caesium bromaurate.
Ppt. (Gutbier, Z. anorg. 1914, 85. 359.)

Samarium bromaurate, $\text{SmAuBr}_6 + 10\text{H}_2\text{O}$.
Very deliquescent. (Cleve, Bull. Soc. (2) 43. 165.)

Sodium bromaurate, NaAuBr_4 .
Slowly sol. in H_2O . (v. Bonsdorff.)

Zinc bromaurate, $\text{Zn}(\text{AuBr}_4)_2$.
Very deliquescent. (v. Bonsdorff.)

Bromauricyanhdric acid.
Not known in free state.

Barium bromauricyanide, $\text{Ba}[\text{Au}(\text{CN})_2\text{Br}_2]_2 + 10\text{H}_2\text{O}$.
Very sol. in hot or cold H_2O , also in alcohol. (Lindbom, Lund. Univ. Arsk. 12. No. 6.)

Cadmium bromauricyanide, $\text{Cd}[\text{Au}(\text{CN})_2\text{Br}_2]_2 + 6\text{H}_2\text{O}$.
Very sol. in hot or cold H_2O , but solution is unstable. (Lindbom.)

Calcium bromauricyanide, $\text{Ca}[\text{Au}(\text{CN})_2\text{Br}_2]_2 + 10\text{H}_2\text{C}$.
Extremely sol. in H_2O and alcohol. (Lindbom.)

Cobalt bromauricyanide, $\text{Co}[\text{Au}(\text{CN})_2\text{Br}_2]_2 + 9\text{H}_2\text{O}$.
Moderately sol. in H_2O . Less sol. than other bromauricyanides. (Lindbom.)

Potassium bromauricyanide, $\text{KAu}(\text{CN})_2\text{Br}_2 + 3\text{H}_2\text{O}$.
Sol. in H_2O and alcohol.

Sodium bromauricyanide, $\text{NaAu}(\text{CN})_2\text{Br}_2 + 2\text{H}_2\text{O}$.
Very sol. in H_2O or alcohol.

Strontium bromauricyanide, $\text{Sr}[\text{Au}(\text{CN})_2\text{Br}_2]_2 + x\text{H}_2\text{O}$.
Very sol. in H_2O or alcohol.

Zinc bromauricyanide, $\text{Zn}[\text{Au}(\text{CN})_2\text{Br}_2]_2 + 8\text{H}_2\text{O}$.
Easily sol. in cold or hot H_2O .

Bromhydric acid, HBr .
Very sol. in H_2O .
The most concentrated $\text{HBr} + \text{Aq}$ has a sp. gr. of 1.78, and contains 82.02% HBr . (Champion and Pellat, C. R. 70. 620.) This, or a weak acid on heating leaves a residue, which

distils unchanged at 125–125.5° under 785 mm. pressure, and contains 48.17% HBr (Topsoë); at 126° under 758 mm. pressure, and contains 46.83% HBr (Bineau); and has sp. gr. = 1.486 at 20° (Bineau); sp. gr. = 1.48 at 20° (Champion and Pellat); sp. gr. = 1.49 at 20° (Topsoë).

According to Roscoe (A. 116. 214) an acid of constant composition, obtained by boiling a stronger or a weaker acid, if distilled under 752–762 mm. pressure, contains 47.38–47.86% HBr , and boils at 126° at 760 mm. pressure; but the composition is dependent on the pressure, as, for example, under 1952 mm. pressure, the residue boils at 153°, and contains 46.3% HBr . (Roscoe.)

By conducting dry air through $\text{HBr} + \text{Aq}$ an acid is obtained containing 51.65% HBr if at 16°, and 49.35% HBr if at 100° (Roscoe).

1 vol. H_2O dissolves 600 ± vols. HBr at 10°. (Berthelot, C. R. 76. 679.)

1 pt. H_2O at t° and 760 mm. pressure dissolves pts. HBr .

t°	Pts. HBr	t°	Pts. HBr	t°	Pts. HBr
—25	2.550	—5	2.280	+50	1.715
—20	2.473	0	2.212	+75	1.505
—15	2.390	+10	2.103	+100	1.300
—10	2.335	+25	1.930

(Roozeboom, R. t. c. 4. 107.)

Absorption by 1 pt. H_2O at t° and p pressure in mm.

t° = —25°.

p	Pts. HBr	p	Pts. HBr
760	2.550	100	2.056
300	2.263	1	1.755
140	2.120	0.5	1.10

t° = —20°.

p	Pts. HBr	p	Pts. HBr
760	2.473	130	2.056
375	2.267	20	1.850
180	2.119

t° = —15°.

p	Pts. HBr	p	Pts. HBr
760	2.390	175	2.056
470	2.266	102	1.980
250	2.119

t° = —11.3°.

p	Pts. HBr	p	Pts. HBr
760	2.350	310	2.118
570	2.265	216	2.055

$t^{\circ} = -5^{\circ}$.

p	Pts. HBr	p	Pts. HBr
760	2.280	430	2.117
730	2.264	298	2.055

 $t^{\circ} = 0^{\circ}$.

p	Pts. HBr	p	Pts. HBr
760	2.212	380	2.054
540	2.116	5	1.085

(Roozeboom, R. t. c. 4. 107.)

Sp. gr. of HBr + Aq.

Sp. gr.	% HBr	Temp.	Sp. gr.	% HBr	Temp.
1.055	7.67	14°	1.335	36.67	13°
1.075	10.19	14°	1.349	37.86	13°
1.089	11.94	14°	1.368	39.13	13°
1.097	12.96	14°	1.419	43.12	13°
1.118	15.37	14°	1.431	43.99	13°
1.131	16.92	14°	1.438	44.62	13°
1.164	20.65	14°	1.451	45.45	14°
1.200	24.35	13°	1.460	46.09	13°
1.232	27.62	13°	1.485	47.87	14°
1.253	29.68	13°	1.490	48.17	14°
1.302	33.84	13°

(Topsoë, B. 3. 404.)

Sp. gr. of HBr + Aq at 14°.

% HBr	Sp. gr.	% HBr	Sp. gr.	% HBr	Sp. gr.
1	1.007	18	1.140	35	1.314
2	1.014	19	1.149	36	1.326
3	1.021	20	1.158	37	1.338
4	1.028	21	1.167	38	1.351
5	1.035	22	1.176	39	1.363
6	1.043	23	1.186	40	1.376
7	1.050	24	1.196	41	1.389
8	1.058	25	1.206	42	1.403
9	1.065	26	1.215	43	1.417
10	1.073	27	1.225	44	1.431
11	1.081	28	1.235	45	1.445
12	1.089	29	1.246	46	1.459
13	1.097	30	1.257	47	1.473
14	1.106	31	1.268	48	1.487
15	1.114	32	1.279	49	1.502
16	1.122	33	1.290
17	1.131	34	1.302

(Topsoë, calculated by Gerlach, Z. anal. 27. 316.)

Sp. gr. of HBr + Aq at 15°.

% HBr	Sp. gr.	% HBr	Sp. gr.	% HBr	Sp. gr.
5	1.038	25	1.204	45	1.435
10	1.077	30	1.252	50	1.515
15	1.177	35	1.305
20	1.159	40	1.365

Only a "moderate degree of accuracy" is claimed for this table. (Wright, C. N. 23. 242.)

Sp. gr. of HBr + Aq at 15°.

% HBr	Sp. gr.	% HBr	Sp. gr.	% HBr	Sp. gr.
1	1.0082	18	1.145	35	1.314
2	1.0155	19	1.154	36	1.326
3	1.0230	20	1.163	37	1.338
4	1.0305	21	1.172	38	1.350
5	1.038	22	1.181	39	1.362
6	1.046	23	1.190	40	1.375
7	1.053	24	1.200	41	1.388
8	1.061	25	1.209	42	1.401
9	1.069	26	1.219	43	1.415
10	1.077	27	1.229	44	1.429
11	1.085	28	1.239	45	1.444
12	1.093	29	1.249	46	1.459
13	1.102	30	1.260	47	1.474
14	1.110	31	1.270	48	1.490
15	1.119	32	1.281	49	1.496
16	1.127	33	1.292	50	1.513
17	1.136	34	1.303

(Biel, C. C. 1882. 148.)

Absorbed by alcohol with formation of C_2H_5Br .

The composition of the hydrates formed by HBr at different dilutions is calculated from determinations of the lowering of the fr. pt. produced by HBr and of the conductivity and sp. gr. of HBr + Aq. (Jones, Am. Ch. J. 1905, 34. 326.)

+ H_2O . (Roozeboom, R. t. c. 5. 363.)+ $2H_2O$. (Berthelot, A. ch. (5) 14. 369.)

(Pickering Chem. Soc. 1894, 64 (2) 232.

Mpt. -11.2° . (Pickering, l. c.)+ $3H_2O$. Mpt. -48.0° . (Pickering.)+ $4H_2O$. Mpt. -55.8° . (Pickering.)+ $5H_2O$. (Pickering.)**Bromhydric cyanhydric acid**, $3HBr$, $2HCN$.Decomp. by H_2O and alcohol.

Insol. in ether. (Gautier, A. ch. (4) 17. 141.)

Bromic acid, $HBrO_3$.

Known only in aqueous solution.

Solution evaporated on water bath decomposes when it contains 4.26% $HBrO_3$. In vacuo, an acid containing 50.59% $HBrO_3$ corresponding to formula $HBrO_3 + 7H_2O$ can be obtained.Not decomp. by dil. HNO_3 , or H_2SO_4 + Aq. Conc. H_2SO_4 decomposes.Alcohol and ether are quickly oxidized by $HBrO_3$.**Bromates**.Most of the bromates are very sol. in H_2O , a few are sl. sol., but none are insol., the least sol. being $AgBrO_3$ and $Hg_2(BrO_3)_2$.**Aluminum bromate**, $Al(BrO_3)_3$.

Deliquescent. (Rammelsberg, Pogg. 55. 63.)

+ $9H_2O$. Mpt. 62.3° . Less hygroscopic

than $\text{Al}(\text{ClO}_3)_3$. (Dobrosserdow, C. C. 1907. I, 1723.)

Ammonium bromate, NH_4BrO_3 .

Decomposes spontaneously; sol. in H_2O . (Rammelsberg, Pogg. 52. 85.)

Barium bromate, $\text{Ba}(\text{BrO}_3)_2$.

Solubility of $\text{Ba}(\text{BrO}_3)_2$ in H_2O . 100 g. sat.

$\text{Ba}(\text{BrO}_3)_2 + \text{Aq}$ at t° contain g. anhydrous $\text{Ba}(\text{BrO}_3)_2$.

t°	grams $\text{Ba}(\text{BrO}_3)_2$	t°	Grams $\text{Ba}(\text{BrO}_3)_2$
Eutectic point			
$-0.034^\circ \pm 0.002^\circ$	0.280	50°	1.72
0°	0.286	60°	2.271
$+10^\circ$	0.439	70°	2.922
20°	0.652	80°	3.521
25°	0.788	90°	4.26
30°	0.95	98.7°	5.256
40°	1.31	$*99.65^\circ$	5.39

* 99.65° is bpt. at 740 mm. = 100.39° at 760 mm.

(Anschütz, Z. phys. Ch. 1906, 56. 240.)

100 g. sat. $\text{Ba}(\text{BrO}_3)_2 + \text{Aq}$ contain 0.793 g. $\text{Ba}(\text{BrO}_3)_2$ at 25° . Sp. gr. of the solution at $25^\circ/4^\circ = 1.0038$. (Harkins J. Am. Chem. Soc. 1911, 33. 1815.)

Solubility of $\text{Ba}(\text{BrO}_3)_2$ in salts + Aq at 25° . C = concentration of salt in salt + Aq in milliequivalents per l.

d_1 = Sp. gr. at $25^\circ/4^\circ$ of salt + Aq.

S = solubility of $\text{Ba}(\text{BrO}_3)_2$ in salt + Aq expressed in milliequivalents per l.

d_2 = sp. gr. at $25^\circ/4^\circ$ of $\text{Ba}(\text{BrO}_3)_2 + \text{salt} + \text{Aq}$.

Salt	C	d_1	S	d_2
None			40.18	1.0038
KNO_3	25.018	0.9985	43.86	1.0059
	50.032	1.0030	47.03	1.0081
	99.970	1.0033	52.13	1.0120
$\text{Ba}(\text{NO}_3)_2$	25.018	1.0003	36.77	1.0059
	50.039	1.0025	34.74	1.0083
	99.97	1.0073	32.63	1.0132
	199.95	1.0183	30.95	1.0233
KBrO_3	24.988	1.0001	26.53	1.0046
	49.971	1.0031	17.37	1.0062
	99.85	1.0093	8.76	1.0109
$\text{Mg}(\text{NO}_3)_2$	100.0		52.57	1.0114

(Harkins, J. Am. Chem. Soc. 1911, 33. 1815.)

+ H_2O . Sol. in 130 pts. cold, and 24 pts. boiling H_2O . (Rammelsberg, Pogg. 52. 81.)

Decomp. by H_2SO_4 or $\text{HCl} + \text{Aq}$.

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Bismuth bromate.

Known only in solution, which decomp. on evaporation. (Rammelsberg, Pogg. 55. 76.)

Cadmium bromate, $\text{Cd}(\text{BrO}_3)_2 + \text{H}_2\text{O}$.

Sol. in 0.8 pt. cold H_2O . (Rammelsberg, Pogg. 55. 74.)

+ $2\text{H}_2\text{O}$. (Topsoë, J. B. 1872, 164.)

Cadmium bromate ammonia, $\text{Cd}(\text{BrO}_3)_2, 3\text{NH}_3$.

Decomp. by H_2O . (Rammelsberg, Pogg. 55. 74.)

$\text{Cd}(\text{BrO}_3)_2, 4\text{NH}_3$. Ppt. (Ephraim, B. 1915, 48. 51.)

Calcium bromate, $\text{Ca}(\text{BrO}_3)_2 + \text{H}_2\text{O}$.

Sol. in 1.1 pts. cold H_2O . (Rammelsberg, Pogg. 52. 98.)

Cerous bromate, $\text{Ce}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Easily sol. in H_2O . (Rammelsberg, Pogg. 55. 63.)

Mpt. 49° ; very sol. in H_2O with decomp. (James, J. Am. Chem. Soc. 1909, 31. 914.)

Cobaltous bromate, $\text{Co}(\text{BrO}_3)_2 + 6\text{H}_2\text{O}$.

Sol. in 2.2 pts. cold H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg, Pogg. 55. 71.)

Cupric bromate, basic, $6\text{CuO}, \text{Br}_2\text{O}_3 + 10\text{H}_2\text{O}$.

Ppt. (Rammelsberg, Pogg. 55. 78.)

Cupric bromate, $\text{Cu}(\text{BrO}_3)_2 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Rammelsberg, Pogg. 52. 92.)

Cupric bromate ammonia, $\text{Cu}(\text{BrO}_3)_2, 4\text{NH}_3$.

Completely sol. in a little H_2O , but decomp. by dilution.

Insol. in alcohol. (Rammelsberg, Pogg. 52. 92.)

Didymium bromate, $\text{Di}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac.)

Dysprosium bromate, $\text{Dy}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Mpt. 78° . Easily sol. in H_2O . Difficultly sol. in alcohol. (Jantsch, B. 1911, 44. 1275.)

Erbium bromate, $\text{Er}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Very sol. in alcohol and H_2O .

Glucinum bromate.

Deliquescent.

Iron (ferrous) bromate, $\text{Fe}(\text{BrO}_3)_2$.

Sol. in H_2O , but solution decomp. very easily.

Iron (ferric) bromate, $5\text{Fe}_2\text{O}_3, \text{Br}_2\text{O}_3 + 30\text{H}_2\text{O}$.

Partially sol. in H_2O , with separation of a more basic salt. Sol. in $\text{HBO}_3 + \text{Aq}$. (Rammelsberg, Pogg. 55. 68.)

Lanthanum bromate, $\text{La}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Sol. in $3\frac{1}{2}$ pts. H_2O at 15° . (Marignac, Ann. Min. (5) 15. 274.)

Mpt. 37.5° in its water of crystallization. 416 pts. are sol. in 100 pts. H_2O at 25° . (James, J. Am. Chem. Soc. 1909, 31. 913.)

ate, basic, 3PbO , $\text{Pb}(\text{BrO}_3)_2 +$
trömholm, Z. anorg. 1904, **38**. 441.)

ate, $\text{Pb}(\text{BrO}_3)_2$.

H_2O . $13.37 \times 10^{-1}\text{g.}$ are contained
of sat. solution at 20° . (Böttger,
1. 1903, **46**. 603.)

Sol. in 75 pts. cold H_2O . (Ram-
pogg. **52**. 96.)

omate, LiBrO_3 .

iquescent, and sol. in H_2O . (Ram-
pogg. A. **55**. 63.)

iquescent. (Politilitzin, B. **23**.

f solution sat. at $18^\circ = 1.833$, and
0.4% LiBrO_3 . (Mylius, B. 1897,

Not deliquescent. (Potilitzin.)

a bromate, $\text{Mg}(\text{BrO}_3)_2 + 6\text{H}_2\text{O}$.

ent. Sol. in 1.4 pts. cold H_2O at
in its water of crystallization when
ammelsberg, Pogg. **52**. 89.)

bromate, basic, $2\text{Hg}_2\text{O}$, Br_2O_3 .

warm H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$.
erg, Pogg. **55**. 79.)

bromate, $\text{Hg}_2(\text{BrO}_3)_2$.

by H_2O into basic salt. Difficultly
 $\text{O}_3 + \text{Aq}$; easily sol. in $\text{HCl} + \text{Aq}$.
erg.)

romate, basic, 2HgO , $\text{Br}_2\text{O}_3 + \text{H}_2\text{O}$.
ecomp. by cold, quickly by hot
tide and an acid salt.

l. in dil. acids. (Topsoë, W. A. B.

romate, $\text{HgBrO}_3 + 2\text{H}_2\text{O}$.

50 pts. cold, and 64 pts. boiling
ol. in $\text{HNO}_3 + \text{Aq}$. Easily sol. in
(Rammelsberg, Pogg. **55**. 79.)

romate ammonia.

decomp. in $\text{HCl} + \text{Aq}$. (Storer's

a bromate, $\text{Nd}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

7° . 146 pts. are sol. in 100 pts.
 $^\circ$. (James, J. Am. Chem. Soc.
5.)

ate, $\text{Ni}(\text{BrO}_3)_2 + 6\text{H}_2\text{O}$.

58 pts. cold H_2O . (Rammelsberg,
3.)

ate ammonia, $\text{Ni}(\text{BrO}_3)_2, 2\text{NH}_3$.

H_2O , with decomposition of the
on. Insol. in alcohol. (Rammels-

, 6NH_3 . Ppt. (Ephraim, B.
1.)

romate, KBrO_3 .

H_2O dissolve 6.58 pts. KBrO_3 at
elsberg). 100 pts. H_2O dissolve
 BrO_3 at 17.1° (Pohl. W. A. B. **6**.

595); at 0° , 3.11 pts.; at 20° , 6.92 pts.; at 40° ,
13.24 pts.; at 60° , 22.76 pts.; at 80° , 33.90 pts.;
at 100° 49.75 pts. KBrO_3 . Sat. solution boils
at 104° . (Kremers, Pogg. **97**. 5.)

1 l. H_2O at 25° dissolves 0.4715 moles
 KBrO_3 . (Geffcken, Z. phys. Ch. 1904, **49**.
296.)

1 l. H_2O dissolves 0.478 mol. KBrO_3 at 25° .
(Rothmund, Z. phys. Ch. 1909, **69**. 539.)

Sp. gr. of $\text{KBrO}_3 + \text{Aq}$ at 19.5° .

% KBrO_3	1	2	3	4	5
Sp. gr. .	1.009	1.016	1.024	1.031	1.039

% KBrO_3	6	7	8	9	10
Sp. gr. .	1.046	1.054	1.062	1.070	1.079

(Gerlach, Z. anal. **8**. 290.)

Solubility of KBrO_3 in salts + Aq at 25° .

Salt	Moles of KBrO_3 sol. in 1 liter of				
	.5-N solution	N solution	2-N solution	3-N solution	4-N solution
NaNO_3	0.5745	0.6497	0.7680	0.9026	1.031
NaCl	0.5220	0.5616	0.6042	0.6244	0.640

(Geffcken, Z. phys. Ch. 1904, **49**. 296.)

Easily sol. in liquid HF. (Franklin, Z.
anorg. 1905, **46**. 2.)

Sl. sol. in alcohol. (Rammelsberg.)

Insol. in absolute alcohol.

Solubility in organic compds. + Aq at 25° .

Solvent	Mol. KBrO_3 sol. in 1 litre
Water	0.478
0.5-N Methyl alcohol	0.444
" Ethyl alcohol	0.421
" Propyl alcohol	0.409
" Tert. amyl alcohol	0.383
" Acetone	0.425
" Ether	0.395
" Formaldehyde	0.397
" Glycol	0.448
" Glycerine	0.451
" Mannitol	0.451
" Glucose	0.463
" Sucrose	0.431
" Urea	0.477
" Dimethyl pyrone	0.478
" Ammonia	0.445
" Diethylamine	0.384
" Pyridine	0.415
" Piperidine	0.396
" Urethane	0.433
" Formamide	0.473
" Acetamide	0.445
" Glycocoll	0.501
" Acetic acid	0.456
" Phenol	0.426
" Methylal	0.405
" Methyl acetate	0.420

(Rothmund, Z. phys. Ch. 1909, **69**. 539.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Praseodymium bromate, $\text{Pr}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Mpt. 56.5° . 190 pts. are sol. in 100 pts. H_2O at 25° . (James, J. Am. Chem. Soc. 1909, 31. 914.)

Samarium bromate, $\text{Sm}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Mpt. 75° . 114 pts. are sol. in 100 pts. H_2O at 25° . Very sl. sol. in alcohol. (James, J. Am. Chem. Soc. 1909, 31. 915.)

Scandium bromate.

(Crookes, Roy. Soc. Proc. 1908, 80, A, 518.)

Silver bromate, AgBrO_3 .

1 pt. H_2O dissolves 0.00810 pt. AgBrO_3 at 24.5° . (Noyes, Z. phys. Ch. 6. 246.)

Sol. in 595.3 pts. H_2O at 25° .

Sol. in 320.4 pts. $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.21) at 25° .

Sol. in 2.2 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (sp. gr. 0.96) at 25° . (Longi, Gazz. ch. it. 13. 87.)

1 l. H_2O dissolves 1.71 g. AgBrO_3 at 27° . (Whitby, Z. anorg. 1910, 67. 108.)

Sl. sol. in H_2O . 1.59×10^{-4} g. are contained in 1 liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

Insol. in HNO_3 . (Löwig.) Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Silver bromate ammonia, $\text{AgBrO}_3, 2\text{NH}_3$.

Decomp. in air or by H_2O . (Rammelsberg, Pogg. 52. 94.)

Sodium bromate, NaBrO_3 .

Sol. in 2.7 pts. H_2O at 15° . (Rammelsberg.)

100 pts. H_2O dissolve at—

0° 20° 40° 60° 80° 100°

27.54 34.48 50.25 62.5 75.75 90.9 pts. NaBrO_3 .

(Kremers, Pogg. 94. 271.)

Easily forms supersaturated solutions.

Sat. solution boils at 109° . (Kremers.)

$\text{NaBrO}_3 + \text{Aq}$ containing 10.10% NaBrO_3 has sp. gr. $20^\circ/20^\circ = 1.0818$.

$\text{NaBrO}_3 + \text{Aq}$ containing 11.09% NaBrO_3 has sp. gr. $20^\circ/20^\circ = 1.0900$.

(Le Blanc and Rohland, Z. phys. Ch. 1896 19. 278.)

Sp. gr. of $\text{NaBrO}_3 + \text{Aq}$ at 19.5° .

% NaBrO_3	5	10	15
Sp. gr.	1.041	1.083	1.129

% NaBrO_3	20	25	30
Sp. gr.	1.178	1.231	1.289

(Kremers, Pogg. 97. 5, calculated by Gerlach, Z. anal. 8. 290.)

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Sodium bromate bromide, $3\text{NaBrO}_3, 2\text{NaBr} + 3\text{H}_2\text{O}$.

Decomp. by H_2O or alcohol. (Fritzsche.)

Strontium bromate, $\text{Sr}(\text{BrO}_3)_2 + \text{H}_2\text{O}$.

Sol. in 3 pts. H_2O (Rammelsberg, Pogg. 52. 84); less sol. in H_2O than $\text{SrBr}_2 + 6\text{H}_2\text{O}$. (Löwig.)

Thallous bromate, TlBrO_3 .

Sl. sol. in hot H_2O ; easily sol. in $\text{HNO}_3 + \text{Aq}$. (Oettinger.)

Easily sol. in H_2O and dil. acids. (Ditte, A. ch. (6) 21. 145.)

Terbium bromate, $\text{Tb}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

Not deliquescent. (Potratz, C. N. 1905, 92, 3.)

Thallous bromate, TlBrO_3 .

1 l. H_2O at 39.75° dissolves 2.216×10^{-2} g. mol. (Noyes and Abbott, Z. phys. Ch. 1895, 16. 130.)

Sl. sol. in H_2O . 3.46×10^{-1} gram are contained in 1 liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

Thallic bromate, $\text{Tl}(\text{BrO}_3)_3 + 3\text{H}_2\text{O}$.

Very hygroscopic. Easily decomp. by H_2O . (Gewecke, Z. anorg. 1912, 75. 275.)

Thulium bromate, $\text{Tm}_2(\text{BrO}_3)_6 + 18\text{H}_2\text{O}$.

Pptd. from sat. aqueous solution by 95% alcohol.

NH_4OH is the best precipitant. (James, J. Am. Chem. Soc. 1911, 33. 1342.)

Tin (stannous) bromate (?).

Insol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$.

Uranyl bromate, $4\text{UO}_3, 3\text{Br}_2\text{O}_3 + 16\text{H}_2\text{O}$.

Sol. in H_2O . (Rammelsberg.)

Yttrium bromate, $\text{Y}(\text{BrO}_3)_3 + 9\text{H}_2\text{O}$.

More easily sol. in H_2O than $\text{Y}(\text{IO}_3)_3$. Sl. sol. in alcohol. Insol. in ether. (Clève.)

Mpt. 74° . 168 pts. are sol. in 100 pts. H_2O at 25° .

Sl. sol. in alcohol. (James, J. Am. Chem. Soc. 1909, 31. 916.)

Zinc bromate, $\text{Zn}(\text{BrO}_3)_2 + 6\text{H}_2\text{O}$.

Sol. in 1 pt. cold H_2O . (Rammelsberg, Pogg. 52. 90.)

Zinc bromate ammonia, $\text{Zn}(\text{BrO}_3)_2, 2\text{NH}_3 + 3\text{H}_2\text{O}$.

Decomp. by H_2O and alcohol. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg, Pogg. 52. 90.)

$\text{Zn}(\text{BrO}_3)_2, 4\text{NH}_3$. Ppt. (Ephraim, B. 1915, 48. 51.)

Perbromic acid.

See Perbromic acid.

les.

bromides are sol. in H_2O , many in and some in ether.
and Hg_2Br_2 are insol. in H_2O or acids; and $TlBr$ are sl. sol. therein. Cu_2Br_2 in H_2O , sol. in acids.
nder each element.

ie, Br_2 .

Br dissolves at 15° in 33 pts. H_2O . (Pogg. 14. 485.)
 Br dissolves at 15° in 31 pts. H_2O . (r, Chem. Soc. 15. 477.)

ibility of Br in 100 pts. H_2O at t° .

Pts. Br.	t°	Pts. Br.	t°	Pts. Br.
3.600	15	3.226	25	3.167
3.327	20	3.208	30	3.126

(Dancer, l. c.)

t. aqueous solution of Br contains Br at 0° ; 3.80% Br at 3° ; 3.33% Br at Roozeboom, R. t. c. 3. 29, 59, 73, 84.)
 H_2O dissolves 34 g. Br at 25° . (Jakow-phys. Ch. 1896, 20. 25.)
is sol. in 30 pts. H_2O . (Dietze, Chem. 99, 76 (2) 150.)

pts. H_2O dissolve at:
 10.34° 19.96° 30.17° 40.03° 49.85°
3.740 3.578 3.437 3.446 3.522
mine.

and bromine as such is insol. in H_2O ; the vapor dissolves. (Winkler, Ch. Z. 3. 688.)
 H_2O dissolves 33.95 g. Br_2 at 25° . (Mc-an, Z. phys. Ch. 1903, 44. 617.)

ility of bromine vapor in H_2O at t° .
 a = coefficient of absorption.

a	t°	a
60.5	42	8.6
54.1	44	7.9
48.3	46	7.4
43.3	48	6.9
38.9	50	6.5
35.1	52	6.1
31.5	54	5.8
28.4	56	5.4
25.7	58	5.1
23.4	60	4.9
21.3	62	4.6
19.4	64	4.4
17.7	66	4.2
16.3	68	4.0
15.0	70	3.8
13.8	72	3.6
12.7	74	3.4
11.7	76	3.3
10.9	78	3.1
10.1	80	3.0
9.4

(Winkler, Ch. Z. 1899, 23. 688.)

Solubility of bromine vapor.
(Mean of many determinations)

Temp.	Pressure	Absorption coefficient
0.0	56-13mm.	60.53
9.94°	80-16	35.22
20.46	138-9	20.87
30.38	179-12	13.65
40.31	229-26	9.22
50.25	274-53	6.50
60.04	314-46	4.84
69.98	154-54	3.82
80.22	396-74	2.94

Solubility of liquid bromine.
(The mean of many determinations)

Temp.	0°	10.34°	19.96°	30.17°	40.03°	49.85°
Pts. H_2O that dissolve 1 pt. Br_2	24.0	26.74	27.94	29.10	29.02	28.38

Much less Br_2 is sol. in ice cold H_2O in the presence of bromine hydrate.

Solubility in presence of bromine hydrate.
(The mean of many determinations)

Temp.	0°	5.12°
Pts. H_2O that dissolve 1 pt. Br_2	42.39	26.26

(Winkler, Ch. Z. 1899, 23. 688-689.)

Solubility of Br_2 in H_2O at $25^\circ = 0.21$ mols. in 1 l. (Bray, J. Am. Chem. Soc. 1910, 32. 398.)

Sp. gr. of $Br_2 + Aq$ containing pts. Br in 1000 pts. solution.

Pts. Br.	Sp. gr.	Pts. Br.	Sp. gr.
10.72	1.00901	18.74-19.06	1.01491
10.68	1.00931	19.52-20.09	1.01585
12.05	1.00995	20.89-21.55	1.01807
12.21	1.01223	31.02-31.69	1.02367

(Slessor, N. Edin. Phil. J. 7. 287.)

Sp. gr. of $Br_2 + Aq$ at 32.5° .

% Br_2 by weight	Sp. gr.
0.7214	0.999814
1.1172	1.002520
1.6448	1.006100
1.9956	1.008870
2.5960	1.013200

(Joseph, Chem. Soc. 1915, 107. 3.)

Sol. in conc. HCl , HBr , conc. solutions of bromides, and in liquid SO_2 . (Sestini, Zeit. Chem. 1868. 718.)

Much more sol. in $HCl + Aq$ than in H_2O . 100 ccm. $HCl + Aq$ of 1.153 sp. gr. dissolve 36.4 g. Br at 12° .

More sol. in $SrCl_2$, and $BaCl_2 + Aq$ than in H_2O . (Berthelot, C. R. 100. 761.)

Bromine is not more sol. in KBr + Aq than in H₂O (?). (Balard.)

KBr + Aq containing 1 pt. KBr to 6 pts. H₂O takes up as much Br as it already contains; when this solution is heated the dissolved Br is separated. 1 pt. KBr + 1 pt. H₂O takes up twice as much Br as it already contains, much heat being evolved. This solution loses Br on exposure to the air or when heated. (Löwig.)

Solubility of Br₂ in KBr + Aq.

g. Mols. KBr per l.	g. at. Br dissolved per l at 18.5°	g. at. Br dissolved per l. at 26.5°.
0.00	0.4448	0.4282
0.01	0.4634	0.4490
0.02	0.4823	0.4671
0.03	0.5049	0.4925
0.04	0.5243	0.5101
0.05	0.5431	0.5301
0.06	0.5668	0.5530
0.07	0.5895	0.5636
0.08	0.6059	0.5920
0.09	0.6301	0.5981
0.1	0.6533	0.6488
0.2	0.8718	0.8591
0.3	1.0549	1.0787
0.4	1.3124	1.2704
0.5	1.5436	1.4731
0.6	1.7712	1.6717
0.7	2.0006	1.9197
0.8	2.2354	2.1029
0.9	2.4851	2.3349

The above figures indicate that below a concentration of 0.1 g. mol. KBr per l. just enough Br is dissolved to form KBr₃, while above that concentration somewhat larger amounts of Br are dissolved, which is greater at the lower temp.

(Worley, Chem. Soc. 1905, 87. 1109.)

Solubility of Br₂ in NaBr + Aq at 25°.

g. NaBr per l.	g.-atoms Br ₂ per l.	Sp. gr.
92.6	2.479	1.213
160.5	4.345	1.372
205.8	6.195	1.515
255.8	8.575	1.678
319.7	13.65	1.997
359.0	16.04	2.137
...	19.23	2.327
408.3	20.85	2.420

(Bell, J. Am. Chem. Soc. 1912, 34. 14.)

Solubility in salts + Aq.

Solubility in 1 liter K₂SO₄ + Aq at 25°.

K ₂ SO ₄ + Aq	g. Bromine
1-N	25.14
1/2-N	29.44
1/4-N	31.46
1/8-N	32.70
1/16-N	33.10

(Jakowkin, Z. phys. Ch. 1896, 20, 26.)

Solubility in 1 liter Na₂SO₄ + Aq at 25°.

Na ₂ SO ₄ + Aq	g. Bromine
1-N	25.07
1/2-N	29.20
1/4-N	31.33
1/8-N	32.94
1/16-N	33.26

(Jakowkin, l. c.)

Solubility in 1 liter NaNO₃ + Aq at 25°.

NaNO ₃ + Aq	g. Bromine
1-N	28.80
1/2-N	31.35
1/4-N	32.62
1/8-N	33.33
1/16-N	33.74

(Jakowkin, l. c.)

Solubility in salts + Aq at 25°.

Salt + Aq.	g. Br ₂ sol. in 1 liter
1/2-N. Na ₂ SO ₄	23.90
1/2-N. K ₂ SO ₄	24.80
1/2-N. (NH ₄) ₂ SO ₄	77.7
N. NaNO ₃	28.00
N. KNO ₃	28.95
N. NH ₄ NO ₃	55.15
N. NaCl	55.90
N. KCl	57.40
N. NH ₄ Cl	82.2

(McLauchlan, Z. phys. Ch. 1903, 44. 617.)

Solubility in HgBr₂ + Aq at 25°.

10 ccm. of the solution contain:—

Millimols. Br ₂	Millimols. Hg
2.125	0.
2.204	0.0560
2.216	0.0793
2.226	0.1284
2.231	0.2120

(Herz and Paul, Z. anorg. 1914, 85. 215.)

1 l. N.NH₄C₂H₃O₂ + Aq dissolves 340.5 g. Br₂ at 25°. (McLauchlan, Z. phys. Ch. 1903, 44. 617.)

Miscible in all proportions with liquid NO₂. (Frankland, Chem. Soc. 1901, 79. 1361.)

More sol. in alcohol than in H₂O; miscible with ether, CS₂, CHCl₃. (Sestini, Zeit. Chem. 1868. 718.)

Somewhat soluble in glycerine. (Pelouze.) Sol. in benzene (Mansfield); insol. in benzene (Moride, A. ch. (3) 39. 452). Sol. in warm chloral, bromal, and iodal. (Löwig, Pogg. 14. 485.) Sol. in SCl₂ (Solly), and SBr₂. Sol. in conc. HC₂H₃O₂ + Aq. (Balard.) Sol. in aqueous solution of potassium, sodium, or calcium acetates. (Cahours.)

Solubility in CS_2 .

100 g. of the sat. solution contain at:

-95°	-110.5°	-116°
45.4	39.0	36.9 g. Br_2 .

(Arctowski, Z. anorg. 1896, 11. 274.)

Cryst. from CS_2 at -90° in fine needles.
(Arctowski, Z. anorg. 1895, 10. 25.)

Sp. gr. of $\text{Br}_2 + \text{CCl}_4$ at 32.5° .

% Br_2 by weight	Sp. gr.
1.5449	1.58014
1.6454	1.58060
1.7990	1.58168
2.6676	1.58312
3.5833	1.59526

(Joseph, Chem. Soc. 1915, 107. 3.)

Sp. gr. of $\text{Br}_2 + \text{nitrobenzene}$ at 32.5° .

% Br_2 by weight	Sp. gr.
1.5643	1.20225
3.2323	1.21449
4.6462	1.22518
6.1826	1.23603

(Joseph, Chem. Soc. 1915, 107. 3.)

Very sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in acetone. (Eidmann, C. C. 1899, II, 1014; Naumann, B. 1904, 37. 4328.)

Partition of Br_2 between water and other solvents.

W = millimols Bromine in 10 cm. of the aqueous layer.

G = millimols Bromine in 10 cm. of the other layer.

Other solvent	G	W	G/W
CCl_4	1.949	0.0853	22.73
	7.008	0.3085	22.71
	12.171	0.5300	23.13
	39.880	1.3132	30.32
	54.574	1.5560	35.01
75% by vol. CCl_4 + 25% by vol. CS_2	3.567	0.0985	37.06
	7.304	0.1910	38.15
	10.833	0.2900	37.36
	13.922	0.3720	37.42
	17.230	0.4580	37.62
	25.637	0.6580	38.96
	40.625	0.9940	40.88
50% by vol. CCl_4 + 50% by vol. CS_2	54.035	1.2080	44.73
	3.592	0.0784	45.82
	6.820	0.1487	46.85
	10.148	0.2206	46.01
	13.866	0.3065	45.24
	16.616	0.3688	45.05
	42.975	0.8086	53.15
25% by vol. CCl_4 + 75% by vol. CS_2	55.965	0.9960	56.19
	5.753	0.0884	65.05
	10.902	0.1682	64.82
	26.724	0.4970	65.65
	41.314	0.6331	65.26
	55.526	0.8520	65.17

Partition of Br_2 , etc.—Continued

Other solvent	G	W	G/W
CS_2	7.750	0.1015	76.35
	10.600	0.1387	76.44
	14.696	0.1910	76.98
	17.999	0.2352	76.54
	26.345	0.3467	75.99
	40.625	0.5194	78.21
	57.038	0.7160	79.66

(Herz, Z. Elektrochem, 1910, 16. 871.)

Partition coefficient for bromine between CS_2 and H_2O at 25°C .

A = concentration of the water layer.

C = concentration of the CS_2 layer.

A	C	N = C/A
7.545	691.9	91.71
4.109	338.6	82.41
2.660	217.4	81.72
2.544	207.7	81.66
1.740	140.38	80.67
1.2878	103.7	80.51
0.8073	64.44	79.83
0.5046	39.64	78.38

Partition coefficient for bromine between CHBr_3 and H_2O at 25°C .

A = concentration of the water layer.

C = concentration of the CHBr_3 layer

A	C	N = C/A
5.424	373.6	68.88
3.838	264.7	68.80
2.368	161.5	68.19
1.348	90.17	66.90
0.766	50.49	65.84
0.366	23.62	64.85

Partition coefficient for bromine between CCl_4 and H_2O at 25°C .

A = concentration of the water layer.

C = concentration of the CCl_4 layer.

A	C	N = C/A
14.42	545.2	37.82
10.80	372.2	34.44
7.901	252.8	32.01
7.163	225.8	31.52
6.803	218.5	32.12
5.651	172.6	30.54
3.216	94.84	29.48
2.054	58.36	28.41
1.266	35.92	28.37
0.7711	21.53	27.92
0.5761	15.72	27.26
0.4476	12.09	27.02
0.3803	10.27	27.00
0.2478	6.691	27.00

(Jakowkin, Z. phys. Ch. 1895, 13. 588.)

Partition of bromine between CCl_4 and salts + Aq.

A = concentration of Br in H_2O layer.

C = concentration of Br in CCl_4 layer.

Partition of Br_2 between CCl_4 and NaNO_3 + Aq at 25° .

NaNO_3 + Aq	A	C
1-N	7.905	316.7
$\frac{1}{2}$ -N	8.763	319.5
$\frac{1}{4}$ -N	9.033	315.7
$\frac{1}{8}$ -N	9.200	316.7
$\frac{1}{16}$ -N	9.399	319.3

(Jakowkin, Z. phys. Ch. 1896, 20, 25.)

Partition of Br_2 between CCl_4 and K_2SO_4 + Aq at 25° .

K_2SO_4 + Aq	A	C
1-N	5.982	255.4
$\frac{1}{2}$ -N	6.843	253.4
$\frac{1}{4}$ -N	7.354	252.8
$\frac{1}{8}$ -N	7.585	250.3
$\frac{1}{16}$ -N	7.498	242.3

(Jakowkin, l. c.)

Partition of Br_2 between CCl_4 and Na_2SO_4 + Aq at 25° .

Na_2SO_4 + Aq	A	C
1-N	5.934	254.6
$\frac{1}{2}$ -N	6.838	253.4
$\frac{1}{4}$ -N	7.402	254.4
$\frac{1}{8}$ -N	7.609	252.8
$\frac{1}{16}$ -N	7.713	251.2

(Jakowkin, l. c.)

Crystallizes at 4° with $10\text{H}_2\text{O}$.

Bromine chloride, BrCl .

Sol. in H_2O , CS_2 , ether, etc.

Bromine fluoride, BrF_3 .

Fumes in the air. Decomp. by H_2O . (Lebeau, C. R. 1905, 141, 1019.)

Bromine oxides.

No oxides of bromine are known in the free state. See hypobromous, bromic, and perbromic acids.

Bromiridic acid.

Ammonium bromiridate, $(\text{NH}_4)_2\text{IrBr}_6$.

Less sol. in cold H_2O than the K salt. (Birnbbaum, Zeit. Chem. 1865, 22.)

Very sol. in cold H_2O . (Gutbier, B. 1909, 42, 3910.)

Cæsium bromiridate, Cs_2IrBr_6 .

Sol. in H_2O . (Gutbier, B. 1909, 42, 3911.)

Potassium bromiridate, K_2IrBr_6 .

Moderately sol. in cold, more easily in hot H_2O .

Insol. in alcohol or ether.

Sol. in cold H_2O and in HBr + Aq. (Gutbier, B. 1909, 42, 3910.)

Rubidium bromiridate, Rb_2IrBr_6 .

Very sol. in cold H_2O . Sol. in hot dil. HBr + Aq. (Gutbier, B. 1909, 42, 3911.)

Sodium bromiridate.

Deliquescent. Easily sol. in H_2O , alcohol, or ether.

Bromiridous acid, $\text{H}_2\text{Ir}_2\text{Br}_{12} + 6\text{H}_2\text{O}$.

Easily sol. in H_2O , alcohol, or ether. (Birnbbaum, 1864.)

Ammonium bromiridite, $(\text{NH}_4)_2\text{Ir}_2\text{Br}_{12} + \text{H}_2\text{O}$.

Difficultly sol. in H_2O . (Birnbbaum.)

Potassium bromiridite, $\text{K}_2\text{Ir}_2\text{Br}_{12} + 6\text{H}_2\text{O}$.

Efflorescent. Sol. in H_2O .

Silver bromiridite, $\text{Ag}_2\text{Ir}_2\text{Br}_{12}$.

Ppt. Insol. in H_2O or acids.

Sodium bromiridite, $\text{Na}_2\text{Ir}_2\text{Br}_{12} + 24\text{H}_2\text{O}$.

Efflorescent. Very sol. in H_2O .

Bromocarbonatoplatindiamine carbonate, $\text{CO}_3[\text{Pt}(\text{N}_2\text{H}_4)_2]_2(\text{CO}_3)_2 + 4\text{H}_2\text{O}$.

Ppt.

Bromocarbonatoplatindiamine carbonate

bromoplatin/diamine nitrate,

$\text{CO}_3[\text{Pt}(\text{N}_2\text{H}_4)_2]_2(\text{CO}_3)_2$, $2\text{Br}_2\text{Pt}(\text{N}_2\text{H}_4)_2(\text{NO}_3)_2$.

Bromochloroplatindiamine chloride,

$\text{Br Pt}(\text{N}_2\text{H}_4)_2\text{Cl}_2$.

Very sl. sol. in H_2O . (Cleve.)

— chlorobromide, $\text{Br Pt N}_2\text{H}_4\text{Cl N}_2\text{H}_4\text{Br}^{(?)}$.

Very sl. sol. in H_2O .

Bromochlororoplatinic acid.

Potassium bromochloroplatinate, $\text{K}_2\text{PtCl}_2\text{Br}_2$ (Pitkin, J. Am. Chem. Soc. 2, 408.)

Mixture. (Herty, J. Am. Chem. Soc. 1896, 18, 136.)

$\text{K}_2\text{PtCl}_4\text{Br}_2$. Sl. sol. in cold H_2O ; much more sol. in hot H_2O . (Pitkin.)

Mixture. (Herty.)

$\text{K}_2\text{PtCl}_2\text{Br}_2$. As above.

$\text{K}_2\text{PtCl}_2\text{Br}_4$. (Pigeon, A. ch. 1894, (7) 2, 488.)

$\text{K}_2\text{PtClBr}_3$. (Pitkin.)

Bromochromic acid.

Potassium bromochromate, $\text{KCrO}_2\text{Br}_2 = \text{CrO}_2(\text{Br})\text{OK}$.

Decomp. by H_2O . (Heintze, J. pr. (2) 4, 225.)

- Dibromochromium chloride**,
 $[\text{Cr}(\text{H}_2\text{O})_4\text{Br}_2]\text{Cl} + 2\text{H}_2\text{O}$.
 Ppt. Nearly insol. in fuming HCl . (Bjerrum, B. 1907, 40. 2918.)
- Bromohydroxyloplatindiamine bromide**,
 $\text{OH} \text{Pt}(\text{N}_2\text{H}_4\text{Br})_2$.
 Very sl. sol. in H_2O . (Cleve.)
- **chloride**, $\text{OH} \text{Pt}(\text{N}_2\text{H}_4\text{Cl})_2$.
 Sol. in H_2O (Cleve.)
- **nitrate**, $\text{OH} \text{Pt}(\text{N}_2\text{H}_4\text{NO}_3)_2$.
 Very sl. sol. in cold, moderately sol. in hot H_2O . (Cleve.)
- Bromohydroxyloplatinmonodiamine nitrate**, $\text{Br} \text{Pt}(\text{NH}_3)_2\text{NO}_3 + \text{H}_2\text{O}$.
 Easily sol. in H_2O . (Cleve.)
- Bromomercurosulphurous acid**.
- Ammonium bromomercurosulphite**,
 $\text{NH}_4\text{SO}_3\text{HgBr}$.
 Sol. in H_2O . (Barth, Z. phys. Ch. 9. 215.)
- Potassium bromomercurosulphite**,
 KSO_3HgBr .
 As above. (B.)
- Bromomolybdenum bromide**,
 $\text{Br}_4\text{Mo}_3\text{Br}_2 = \text{molybdenum dibromide}$,
 MoBr_2 .
 Insol. in H_2O or acids, or even in boiling aqua regia. Easily sol. in dilute, decomp. by conc. alkalis + Aq. (Blomstrand, J. pr. 82. 436.)
- Bromomolybdenum chloride**, $\text{Br}_4\text{Mo}_3\text{Cl}_2 + 3\text{H}_2\text{O}$.
 Insol. in acids. (Blomstrand.)
- Bromomolybdenum chromate**, $\text{Br}_4\text{Mo}_3\text{CrO}_4 + 2\text{H}_2\text{O}$.
 Insol. in dil. acids. Sol. in hot conc. HCl + Aq. Insol. in alkali chromates + Aq. (Atterberg.)
- Bromomolybdenum fluoride**, $\text{Br}_4\text{Mo}_3\text{F}_2 + 3\text{H}_2\text{O}$.
 Insol. in H_2O . (Atterberg.)
- Bromomolybdenum hydroxide**, $\text{Br}_4\text{Mo}_3(\text{OH})_2$.
 Completely sol. in alkalis if not heated over 90° . (Atterberg.)
 $+ 2\text{H}_2\text{O}$.
 $+ 8\text{H}_2\text{O}$.
- Bromomolybdenum iodide hydroxide**,
 $2\text{Br}_4\text{Mo}_3\text{I}_2, \text{Br}_4\text{Mo}_3(\text{OH})_2 + 8\text{H}_2\text{O}$.
 Precipitate. (Blomstrand, J. pr. 77. 92.)
- Bromomolybdenum molybdate**, $\text{Br}_4\text{Mo}_3\text{MoO}_4$.
 Precipitate. (Atterberg.)
- Bromomolybdenum phosphate**,
 $\text{Br}_4\text{Mo}_3\text{H}_4(\text{PO}_4)_2$.
 Precipitate. Insol. in H_2O . (Atterberg.)
- Bromomolybdenum sulphate**, $\text{Br}_4\text{Mo}_3\text{SO}_4 + 3\text{H}_2\text{O}$.
 Precipitate. Sl. sol. in boiling H_2SO_4 . (Atterberg.)
- Dibromomolybdous acid**, $\text{MoOBr}_2(\text{OH}) + 1\frac{1}{2}\text{H}_2\text{O}$.
 Sol. in H_2O . Very hygroscopic. (Weinland, Z. anorg. 1905, 44. 86.)
- Tetrabromomolybdous acid**, $\text{MoBr}_4(\text{OH}) + 2\text{H}_2\text{O}$.
 Sol. in H_2O . Hygroscopic. (Weinland, l. c.)
- Diammonium pentabromomolybdite**,
 $\text{MoBr}_5\text{O}(\text{NH}_4)_2$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Dicæsium pentabromomolybdite**,
 $\text{MoBr}_5\text{OCs}_2$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Calcium tetrabromomolybdite**, $(\text{MoBr}_4\text{O})_2\text{Ca} + 7\text{H}_2\text{O}$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Monolithium tetrabromomolybdite**,
 $\text{MoBr}_4(\text{OLi}) + 4\text{H}_2\text{O}$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Magnesium pentabromomolybdite**,
 $\text{MoBr}_5(\text{OMg}) + 7\text{H}_2\text{O}$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Monopotassium tetrabromomolybdite**,
 $\text{MoBr}_4(\text{OK}) + 2\text{H}_2\text{O}$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Dipotassium pentabromomolybdite**,
 MoBr_5OK_2 .
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Dirubidium pentabromomolybdite**,
 $\text{MoBr}_5\text{ORb}_2$.
 Hygroscopic. Sol. in H_2O . (Weinland, l. c.)
- Bromonitratoplatindiamine nitrate**,
 $\text{Br} \text{Pt} \text{N}_2\text{H}_4\text{NO}_3$.
 $\text{NO}_3 \text{Pt} \text{N}_2\text{H}_4\text{NO}_3$.
 Decomp. by H_2O . (Cleve.)
- **sulphate**, $\text{Br} \text{Pt}(\text{N}_2\text{H}_4)_2\text{SO}_4 + \text{H}_2\text{O}$.
 Sl. sol. in H_2O .

Bromonitritoplatinsemidiamine nitrite,
 $\text{NO}_2\text{Br}_2\text{Pt}(\text{NH}_3)_2\text{NO}_2$.

Sl. sol. in H_2O . (Blomstrand.)

Bromonitrous acid.

Platinum silver bromonitrite, $\text{PtAg}_2\text{Br}_2(\text{NO}_2)_4$.
 Ppt. (Miolati, Gazz. ch. it. 1900, 30. 588.)

Bromopalladic acid.

Ammonium bromopalladate, $(\text{NH}_4)_2\text{PdBr}_6$.

Difficultly sol. in cold H_2O . Decomp. by hot H_2O and by hot conc. H_2SO_4 . (Gutbier, B. 1905, 38. 1907.)

Cæsium bromopalladate, Cs_2PdBr_6 .

Difficultly sol. in cold H_2O . Decomp. by hot H_2O or by hot conc. H_2SO_4 . (Gutbier, l. c.)

Potassium bromopalladate, K_2PdBr_6 .

Difficultly sol. in cold H_2O . Decomp. by hot H_2O or by hot conc. H_2SO_4 . (Gutbier, l. c.)

Rubidium bromopalladate, Rb_2PdBr_6 .

Insol. in cold H_2O . Decomp. by hot H_2O or by hot conc. H_2SO_4 . (Gutbier, l. c.)

Bromopalladious acid.

Ammonium bromopalladite, $(\text{NH}_4)_2\text{PdBr}_4$.

Very stable. Sol. in H_2O . (Smith, Z. anorg. 1894, 6. 381.)

Very sol. in cold H_2O .

Can be cryst. from a very small amount of hot H_2O . (Gutbier, B. 1905, 38. 2387.)

Barium bromopalladite.

Not deliquescent. Sol. in H_2O . (v. Bonsdorff.)

Cæsium bromopalladite, Cs_2PdBr_4 .

Very sol. in H_2O . (Gutbier, B. 1905, 38. 2388.)

Manganese bromopalladite, MnPdBr_4 .

Sol. in H_2O and alcohol. (v. Bonsdorff.)
 $+7\text{H}_2\text{O}$. Very sol. in H_2O . (Smith, Z. anorg. 1894, 6. 382.)

Potassium bromopalladite, K_2PdBr_4 .

Easily sol. in H_2O . (Joannis, C. R. 95. 295.)

Very stable. Sol. in H_2O . (Smith, Z. anorg. 1894, 6. 381.)

$+2\text{H}_2\text{O}$. Unstable in the air. (Smith, l. c.)

Rubidium bromopalladite, Rb_2PdBr_4 .

Can be cryst. from a very small amount of hot H_2O . (Gutbier, B. 1905, 38. 2388.)

Sodium bromopalladite, $\text{Na}_2\text{PdBr}_4 + 4\frac{1}{2}\text{H}_2\text{O}$.

Very deliquescent. Sol. in H_2O . (Smith, l. c.)

Strontium bromopalladite, $\text{SrPdBr}_4 + 6\text{H}_2\text{O}$.

Stable in the air. Very sol. in H_2O . (Smith, l. c.)

Zinc bromopalladite.

Sol. in H_2O . (v. Bonsdorff.)

Bromophosphatoplatinindiamine phosphate, $\text{BrPt}(\text{N}_2\text{H}_5)_2 + 2\text{H}_2\text{O}$.



Sl. sol. in H_2O . (Cleve.)

Bromophosphoric acid.

Thorium bromophosphate, ThBr_4 ,
 $3(3\text{ThO}_2, 2\text{P}_2\text{O}_5)$.

Insol. in most acids and in fused all carbonates. Decomp. by long boiling w conc. H_2SO_4 . (Colani, C. R. 1909, 149. 21)

Bromoplatinamine bromide,



Sl. sol. in H_2C . (Cleve, Sv. V. A. H. 10 31.)

— nitrite, $\text{Br}_2\text{Pt}(\text{NH}_3\text{NO}_2)_2$.

Very sl. sol. in H_2O . (Cleve.)

Bromoplatinindiamine bromide,



Only sl. sol. in hot H_2O . (Cleve.)

— chloride, $\text{Br}_2\text{Pt}(\text{N}_2\text{H}_5)_2\text{Cl}_2$.

Very sl. sol. in H_2O . (Cleve.)

— dichromate, $\text{Br}_2\text{Pt}(\text{N}_2\text{H}_5)_2\text{Cr}_2\text{O}_7$.

Sl. sol. in H_2O .

— nitrate, $\text{Br}_2\text{Pt}(\text{N}_2\text{H}_5\text{NO}_3)_2$.

Sl. sol. in cold, rather easily sol. in hot H_2O . (Cleve.)

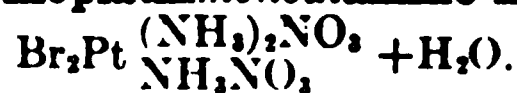
— phosphate, $\text{Br}_2\text{Pt}[\text{N}_2\text{H}_5\text{PO}_2(\text{OH})_2]_2 + 2\text{H}_2\text{O}$.

Rather easily sol. in hot H_2O . (Cleve.)

— sulphate, $\text{Br}_2\text{Pt}(\text{N}_2\text{H}_5)_2\text{SO}_4$.

Very sl. sol. in H_2O .

Bromoplatinmonodiamine nitrate,



Easily sol. in H_2O .

— sulphate, $\text{Br}_2\text{Pt} \begin{array}{c} (\text{NH}_3)_2 \\ \text{NH}_3 \end{array} \text{SO}_4 + \text{H}_2\text{O}$.

Moderately sol. in H_2O . (Cleve.)

Bromoplatinsemidiamine bromide,



Sl. sol. in cold H_2O . (Cleve.)

Bromodiplatinindiamine anhydronitrate,



Sol. in $\text{HNO}_3 + \text{Aq}$.

iplatindiamine chloride,
 $\text{Pt}_2(\text{N}_2\text{H}_4)_4\text{Cl}_4$.

(Cleve.)

rate, $\text{Br}_2\text{Pt}_2(\text{N}_2\text{H}_4)_4(\text{NO}_3)_4 + 2\text{H}_2\text{O}$.
 rarely sol. in hot H_2O .

iphate, $\text{Br}_2\text{Pt}_2(\text{N}_2\text{H}_4)_4(\text{SO}_4)_2 + 2\text{H}_2\text{O}$.
 (Cleve.)

latinic acid, $\text{H}_2\text{PtBr}_6 + 9\text{H}_2\text{O}$.

deliquescent, and sol. in H_2O , alcohol,
 chloroform, or acetic acid. (Topsoë,
 38. 273.)

um bromoplatinate, $(\text{NH}_4)_2\text{PtBr}_6$.

200 pts. H_2O at 15° . (Topsoë.)

s. $(\text{NH}_4)_2\text{PtBr}_6 + \text{Aq sat. at } 20^\circ$ con-
 9 pt. dry salt. (Halberstadt, B. 17.

bromoplatinate, $\text{BaPtBr}_6 + 10\text{H}_2\text{O}$.
 iquescent. Very sol. in H_2O .

bromoplatinate, Cs_2PtBr_6 .

l. in dil. $\text{HBr} + \text{Aq}$. (Obermaier,

bromoplatinate, $\text{CaPtBr}_6 + 12\text{H}_2\text{O}$.
 iquescent. Very sol. in H_2O .

romoplatinate, $\text{CoPtBr}_6 + 12\text{H}_2\text{O}$.
 iescent.

romoplatinate, $\text{CuPtBr}_6 + 8\text{H}_2\text{O}$.
 deliquescent; sol. in H_2O .

omoplatinate, PbPtBr_6 .

sol. in H_2O , but decomp. by large

tetrabromoplatinate, $[\text{PtBr}_4(\text{OH})_2]\text{Pb}$,
 H.

in H_2O . (Miolati, C. C. 1900, II.

um bromoplatinate, $\text{MgPtBr}_6 +$
 H_2O .

deliquescent.

ese bromoplatinate, $\text{MnPtBr}_6 +$
 H_2O .

H_2O . Sol. in H_2O .

tetrabromoplatinate,
 $[\text{PtBr}_4(\text{OH})_2]\text{Hg}$

in H_2O . (Miolati, C. C. 1900, II.

romoplatinate, $\text{NiPtBr}_6 + 12\text{H}_2\text{O}$.
 iescent.

m bromoplatinate, K_2PtBr_6 .

l. in H_2O . Insol. in alcohol. (v.
 T, Pogg. 19. 344.)

Sol. in 10 pts. boiling H_2O . (Pitkin, C. N.
 41. 218.)

100 pts. $\text{K}_2\text{PtBr}_6 + \text{Aq sat. at } 20^\circ$ contain
 2.02 pts. dry salt. (Halberstadt, B. 17. 2962.)

Praseodymium bromoplatinate, $\text{PrBr}_3, \text{PtBr}_6$
 $+ 10\text{H}_2\text{O}$.

Deliquescent; very sol. in H_2O ; sol. in HBr .
 (Von Schule, Z. anorg. 1898, 18. 353.)

Rubidium bromoplatinate, Rb_2PtBr_6 .

Sl. sol. in dil. $\text{HBr} + \text{Aq}$. (Obermaier.
 Dissert.)

Silver bromoplatinate, Ag_2PtBr_6 .

Insol. in H_2O . (Miolati, C. C. 1900, II.
 810.)

Silver tetrabromoplatinate, $[\text{PtBr}_4(\text{OH})_2]\text{Ag}_2$.
 Ppt.; insol. in H_2O . (Miolati, l. c.)

Sodium bromoplatinate, $\text{Na}_2\text{PtBr}_6 + 6\text{H}_2\text{O}$.
 Easily sol. in H_2O and alcohol.

Strontium bromoplatinate, $\text{SrPtBr}_6 + 10\text{H}_2\text{O}$.
 Sl. deliquescent. Very sol. in H_2O .

Thallium tetrabromoplatinate,
 $[\text{PtBr}_4(\text{OH})_2]\text{Tl}_2$.

Insol. in H_2O . (Miolati, C. C. 1900, II.
 810.)

Ytterbium bromoplatinate, $\text{YbBr}_3, 3\text{H}_2\text{PtBr}_6$
 $+ 30\text{H}_2\text{O}$.

Ppt. (Cleve, Z. anorg. 1902, 32. 138.)

Zinc bromoplatinate, $\text{ZnPtBr}_6 + 12\text{H}_2\text{O}$.
 Sol. in H_2O .

Bromoplatinocyanhydric acid,

$\text{H}_2\text{Pt}(\text{CN})_4\text{Br}_2$.

See **Perbromoplatinocyanhydric acid**.

Potassium bromoplatinocyanide, $5\text{K}_2\text{Pt}(\text{CN})_4$,
 $\text{K}_2\text{Pt}(\text{CN})_4\text{Br}_2 + 18\text{H}_2\text{O}$.

Sol. in H_2O .

Bromoplatinous acid.

Potassium bromoplatinite, $\text{K}_2\text{PtBr}_4 + 2\text{H}_2\text{O}$.

Extremely sol. in H_2O . (Billmann and An-
 dersen, B. 1903, 36. 1566.)

Bromopurpureochromium bromide,
 $\text{BrCr}(\text{NH}_3)_5\text{Br}_2$.

Less sol. in H_2O than chloropurpureo-
 chromium chloride. (Jørgensen, J. pr. (2)
 25. 83.)

— **bromoplatinate,** $\text{BrCr}(\text{NH}_3)_5\text{PtBr}_6$.
 (Jørgensen, l. c.)

— **chloride,** $\text{BrCr}(\text{NH}_3)_5\text{Cl}_2$.

More sol. in H_2O than the bromide.
 (Jørgensen, l. c.)

Bromopurpureochromium chromate,
 $\text{BrCr}(\text{NH}_3)_5\text{CrO}_4$.Precipitate. (Jørgensen, *l. c.*)— **nitrate**, $\text{BrCr}(\text{NH}_3)_5(\text{NO}_3)_2$.More sol. than bromide and less than chloride. (Jørgensen, *l. c.*)**Bromopurpureocobaltic bromide,**
 $\text{CoBr}(\text{NH}_3)_5\text{Br}_2$.Sol. in 530 pts. H_2O at 16° . Insol. in alcohol, NH_4Br , KBr , or $\text{HBr} + \text{Aq}$. More sol. in hot H_2O containing a little HBr . (Jørgensen, *J. pr.* (2) 19. 49.)**Bromopurpureocobaltic mercuric bromide,**
 $\text{CoBr}(\text{NH}_3)_5\text{Br}_2, 3\text{HgBr}_2$.More sol. in H_2O than the corresponding HgCl_2 salt. (J.)— **bromoplatinate**.Very sl. sol. in cold H_2O . (J.)— **chloride**, $\text{CoBr}(\text{NH}_3)_5\text{Cl}_2$.Difficultly sol. in cold H_2O , but much more easily than the bromide. Insol. in dil. $\text{HCl} + \text{Aq}$, and in alcohol.— **mercuric chloride**, $\text{CoBr}(\text{NH}_3)_5\text{Cl}_2, 3\text{HgCl}_2$.Sl. sol. in H_2O .— **chloroplatinate**.Nearly or quite insol. in H_2O . (J.)— **chromate**, $\text{CoBr}(\text{NH}_3)_5\text{CrO}_4$.Nearly insol. in H_2O .— **dithionate**, $\text{CoBr}(\text{NH}_3)_5\text{S}_2\text{O}_6$.Nearly insol. in H_2O .— **fluosilicate**, $\text{CoBr}(\text{NH}_3)_5\text{SiF}_6$.Very sl. sol. in cold H_2O ; insol. in alcohol.— **nitrate**, $\text{CoBr}(\text{NH}_3)_5(\text{NO}_3)_2$.More sol. in H_2O than the bromide, but less than the chloride. Wholly insol. in dil. $\text{HN}(\text{O})_2 + \text{Aq}$ or alcohol.— **oxalate**, $\text{CoBr}(\text{NH}_3)_5\text{C}_2\text{O}_4$.Nearly insol. in H_2O .— **sulphate**, $\text{CoBr}(\text{NH}_3)_5\text{SO}_4$.Can be crystallized from very dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Insol. in alcohol. $+6\text{H}_2\text{O}$. Efflorescent.**Bromopurpureorhodium bromide,**
 $\text{BrRh}(\text{NH}_3)_5\text{Br}_2$.Much less easily sol. in H_2O than the chlorochloride. Insol. in dil. $\text{HBr} + \text{Aq}$ and alcohol. (Jørgensen, *J. pr.* (2) 27. 433.)— **bromoplatinate**, $\text{BrRh}(\text{NH}_3)_5\text{PtBr}_6$.Almost insol. in H_2O .— **fluosilicate**, $\text{BrRh}(\text{NH}_3)_5\text{SiF}_6$.Sl. sol. in H_2O . Sol. in boiling $\text{NaOH} + \text{Aq}$ as roseo salt.**Bromopurpureorhodium nitrate,**
 $\text{BrRh}(\text{NH}_3)_5(\text{NO}_3)_2$.Sl. sol. in H_2O , but much more sol. than the bromide.**Bromorhodos acid.****Ammonium bromorhodite**, $(\text{NH}_4)_2\text{RhBr}_4$.Sol. in H_2O . (Goloubkine, *Chem. Sc.* 1911, 100 (2) 45.)Sol. in H_2O . (Gutbier, *B.* 1908, 41. 215.)**Barium bromorhodite**, BaRhBr_4 .Sol. in H_2O . (Goloubkine, *l. c.*)**Cæsium bromorhodite**, Cs_2RhBr_4 .Difficultly sol. in H_2O . (Gutbier, *l. c.*)**Potassium bromorhodite**, K_2RhBr_4 .Very sol. in H_2O . (Goloubkine, *l. c.*)Sol. in H_2O . (Gutbier, *l. c.*)**Rubidium bromorhodite**, Rb_2RhBr_4 .Sol. in H_2O . (Goloubkine, *l. c.*)Difficultly sol. in H_2O . (Gutbier, *l. c.*)**Sodium bromorhodite**, Na_2RhBr_4 .Very sol. in H_2O . (Goloubkine, *l. c.*)**Bromoruthenic acid.****Potassium bromoruthenate**, K_2RuBr_6 .Very sol. in H_2O . (Howe, *J. Am. Ch. Soc.* 1904, 26. 946.)**Potassium aquobromoruthenate,**
 $\text{K}_2\text{Ru}(\text{H}_2\text{O})\text{Br}_6$.Ppt. (Howe, *l. c.*)**Rubidium bromoruthenate**, Rb_2RuBr_6 .Sol. in H_2O . (Howe, *l. c.*)**Rubidium aquobromoruthenate,**
 $\text{Rb}_2\text{Ru}(\text{H}_2\text{O})\text{Br}_6$.Ppt. (Howe, *l. c.*)**Bromoruthenious acid.****Cæsium bromoruthenite**, $\text{CsRuBr}_6 + \text{H}_2\text{O}$.Ppt. (Howe, *J. Am. Chem. Soc.* 1904, 945.)**Potassium bromoruthenite**, K_2RuBr_6 .Very sol. in H_2O with decomp. Very in dil. HBr . (Howe, *l. c.*)**Rubidium bromoruthenite**, $\text{Rb}_2\text{RuBr}_6 + \text{H}_2\text{O}$.Sol. in dil. HBr . (Howe, *l. c.*)**Bromoselenic acid.****Ammonium bromoselenate**, $(\text{NH}_4)_2\text{SeBr}_6$.Sol. in H_2O with decomp. (Muthmann; Schäfer, *B.* 26. 1008.)**Cæsium bromoselenate**, Cs_2SeBr_6 .Sl. sol. in H_2O . (Lenher, *J. Am. Ch. Soc.* 1898, 20. 571.)**Potassium bromoselenate**, K_2SeBr_6 .As NH_4 salt. (M. and S.)

m bromoselenate, Rb_2SeBr_6 .
 ol. in H_2O than K salt. (Lenher, *l. c.*)

pyroselenious acid.

ium bromopyroselenite, $\text{NH}_4\text{Br}, 2\text{SeO}_2, \text{H}_2\text{O}$.
 easily sol. in H_2O than corresponding
 bound. (Muthmann and Schäfer, B. 5. 1014.)

um bromopyroselenite, $\text{KBr}, 2\text{SeO}_2 + \text{H}_2\text{O}$.
 n H_2O . (Muthmann and Schäfer, B. 5.)

smic acid.

ium bromosmate, $(\text{NH}_4)_2\text{OsBr}_6$.
 sl. sol. in H_2O . (Rosenheim, Z. anorg. 1. 135.)

i bromosmate, Cs_2OsBr_6 .
 ly insol. in H_2O and dil. HBr . (Gut- 1913, 46. 2103.)

um bromosmate, K_2OsBr_6 .
 sl. sol. in H_2O . (Rosenheim, *l. c.*)

um bromosmate, Rb_2OsBr_6 .
 ultly sol. in H_2O and in dil. HBr .
 r, *l. c.*)

bromosmate, Ag_2OsBr_6 .
 insol. in H_2O . (Rosenheim, *l. c.*)

bromosmate, $\text{Na}_2\text{OsBr}_6 + 4\text{H}_2\text{O}$.
 n H_2O . (Rosenheim, *l. c.*)

stannic acid, $\text{H}_2\text{SnBr}_6 + 8\text{H}_2\text{O}$.
 deliquescent. Sol. in H_2O . (Seubert, 794.)

ium bromostannate, $(\text{NH}_4)_2\text{SnBr}_6$.
 deliquescent, and sol. in H_2O . (Ray-
 mann and Preis, A. 223. 323.)

a bromostannate.
 n H_2O . (Raymann and Preis.)

n bromostannate, $\text{CaSnBr}_6 + 6\text{H}_2\text{O}$.
 deliquescent. Sol. in H_2O . (Ray-
 mann and Preis.)

bromostannate, $\text{CoSnBr}_6 + 10\text{H}_2\text{O}$.
 uescent. (Raymann and Preis.)

s bromostannate, $\text{FeSnBr}_6 + 6\text{H}_2\text{O}$.
 uescent. (Raymann and Preis.)

a bromostannate, $\text{Li}_2\text{SnBr}_6 + 6\text{H}_2\text{O}$.
 emely deliquescent. (Leteur, C. R. 1.)

sium bromostannate, $\text{MgSnBr}_6 + \text{H}_2\text{O}$.
 uescent. (Raymann and Preis.)

Manganous bromostannate, $\text{MnSnBr}_6 + 6\text{H}_2\text{O}$.
 Deliquescent. (Raymann and Preis.)

Nickel bromostannate, $\text{NiSnBr}_6 + 8\text{H}_2\text{O}$.
 Deliquescent. (Raymann and Preis.)

Potassium bromostannate, K_2SnBr_6 .
 Sol. in H_2O . (Topsoë.)

Rubidium bromostannate.
 Sol. in H_2O . (Raymann and Preis.)

Sodium bromostannate, $\text{Na}_2\text{SnBr}_6 + 6\text{H}_2\text{O}$.
 Not deliquescent, but extremely sol. in
 H_2O . (Seubert, B. 20. 796.)

Strontium bromostannate, $\text{SrSnBr}_6 + 6\text{H}_2\text{O}$.
 Very hygroscopic, and sol. in H_2O . (Ray-
 mann and Preis.)

Bromosulphatoplatindiamine sulphate,

$$\begin{array}{c} \text{Br} > \text{Pt}(\text{N}_2\text{H}_5)_2\text{SO}_4 \\ \text{SO}_4 > \text{Pt}(\text{N}_2\text{H}_5)_2\text{SO}_4 \\ \text{Br} > \text{Pt}(\text{N}_2\text{H}_5)_2\text{SO}_4 \end{array} + \text{H}_2\text{O}.$$

 Rather easily sol. in hot H_2O .

Bromosulphobismuthous acid.

Cuprous bromosulphobismuthite, $2\text{Cu}_2\text{S}, \text{Bi}_2\text{S}_3, 2\text{BiSBr}$.
 Stable in the air and insol. in H_2O at ord.
 temp. Partially decomp. by boiling H_2O .
 Decomp. by mineral acids with the evolution
 of H_2S . (Ducatte, C. R. 1902, 134. 1212.)

Lead bromosulphobismuthite, $\text{PbS}, \text{Bi}_2\text{S}_3, 2\text{BiSBr}$.
 Insol. in H_2O . Decomp. by boiling H_2O .
 Decomp. by dil. mineral acids with evolution
 of H_2S . (Ducatte, *l. c.*)

Bromotantalum bromide, $(\text{Ta}_6\text{Br}_{12})\text{Br}_2 + 7\text{H}_2\text{O}$.
 Stable in the air when in the solid state.
 Sol. in H_2O without decomp. Sol. in propyl
 alcohol. (Chapin, J. Am. Chem. Soc. 1910,
 32. 328.)

Bromotantalum chloride, $(\text{Ta}_6\text{Br}_{12})\text{Cl}_2 + 7\text{H}_2\text{O}$.
 (Chapin, *l. c.*)

Bromotantalum hydroxide, $(\text{Ta}_6\text{Br}_{12})(\text{OH})_2 + 10\text{H}_2\text{O}$.
 Sl. sol. in HCl . Stable in the air below
 100° .
 Sol. in alcohol. Insol. in ether. (Chapin,
l. c.)

Bromotantalum iodide, $(\text{Ta}_6\text{Br}_{12})\text{I}_2 + 7\text{H}_2\text{O}$.
 (Chapin, *l. c.*)

Bromotelluric acid.

Ammonium bromotellurate, $(\text{NH}_4)_2\text{TeBr}_6$.
 Less sol. in H_2O than K salt. (Muthmann
 and Schmidt, B. 1893, 26. 1011.)

Cæsium bromotellurate, Cs_2TeBr_6 .Decomp. by H_2O .100 pts. $\text{HBr} + \text{Aq}$ (sp. gr. 1.49) dissolve 0.02 pt. at 22° .100 pts. $\text{HBr} + \text{Aq}$ (sp. gr. 1.08) dissolve 0.13 pt. at 22° .

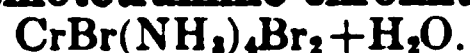
Insol. in alcohol. (Wheeler, Sill. Am. J. 145. 267.)

Potassium bromotellurate, $\text{K}_2\text{TeBr}_6 + 3\text{H}_2\text{O}$.Sol. in little, decomp. by much H_2O . (v. Hauer.)Contains $2\text{H}_2\text{O}$. (Wheeler, Sill. Am. J. 145. 267.)

Efflorescent.

100 pts. $\text{HBr} + \text{Aq}$ (sp. gr. 1.49) dissolve 6.57 pts. at 22° .100 pts. $\text{HBr} + \text{Aq}$ (sp. gr. 1.08) dissolve 62.90 pts. at 22° .

Anhydrous. Stable on air. (Wheeler.)

Rubidium bromotellurate, Rb_2TeBr_6 .Sol. in a little hot H_2O , but H_2TeO_3 separates on cooling.100 pts. $\text{HBr} + \text{Aq}$ (sp. gr. 1.49) dissolve 0.25 pt. at 22° .100 pts. $\text{HBr} + \text{Aq}$ (sp. gr. 1.08) dissolve 3.88 pts. at 22° . (Wheeler.)**Bromotetramine chromium bromide,**Easily sol. in H_2O . (Cleve.)**— chloride, $\text{CrBr}(\text{NH}_3)_4\text{Cl}_2 + \text{H}_2\text{O}$.**Sol. in H_2O . (Cleve.)**— sulphate, $\text{CrBr}(\text{NH}_3)_4\text{SO}_4 + \text{H}_2\text{O}$.**Easily sol. in H_2O . (Cleve.)**Bromotetramine cobaltic sulphate,**Sol. in H_2O . (Vortmann and Blasberg, B. 22. 2652.)**Cadmium, Cd.**Not attacked by H_2O . Sol. in HCl , or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, but more easily in $\text{HNO}_3 + \text{Aq}$. Sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.Chemically pure Cd like Zn is almost insol. in dil. acids, with the exception of HNO_3 . (Weeren, B. 1891, 24. 1798.)Sol. in $\text{HClO}_4 + \text{Aq}$ without evolution of H_2 . (Hendrixson, J. Am. Chem. Soc. 1904, 26. 756.)Cadmium is sol. in molten CdCl_2 and can be recryst. therefrom. (Auerbach, Z. anorg. 1901, 28. 42.)From 4 g. Cd in 32 g. molten CdCl_2 at 650° , 2.197 g. were dissolved in $\frac{1}{2}$ hr. (Helfenstein, Z. anorg. 1900, 23. 295.)Moderately quickly sol. in $\text{K}_2\text{S}_2\text{O}_8 + \text{Aq}$. More slowly sol. in $(\text{NH}_4)_2\text{S}_2\text{O}_8 + \text{Aq}$. (Levi, Gazz. ch. it. 1908, 38 (1) 583.)Sol. in $(\text{NH}_4)_2\text{S}_2\text{O}_8 + \text{Aq}$ without evolution of gas. (Turrentine, J. phys. Chem. 1907, 11. 627.)Sol. in sulphostannates + Aq . (Storch, B. 1883, 16. 2015.) $\frac{1}{2}$ ccm. oleic acid dissolves 0.0293 g. Cd in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Not attacked by sugar solution. (Klein and Berg, C. R. 102. 1170.)

Cadmium amalgam, Cd_2Hg_7 .Stable from 0° – 44° . Can be cryst. from Hg without decomp. if temp. does not exceed 44° . (Kerp. Z. anorg. 1900, 25. 68.)**Cadmium amide, $\text{Cd}(\text{NH}_2)_2$.**Decomp. by H_2O . (Bohart, J. phys. Chem. 1915, 19. 543.)**Cadmium arsenide, Cd_3As_2 .**

(Descamps, C. R. 86. 1022.)

 Cd_3As_2 . Sol. in dil. cold HNO_3 . Attacked by aqua regia. (Granger, C. R. 1904, 138. 575.)**Cadmium azoimide, $\text{Cd}(\text{N}_3)_2$.**

Ppt. (Curtius, J. pr. 1898, (2) 58. 294.)

Cadmium subbromide, Cd_4Br_7 .Decomp. by H_2O . (Morse and Jones, Am. Ch. J. 1890, 12. 490.)**Cadmium bromide, CdBr_2 .**Deliquescent. Very sol. in H_2O .Solubility in H_2O at t°

t°	% CdBr_2	t°	% CdBr_2
—4	32.0	48	60.0
—1	34.7	71	61.2
+1	36.3	104	61.8
2	36.0	155	63.7
9	41.9	170	65.2
14	46.0	215	69.9
25	52.6	232	70.1
35	59.6	245	71.5

Solid phase above 100° is $\text{CdBr}_2 + 1\frac{1}{2}\text{H}_2\text{O}$. (Étard, A. ch. 1894, (7) 2. 541.)See also under $\text{CdBr}_2 + \text{H}_2\text{O}$ and $\text{CdBr}_2 + 4\text{H}_2\text{O}$.Sp. gr. of $\text{CdBr}_2 + \text{Aq}$ at 19.5° containing:

	5	10	15	20	25 % CdBr_2
	1.043	1.090	1.141	1.199	1.260
	30	35	40	45	50 % CdBr_2
	1.326	1.400	1.481	1.578	1.680

(Kremers, calculated by Gerlach, Z. anal. 8. 280.)

 $\text{CdBr}_2 + \text{Aq}$ containing 18.06% CdBr_2 has sp. gr. $20^\circ/20^\circ = 1.1378$. $\text{CdBr}_2 + \text{Aq}$ containing 21.39% CdBr_2 has sp. gr. $20^\circ/20^\circ = 1.1666$.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 282.)

Sp. gr. of $\text{CdBr}_2 + \text{Aq}$ containing 35.84% $\text{CdBr}_2 = 1.4231$ at $19.4^\circ/4^\circ$. (Hallwachs, W. Ann. 1899, 68. 27.)

Sp. gr. of $\text{CdBr}_2 + \text{Aq}$ at $18^\circ/4^\circ$.				
% CdBr_2	33.289	23.973	20.552	11.983
Sp. gr.	1.384	1.252	1.209	1.112
% CdBr_2	6.543	3.734	1.927	
Sp. gr.	1.106	1.030	1.017	
(de Muynck, W. Ann. 1894, 53. 561.)				

Sp. gr. of $\text{CdBr}_2 + \text{Aq}$ at 18° .					
% CdBr_2	1	5	10	15	20
Sp. gr.	1.0072	1.0431	1.0907	1.1432	1.1991
% CdBr_2	25	30	35	40	43
Sp. gr.	1.2605	1.3296	1.4052	1.4915	1.5467
(Grotrian, W. Ann. 1883, 18. 193.)					

Sp. gr. of $\text{CdBr}_2 + \text{Aq}$.			
% CdBr_2	t°	Sp. gr. at t°	Sp. gr. at 18°
0.0324	17.90	0.99901	0.99900
	22.75	0.99702	
0.0748	17.23	0.99949	0.99935
	21.50	0.99863	
0.154	17.67	1.00008	1.00002
	23.10	0.99896	
0.253	17.23	1.00119	0.00100
	22.95	0.99986	
0.506	18.07	0.00308	1.00310
	22.65	1.00212	
1.013	18.00		1.00750

(Wershofen, Z. phys. Ch. 1890, 5. 493.)

Sp. gr. of $\text{CdBr}_2 + \text{Aq}$ at 20° .		
Normality of $\text{CdBr}_2 + \text{Aq}$	% CdBr_2	Sp. gr.
2.774	46.574	1.6198
1.997	37.53	1.4469
0.973	22.53	1.2293
0.5138	12.46	1.1211

(Forchheimer, Z. phys. Ch. 1900, 34. 29.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913, 84, 27.)

Sol. in $\text{HCl} + \text{Aq}$, $\text{HC}_2\text{H}_3\text{O}_2$, alcohol, or ether. (Berthelot, A. ch. 44. 387.)

Sol. in 0.94 pt. H_2O , 3.4 pts. abs. alcohol, 250 pts. ether, and 16 pts. alcohol-ether (1 : 1) (Eder, Dingl. 221. 89.)

Anhydrous CdBr_2 is sol. in acetone. (Krug and M'Elroy.)

1 g. CdBr_2 is sol. in 64.5 g. acetone at 18° . Sp. gr. of sat. solution $18^\circ/4^\circ = 0.8073$. (Naumann, B. 1904, 37. 4337.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

Insol. in mustard oil. (Mathews, J. phys. Chem. 1905, 9, 647.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42, 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Sol. in chinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

100 g. benzonitrile dissolve 0.857 g. CdBr_2 at 18° . (Naumann, B. 1914, 47. 1370.)

Mol. weight determined in piperidine. (Ferchland, Z. anorg. 1897, 15, 17.)

+ H_2O . Solubility in H_2O .

100 g. of the sat. solution contain at:
 35° 40° 45° 60° 80° 100°
 60.29 60.65 60.75 61.10 61.29 61.63 g. CdBr_2 .
 (Dietz, Z. anorg. 1899, 20. 261.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. (Étard, A. ch. 1894, (7) 2. 541.)
 + $4\text{H}_2\text{O}$. Efflorescent. (Rammelsberg, Pogg. 55. 241.)

Solubility in H_2O .

100 g. of the sat. solution contain at:
 0° 18° 30° 38°
 37.92 48.90 56.90 61.84 g. CdBr_2 .
 Sp. gr. of sat. solution at $18^\circ = 1.683$.
 (Dietz, Z. anorg. 1899, 20. 261.)

100 g. sat. solution of $\text{CdBr}_2 + 4\text{H}_2\text{O}$ in absolute alcohol contain 20.93 g. CdBr_2 at 15° .

100 g. sat. solution of $\text{CdBr}_2 + 4\text{H}_2\text{O}$ in absolute ether contain 0.4 g. CdBr_2 at 15° . (Eder, Dingl. 221. 89.)

Cadmium hydrogen bromide.

Decomp. by H_2O . (Berthelot, C. R. 91. 1024.)

Cadmium caesium bromide, $\text{CdBr}_2, \text{CsBr}$.

Easily sol. in H_2O . (Wells and Walden, Z. anorg. 5. 270.)

$\text{CdBr}_2, 2\text{CsBr}$. Decomp. by H_2O into above comp. (W. and W.)

$\text{CdBr}_2, 3\text{CsBr}$. Decomp. by H_2O into $\text{CdBr}_2, \text{CsBr}$. (W. and W.)

Cadmium potassium bromide, $\text{CdBr}_2, \text{KBr} + \frac{1}{2}\text{H}_2\text{O}$.

Sol. in 0.79 pt. H_2O at 15° ; pptd. by alcohol and ether. (Eder, Dingl. 221. 89.)

+ H_2O . Sol. in H_2O without decomp. from 0.4° – 112.5° . (Rimbach, B. 1905, 38, 1554.)

100 pts. of the solution contain at:
 0.4° 15.8° 50° 112.5°
 53.75 58.68 68.25 78.10 pts. of the salt.

$\text{CdBr}_2, 4\text{KBr}$. Sol. in 1.40 pts. H_2O at 15° ; pptd. by alcohol and ether. (Eder, Dingl. 221. 89.)

Cannot be prepared in a pure state as it is decomp. by H_2O below 160° . (Rimbach, B. 1905, 38. 1560.)

Cadmium rubidium bromide, $\text{CdBr}_2, \text{RbBr}$.

Sol. in H_2O without decomp. from 0.4° to 107.5° .

100 pts. of the solution contain at:
 0.4° 14.5° 49.2° 107.5°
 32.65 41.87 58.54 75.77 pts. of the salt.
 (Rimbach, B. 1905, 38. 1556.)

$\text{CdBr}_2, 4\text{RbBr}$. Sol. in H_2O without decomp. from 0.5° to 114.5° .

100 pts. of the solution contain at:
 0.5° 13.5° 51.5° 114.5°
 47.95 55.17 68.82 79.04 pts. of the salt.
 (Rimbach, B. 1905, 38. 1561.)

Cadmium sodium bromide, $\text{CdBr}_2, \text{NaBr} + 2\frac{1}{2}\text{H}_2\text{O}$.

Sol. at 15° in 1.04 pts. H_2O , 3.7 pts. abs. alcohol, and 190 pts. ether (sp. gr. 0.729). (Eder, Dingl. 221. 89.)

$3\text{CdBr}_2, 2\text{NaBr} + 6\text{H}_2\text{O}$. Stable in conc. solutions and decomp. only by great dilution. (Jones and Knight, Am. Ch. J. 1899, 22. 134.)

Cadmium bromide ammonia, $\text{CdBr}_2, 2\text{NH}_3$.

Can be crystallized out of warm $\text{NH}_4\text{OH} + \text{Aq}$. (Croft, Phil. Mag. 21. 356.)

$\text{CdBr}_2, 3\text{NH}_3$. (Tassily, C. R. 1897, 124. 1022.)

$\text{CdBr}_2, 4\text{NH}_3$. Decomp. by H_2O . (Croft.)

Cadmium bromide cupric oxide, $\text{CdBr}_2, 3\text{CuO} + 3\text{H}_2\text{O}$. (Mailhe, A. ch. 1902, (7) 27. 383.)

Cadmium bromide hydrazine, $\text{CdBr}_2, 2\text{N}_2\text{H}_4$.

Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Franzen, Z. anorg. 1908, 60. 280.)

Cadmium bromide hydroxylamine, $\text{CdBr}_2, 2\text{NH}_2\text{OH}$.

Sol. in hot H_2O with formation of a basic salt. Sol. in dil. acids. Insol. in alcohol and ether. (Adams, Am. Ch. J. 1902, 28. 218.)

Cadmium subchloride, Cd_4Cl_7 .

Decomp. by H_2O and by acids. (Morse and Jones, Am. Ch. J. 1890, 12. 490.)

Cadmium chloride, CdCl_2 .

Sol. at 20° 40° 60° 80° 100°
in 0.71 0.72 0.72 0.70 0.67 pts. H_2O .
(Kremers. Pogg: 103. 57.)

Sat. $\text{CdCl}_2 + \text{Aq}$ contains % CdCl_2 at t°.

t°	%CdCl ₂	t°	%CdCl ₂
—7	43.5	120	63.0
+1	47.6	150	64.8
6	49.7	165	68.2
7	51.3	170	68.4
10	51.6	180	70.1
19	52.7	190	71.9
25	52.9	200	72.0
61	57.9	235	76.0
82	58.8	270	77.7

(Étard, A. ch. 1894, (7) 2. 536.)

100 mol. H_2O dissolve at:

19.3° 29.7° 40.1° 54.5°
10.94 12.74 13.15 13.16 mol. CdCl_2 .

(Sudhaus, Miner. Jahrb. Beil.-Bd. 1914, 37. 19.)

See also under $\text{CdCl}_2 + \text{H}_2\text{O}$, $\text{CdCl}_2 + 2\frac{1}{2}\text{H}_2\text{O}$, and $\text{CdCl}_2 + 4\text{H}_2\text{O}$.

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$ containing pts. CdCl_2 to 100 pts. H_2O .

13	26.9	41	pts. CdCl_2 ,
1.1068	1.2106	1.3100	
55.8	72.5	114.2	pts. CdCl_2 .
1.4060	1.5060	1.7266	

(Kremers, Pogg. 103. 57.)

$\text{CdCl}_2 + \text{Aq}$ containing 8.91% CdCl_2 has sp. gr. 20°/20° = 1.0715. (Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 282.)

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$ at room temp. containing:

% CdCl_2	11.09	16.30	24.786
Sp. gr.	1.1093	1.1813	1.3199

(Wagner, W. Ann. 1883, 18. 266.)

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$ at 18°/4°.

% CdCl_2	57.524	41.547	29.977
Sp. gr.	1.852	1.515	1.330

% CdCl_2	21.431	14.761
Sp. gr.	1.210	1.142

(de Muynck, W. Ann. 1894, 53. 561.)

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$ at 18°.

% CdCl_2	1	5	10	15
Sp. gr.	1.0063	1.0436	1.0919	1.1443
% CdCl_2	20	25	30	35
Sp. gr.	1.2007	1.2620	1.3305	1.4071
% CdCl_2	40	45	50	
Sp. gr.	1.4878	1.5775	1.6799	

(Grotrian, W. Ann. 1883, 18. 193.)

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$ at 25°.

Concentration of $\text{CdCl}_2 + \text{Aq}$	Sp. gr.
1-normal	1.0779
$\frac{1}{2}$ -normal	1.0394
$\frac{1}{4}$ -normal	1.0197
$\frac{1}{8}$ -normal	1.0098

(Wagner, Z. phys. Ch. 1890, 5. 36.)

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$.

% CdCl_2	t°	Sp. gr. at t°	Sp. gr. at 1
0.0503	17.59	0.99920	0.999
	24.27	0.99781	
0.0999	17.70	0.99964	0.999
	22.06	0.99833	
0.200	18.31	1.00038	1.000
	24.00	0.99920	
0.399	16.86	1.00239	1.002
	24.21	1.00083	
0.599	17.49	1.00406	1.003
	25.12	1.00238	
0.769	17.58	1.00580	1.005
	21.76	1.00496	
0.997	17.55	1.00754	1.007
	19.65	1.00713	

(Wershofen, Z. phys. Ch. 1890, 5. 492.)

Sp. gr. of $\text{CdCl}_2 + \text{Aq}$ at t°.

t°	Normality of $\text{CdCl}_2 + \text{Aq}$	g. CdCl_2 in 100 g. of solution	Sp. gr. t°/4°
20.5	3.80	44.42	1.56
"	2.61	34.22	1.30
"	1.76	25.90	1.24
"	1.29	19.91	1.16
"	0.93	14.88	1.14
"	0.52	8.84	1.06

(Oppenheimer, Z. phys. Ch. 1898, 27. 454)

Sp. gr. of CdCl ₂ +Aq at t°.		
t°	Concentration of CdCl ₂ +Aq	Sp. gr.
22	1 pt. CdCl ₂ in 1.3458 pts. H ₂ O	1.6128
18.7	1 " " " 2.7005 " "	1.2896
17.2	1 " " " 53.988 " "	1.0155
16	1 " " " 54.18 " "	1.0152
17	1 " " " 57.479 " "	1.0136
22	1 " " " 77.232 " "	1.0076

(Hittorf, Z. phys. Ch. 1902, 39. 628.)

Solubility in KCl+Aq at t°.			
t°	100 g. H ₂ O dissolve		Solid phase
	g. CdCl ₂	g. KCl	
9.3	111.30	...	CdCl ₂ +2½H ₂ O
	59.59	6.70	CdCl ₂ +2½H ₂ O+CdCl ₂ , KCl+H ₂ O
	26.98	11.09	CdCl ₂ , KCl+H ₂ O
	11.61	30.04	CdCl ₂ , KCl+H ₂ O+CdCl ₂ , 4KCl
	1.44	34.76	CdCl ₂ , 4KCl+KCl
17	...	33.94	KCl
	129.65	...	CdCl ₂ +3½H ₂ O
	97.62	0.70	CdCl ₂ +2½H ₂ O
	68.23	7.08	CdCl ₂ +2½H ₂ O+CdCl ₂ , KCl+H ₂ O
	47.12	9.89	CdCl ₂ , KCl+H ₂ O
	32.67	13.06	"
	24.26	16.10	"
	15.99	25.97	"
	15.47	33.58	CdCl ₂ , KCl+H ₂ O+CdCl ₂ , 4KCl
	2.42	37.66	CdCl ₂ , 4KCl+KCl
34.5	...	37.21	KCl
	133.85	...	CdCl ₂ +H ₂ O
	92.15	2.70	CdCl ₂ +H ₂ O+CdCl ₂ , KCl+H ₂ O
	51.90	11.50	CdCl ₂ , KCl+H ₂ O
	37.91	15.21	"
	24.45	21.73	"
	18.97	35.51	"
	19.92	37.63	CdCl ₂ , KCl+H ₂ O+CdCl ₂ , 4KCl
	2.98	40.45	CdCl ₂ , 4KCl+KCl
	...	40.36	KCl
54.5	133.90	...	CdCl ₂ +H ₂ O
	102.15	2.32	CdCl ₂ +H ₂ O+CdCl ₂ , KCl+H ₂ O
	44.01	18.39	CdCl ₂ , KCl+H ₂ O
	26.13	43.78	CdCl ₂ , KCl+H ₂ O+CdCl ₂ , 4KCl
	4.20	45.52	CdCl ₂ , 4KCl+KCl
77.2	...	43.00	KCl

(Sudhaus, Miner. Jahrb. Beil.-Bd. 1914, 37. 34.)

Solubility in NaCl+Aq at t°.			
t°	100 g. H ₂ O dissolve		Solid phase
	g. CdCl ₂	g. NaCl	
19.3	111.30	...	CdCl ₂ +2½H ₂ O
	116.64	7.52	CdCl ₂ +2½H ₂ O+CdCl ₂ , 2NaCl+3H ₂ O
	85.15	12.19	CdCl ₂ , 2NaCl+3H ₂ O
	40.01	25.67	"
	5.96	36.76	CdCl ₂ , 2NaCl+3H ₂ O+ NaCl
	...	35.84	NaCl
29.7	129.65	...	CdCl ₂ +2½H ₂ O
	132.67	9.63	CdCl ₂ +2½H ₂ O+CdCl ₂ , 2NaCl+3H ₂ O
	123.54	10.10	CdCl ₂ , 2NaCl+3H ₂ O
	106.16	12.92	"
	91.10	15.41	"
	43.74	27.46	"
40.1	9.43	37.54	CdCl ₂ , 2NaCl+3H ₂ O+ NaCl
	...	35.88	NaCl
54.5	133.85	...	CdCl ₂ +H ₂ O
	137.03	15.14	CdCl ₂ +H ₂ O+CdCl ₂ , 2NaCl+3H ₂ O
	48.17	29.50	CdCl ₂ , 2NaCl+3H ₂ O
	13.31	38.16	CdCl ₂ , 2NaCl+3H ₂ O+ NaCl
	...	36.18	NaCl

At 34.5°, CdCl₂+2½H₂O → CdCl₂+H₂O and water.
(Sudhaus, Miner. Jahrb. Beil.-Bd. 1914, 37. 28.)

Insol. in SbCl₃. (Klemensiewicz, C. A. 1909, 269.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

Insol. or sl. sol. in ethyl alcohol, furfurol, acetophenone, ethyl monochloracetate, ethyl cyanacetate, ethyl oxalate, ethyl nitrate, amyl nitrite, o-nitrotoluene, pyridine, piperidine, and quinoline. Sol. in salicylic aldehyde. (Lincoln, J. phys. Chem. 1899, 3. 461.)

Insol. in anhydrous ether. (Hampe, Ch. Z. 1887, II, 847.)

Readily sol. in alcohol.
100 pts. absolute methyl alcohol dissolve 1.71 pts. CdCl₂ at 15.5°.

100 pts. absolute ethyl alcohol dissolve 1.52 pts. CdCl₂ at 15.5°. (de Bruyn, Z. phys. Ch. 10. 783.)

100 g. CdCl₂+CH₃OH contain 1.5 g. CdCl₂.

at the critical temp. (Centnerszwer, Z. phys. Ch. 1910, **72**. 437.)

Somewhat sol. in acetone. (Krug and M'Elroy.)

Sol. in acetone; insol. in methylal. (Eidmann, C. C. **1899**, II, 1014.)

Insol. in methyl acetate. (Naumann, B. **1909**, **42**. 3790.)

Sol. in ethyl acetate. (Naumann, B. **1904**, **37**. 3601.)

Difficultly sol. in ethylacetate. (Naumann, B. **1910**, **43**. 314.)

Sol. in urethane. (Castoro, Z. anorg. **1899**, **20**. 61.)

At 18°, 100 g. benzonitrile dissolve 0.06332 g. CdCl₂. (Naumann, B. **1914**, **47**, 1370.)

Insol. in toluene. (Baxter and Hines, Am. Ch. J. **1904**, **31**. 222.)

Sol. in chinolin. (Beckmann and Gabel, Z. anorg. **1906**, **51**. 236.)

+H₂O. Solubility in H₂O.

100 g. of the sat. solution contain at:

10°	20°	40°	60°
57.47	57.35	57.51	57.77

80° 100°

58.41 59.52 g. CdCl₂.

110° is bpt. of the sat. solution.

(Dietz, Z. anorg. **1899**, **20**. 257.)

+2½H₂O. Solubility in H₂O.

100 g. of the sat. solution contain at:

-10°	0°	18°	30°	36°
44.35	47.37	52.53	56.27	57.91 g. CdCl ₂ .

Sp. gr. of sat. solution = 1.741.

(Dietz, Z. anorg. **1899**, **20**. 257.)

+4H₂O. Solubility in H₂O.

100 g. of the sat. solution contain at:

-9°	0°	+10°	+15°
43.58	49.39	55.58	59.12 g. CdCl ₂ .

(Dietz, Z. anorg. **1899**, **20**. 257.)

+5H₂O. (Worobieff, Z. anorg. **1898**, **18**. 386.)

Cadmium hydrogen chloride, CdCl₂, 2HCl + 7H₂O.

Decomp. in air. (Berthelot, C. R. **91**. 1024.)

Cadmium caesium chloride, CdCl₂, 2CsCl.

Easily sol. in H₂O and dil. HCl + Aq; insol. in conc. HCl + Aq. (Godeffroy, B. **8**. 9.)

Nearly insol. in CsCl + Aq. (Wells and Walden, Z. anorg. **5**. 266.)

CdCl₂, CsCl. Sl. sol. in H₂O; nearly insol. in CdCl₂ + Aq. (Wells and Walden.)

Cadmium calcium chloride, 2CdCl₂, CaCl₂ + 7H₂O.

Rather deliquescent, and very sol. in H₂O. When ignited is only sl. sol. in H₂O with evolution of heat. (v. Hauer, J. pr. **63**. 432.)

CdCl₂, 2CaCl₂ + 12H₂O. Very deliquescent. (v. Hauer.)

Cadmium cobaltous chloride, 2CdCl₂, CoCl₂ + 12H₂O.

Deliquescent. Sol. in H₂O. (v. Hauer, W. A. B. **17**. 331.)

Cadmium cupric chloride, CdCl₂, CuCl₂ + 4H₂O.

Sol. in H₂O. (v. Hauer, W. A. B. **17**. 331.)

Cadmium hydrazine chloride, CdCl₂, N₂H₄HCl.

Unstable in the air when moist. Very sol. in H₂O; sl. sol. in alcohol; sol. in NH₃ + Aq. (Curtius, J. pr. **1894**, (2) **50**. 334.)

CdCl₂, 2N₂H₄HCl + 4H₂O. Very sol. in H₂O; sl. sol. in alcohol. (Curtius, J. pr. **1894**, (2) **50**. 335.)

Cadmium iron (ferrous) chloride, 2CdCl₂, FeCl₂ + 12H₂O.

Sol. in H₂O. (v. Hauer, W. A. B. **17**. 331.)

Cadmium lithium chloride, CdCl₂, LiCl + 3½H₂O.

Very deliquescent. Decomp. by solution in H₂O, but not in alcohol. (Chassevant, A. ch. (6) **30**. 39.)

Cadmium magnesium chloride, 2CdCl₂, MgCl₂ + 12H₂O.

Deliquescent in moist, stable in dry air. Easily sol. in H₂O with absorption of heat. Much more sol. in hot than in cold H₂O. (v. Hauer.)

Solubility in H₂O at t°.

t°	G. Cd ₂ MgCl ₂ in 100 g. solution	G. Cd ₂ MgCl ₂ in 100 g. H ₂ O
2 4	45.61	83.86
20.8	49.69	98.77
45.5	53.51	115.10
67.2	58.14	138.90
121.8	65.48	189.69

(Rimbach, B. **1897**, **30**. 3084.)

CdCl₂, 2MgCl₂ + 12H₂O. Very deliquescent. (v. Hauer.)

Cadmium manganese chloride, 2CdCl₂, MnCl₂ + 12H₂O.

Deliquescent in moist, efflorescent in dry air. Sol. in H₂O. (v. Hauer.)

Cadmium nickel chloride, CdCl₂, 2NiCl₂ + 12H₂O.

Sol. in H₂O. (v. Hauer, W. A. B. **20**. 40.)
2CdCl₂, NiCl₂ + 12H₂O. Sol. in H₂O. (v. Hauer.)

Cadmium potassium chloride, CdCl₂, KCl + ½H₂O.

Sol. in H₂O without decomp. (v. Hauer.)

H₂O. 100 mol. H₂O dissolve at:
 29.7° 40.1° 54.5°
 3.21 3.72 4.33 mol. CdCl₂, KCl+H₂O.
 haus, Miner. Jahrb. Beil.-Bd. 1914, **37**.
 26.)

Solubility in H₂O at t°.

t°	G. CdKCl ₂ in 100 g. solution	G. CdKCl ₂ in 100 g. H ₂ O
2.6	21.87	27.99
15.9	26.60	36.4
41.5	35.66	55.34
60.6	40.67	68.55
105.1	51.67	106.91

(Rimbach, B. 1897, **30**. 3079.)

lCl₂, 2KCl. 100 pts. H₂O at 15.5° dis-
 33.45 pts. Sl. sol. in alcohol. (Croft,
 Mag. (3) **21**. 356.)

lubility in salts+Aq at 16°.
 lCl₂, 2KCl is sol. without decomp. in the
 wing salt solutions at 16°.

	Mols. salt in 100 mole H ₂ O	In 1 litre of the solution mole			Sp. gr. of the solution
		CdCl ₂	KCl	RCl	
	9.3	0.166	0.663	4.483	1.1380
l ₂	3.8	0.270	1.080	1.887	1.2333
	2.378	0.507	3.195	...	1.214

(Rimbach, B. 1905, **38**. 1568,)

lCl₂, 4KCl. More sol. in H₂O than
 l₂, KCl. (v. Hauer.)

0 g. H₂O dissolve at:
 29.7° 40.1° 54.5°
 49.05 57.55 69.91 g. CdCl₂, 4KCl.
 haus, Miner. Jahrb. Beil.-Bd. 1914, **37**.
 24.)

Solubility in H₂O at t°.

t°	100 pts. solution contain pts.		
	Cd	Cl	K
4.0	3.64	9.84	8.31
23.6	5.66	14.02	11.52
50.2	9.10	18.09	13.60
08.8	11.97	23.08	17.10
09.0	11.91	23.15	17.22

(Rimbach, B. 1897, **30**. 3080.)

decomp. by H₂O.
 can be recryst. without decomp. from LiCl,
 l₂, or MgCl₂+Aq. (Rimbach, B. 1905,
 565.)

The salt is sol. without decomp. in HCl+Aq
 gaining 19.8 mole HCl per 100 mole H₂O
 at 16°.

1. of the solution contains 0.033 mole
 l₂, 0.132 mole KCl and 8.828 mole HCl;
 sp. gr. of the solution = 1.1403. (Rimbach,
 1905, **38**. 1568.)

Cadmium rubidium chloride, CdCl₂, 2RbCl.
 Sol. in H₂O and HCl+Aq. (Godeffroy, B.
 8. 9.)

CdCl₂, RbCl. Solubility in H₂O at t°.
 100 pts. by wt. of the solution contain pts.
 by wt. RbCl, CdCl₂.

t°	Pts. RbCl, CdCl ₂
1.2	12.97
14.5	16.80
41.4	25.31
57.6	30.83
103.9	46.62

CdCl₂, RbCl is sol. in H₂O without decomp.
 from 0-104°. (Rimbach, B. 1902, **35**. 1303.)

CdCl₂, 4RbCl.

Solubility of CdCl₂, 4RbCl and CdCl₂, RbCl
 in H₂O at t°.

t°	In 100 pts. by wt. of the solution			Composition of the solid phase	
	Pts. by wt. Cd	Pts. by wt. Cl	Pts. by wt. Rb	Mol.-% mono- salt	Mol.-% tetra- salt
0.7	0.65	6.52	14.73	30	70
8.8	1.07	7.37	16.13	24	76
13.8	1.32	7.86	16.93	16	84
42.4	3.21	11.35	22.45	14	86
59.0	4.61	13.41	25.31	33	67
108.4	8.94	18.57	31.15

(Rimbach, B. 1902, **35**. 1305.)

Decomp. by H₂O between 0° and 108°.
 (Rimbach, B. 1905, **38**. 1571.)

Sol. in conc. HCl without decomp. (Rim-
 bach, B. 1905, **38**. 1571.)

Not sol. in CaCl₂+Aq and LiCl+Aq with-
 out decomp. (Rimbach, B. 1905, **38**. 1571.)

**Cadmium sodium chloride, CdCl₂, 2NaCl+
 3H₂O.**

Sol. in 1.4 pts. H₂O at 16°. (Croft.)
 100 mol. H₂O dissolve at:
 19.3° 29.7° 40.1° 54.5°
 3.93 4.29 4.73 5.18 mol. CdCl₂,
 2NaCl+3H₂O.

Stable between 19° and 55°.
 (Sudhaus, Miner. Jahrb. Beil.-Bd. 1914, **37**.
 25.)

Sl. sol. in alcohol or wood alcohol. (Croft.)

**Cadmium strontium chloride, 2CdCl₂, SrCl₂+
 7H₂O.**

Sol. in H₂O. (v. Hauer.)

Cadmium chloride ammonia, CdCl₂, 2NH₃.

Nearly insol. in H₂O. (v. Hauer.)
 CdCl₂, 3NH₃+¼H₂C.
 CdCl₂, 4NH₃+½H₂O.
 CdCl₂, 5NH₃. (André, C. R. **104**. 908.)
 CdCl₂, 6NH₃. Difficultly sol. in cold H₂O.
 (Schüler, A. **87**. 34.)

Cadmium chloride cupric oxide, $\text{CdCl}_2 \cdot 3\text{CuO} + 3\text{H}_2\text{O}$.

Not decomp. by H_2O . (Mailhe, A. ch. 1902, (7) 27. 378 and 174.)

Cadmium chloride hydrazine, $\text{CdCl}_2 \cdot 2\text{N}_2\text{H}_4$.
Insol. in H_2O .

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Franzen, Z. anorg. 1908, 60. 279.)

+ H_2O . Insol. in H_2O ; easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Curtius, J. pr. 1894, (2) 50. 345.)

Cadmium chloride hydroxylamine, $\text{CdCl}_2 \cdot 2\text{NH}_2\text{OH}$.

Sl. sol. in cold, somewhat more in warm H_2O . Very sol. in hydroxylamine + Aq . Very sl. sol. in alcohol and other organic solvents. (Crismer, Bull. Soc. (3) 3. 116.)

Aq solution sat. at 20° contains about 1%. (Antonoff, C. C. 1905, II. 810.)

Cadmium fluoride, CdF_2 .

Difficultly sol. in H_2O . Easily sol. in $\text{HF} + \text{Aq}$. (Berzelius, Pogg. 1. 26.)

Very sol. in H_2O ; insol. in 95% alcohol; sol. in HCl , H_2SO_4 , or $\text{HNO}_3 + \text{Aq}$ with evolution of HF . (Poulenc, C. R. 116. 582.)

1 l. H_2O dissolves 0.289 mol. CdF_2 at 25° , or 100 cc. sat. aqueous solution contains 4.36 g. CdF_2 at 25° . (Jaeger, Z. anorg. 1901, 27. 35.)

1 l. of 1.08-N HF dissolves 0.372 mol. CdF_2 at 25° . (Jaeger, Z. anorg. 1901, 27. 35.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

Cadmium ceric fluoride, $\text{CdF}_2 \cdot 2\text{CeF}_4 + 7\text{H}_2\text{O}$.

Ppt. Decomp. by H_2O . (Rimbach, A. 1909, 368. 106.)

Cadmium columbium fluoride.

See Fluocolumbate, cadmium.

Cadmium molybdenyl fluoride.

See Fluoxymolybdate, cadmium.

Cadmium silicon fluoride.

See Fluosilicate, cadmium.

Cadmium stannic fluoride.

See Fluostannate, cadmium.

Cadmium titanium fluoride.

See Fluotitanate, cadmium.

Cadmium zirconium fluoride.

See Fluozirconate, cadmium.

Cadmous hydroxide, CdOH .

Insol. in H_2O . Decomp. by acids into cadmic salt. (Morse and Jones, Am. Ch. J. 12. 488.)

Cadmium hydroxide, CdO_2H_2 .

Insol. in H_2O .

1 l. $\text{CdO}_2\text{H}_2 + \text{Aq}$ contains 0.0026 g. CdO_2H_2

at 25° . (Bodländer, Z. phys. Ch. 1898, 27. 66.)

Solubility in $\text{H}_2\text{O} = 2.6 \times 10^{-4}$. (Herz, Z. anorg. 1900, 24. 126.)

Sol. in acids; very sol. in $\text{NH}_4\text{OH} + \text{Aq}$; insol. in KOH , NaOH , Na_2CO_3 , K_2CO_3 , and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.

Easily sol. in $(\text{NH}_4)_2\text{SO}_4$, NH_4Cl , NH_4NO_3 , and NH_4 succinate + Aq . (Wittstein.)

Freshly pptd. CdO_2H_2 is sol. in alkali haloids + Aq . (Bersch, Z. phys. Ch. 1891, 8. 392.)

Solubility in $\text{NH}_4\text{OH} + \text{Aq}$ increases with increase in concentration of NH_4OH . (Euler, B. 1903, 36. 3401.)

Solubility in $\text{NH}_4\text{OH} + \text{Aq}$ at 25° .

NH_3 norm.	g. CdO per l.
0.5	0.24
1.0	0.62
1.8	1.33
4.6	4.92

(Bonsdorff, Z. anorg. 1904, 41. 187.)

Insol. in ethyl, and methyl amine + Aq . (Wurtz.)

Very sl. sol. in $\text{HCN} + \text{Aq}$ even when freshly pptd. (Schüler, A. 87. 48.)

Not pptd. in presence of Na citrate (Spiller), and many non-volatile organic substances. (Rose.)

Cadmium iodide, CdI_2 .

Sol. in 1.13 pts. H_2O at 15° . (Eder, Dingl. 221. 89.)

Sol. at 20° 40° 60° 80° 100°
in 1.08 1.00 0.93 0.86 0.75 pts. H_2O .

(Kremers, Pogg. 103. 57.)

Sat. $\text{CdI}_2 + \text{Aq}$ contains at:

-4°	$+2^\circ$	$+10^\circ$	13°	24°	32°
42.4	43.7	45.2	44.8	46.5	47.4% CdI_2
54°	64°	76°	94°	95°	135°
49.5	50.1	52.4	55.1	54.7	62.9% CdI_2
140°	165°	185°	202°	202°	255°
63.1	68.1	70.7	73.4	73.2	84.5% CdI_2

(Étard, A. ch. 1894, (7) 2. 545.)

Solubility in H_2O .

100 g. of the sat. solution contain at:

0°	18°	50°	75°	100°
44.39	46.02	49.35	52.65	56.08 g. CdI_2

(Dietz, Z. anorg. 1899, 20. 262.)

Sp. gr. of $\text{CdI}_2 + \text{Aq}$ containing pts. CdI_2 to 100 pts. H_2O .

21.4	43.7	88.5 pts. CdI_2
1.1681	1.328	1.6139

(Kremers, Pogg. 111. 60.)

Sp. gr. of $\text{CdI}_2 + \text{Aq}$ at 19.5° containing:

5	10	15	20	25 % CdI_2 ,
1.044	1.088	1.138	1.194	1.253

30	35	40	45	50 % CdI_2 .
1.319	1.395	1.476	1.575	1.680

(Kremers, calculated by Gerlach, Z. anal. 8. 285.)

Sp. gr. of $\text{CdI}_2 + \text{Aq}$ at 18° .

% CdI_2	1	5	10	15	20
Sp. gr.	1.0071	1.0425	1.0883	1.1392	1.1943

% CdI_2	25	30	35	40	45
Sp. gr.	1.2550	1.3228	1.4000	1.4816	1.5741

(Grottrian, W. Ann. 1883, 18. 193.)

Sp. gr. of $\text{CdI}_2 + \text{Aq}$.

g. CdI_2 per l.	Sp. gr.	g. CdI_2 per l.	Sp. gr.
98.85	1.08	289.5	1.237
197.7	1.162	400	1.328

(Barbier and Roux, Bull. Soc. 1890, (3) 3. 425.)

Sp. gr. of $\text{CdI}_2 + \text{Aq}$.

% CdI_2	t°	Sp. gr. at t°	Sp. gr. at 18°
0.0429	17.68	0.99915	0.99908
	22.88	0.99807	
0.100	17.55	0.99965	0.99956
	22.91	0.99363	
0.204	17.76	1.00052	1.0005
	22.79	0.99948	
0.399	17.40	0.00223	1.0021
	24.30	1.00082	
0.600	18.00		1.0038
0.800	17.44	1.00564	1.0056
	23.11	1.00442	
1.00	18.00		1.0072

(Wershofen, Z. phys. Ch. 1890, 5. 493.)

Sp. gr. $\text{CdI}_2 + \text{Aq}$ at $18^\circ/4^\circ$ containing:

31.123	13.677	9.559 % CdI_2 .
1.338	1.125	1.086

(de Muynck, W. Ann. 1894, 53. 561.)

$\text{CdI}_2 + \text{Aq}$ containing 10.97% CdI_2 has sp. gr. $20^\circ/20^\circ = 1.0982$.

$\text{CdI}_2 + \text{Aq}$ containing 16.53% CdI_2 has sp. gr. $20^\circ/20^\circ = 1.1562$.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 282.)

Sp. gr. of $\text{CdI}_2 + \text{Aq}$ at 20° .

Normality of $\text{CdI}_2 + \text{Aq}$	% CdI_2	Sp. gr.
1.924	44.53	1.5807
0.951	27.07	1.2837
0.447	14.40	1.1355
0.211	7.26	1.0630

(Forchheimer, Z. phys. Ch. 1900, 34. 29.)

$\text{CdI}_2 + \text{Aq}$ containing 1 pt. CdI_2 in 2.2691 pts. H_2O at 17° has sp. gr. = 1.3341. (Hit-
torf, Z. phys. Ch. 1902, 39. 628.)

Sol. in sat. $\text{HI} + \text{Aq}$.

Sol. in warm $\text{NH}_4\text{OH} + \text{Aq}$.

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, 25. 217.)

Difficultly sol. in POCl_3 . (Walden, Z. anorg. 1900, 25. 212.)

Nearly insol. in AsBr_3 . (Walden, Z. anorg. 1902, 29. 374.)

Sol. in SO_2Cl_2 . (Walden, Z. anorg. 1900, 25. 215.)

Sol. in 15 pts. alcohol. (Vogel, N. Rep. Pharm. 12. 393.)

Sol. in 0.98 pt. abs. alcohol. (Eder, Dingl. 221. 89.)

Sp. gr. of $\text{CdI}_2 + \text{alcohol}$.

% CdI_2 Sp. gr. $20^\circ/20^\circ$

0 0.7949

7.28 0.8470

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 284.)

Sol. in 5.2 mols. methyl, 7 mols. ethyl, and 9.8 mols. propyl alcohol at 20° . (Timofejew, C. R. 112. 1224.)

Sol. in 3.6 pts. ether. (Eder, l. c.)

Sol. in 2.0 pts. alcohol-ether (1 : 1). (Eder, l. c.)

Very sl. sol. in anhydrous abs. ether. (Hampe, Ch. Z. 1887, 11. 847.)

100 g. of sat. solution in abs. ether contain 0.143 g. CdI_2 at 12° . (Tyrer, Proc. Chem. Soc. 1911, 27. 142.)

Solubility in ether + Aq at 12° .

% H_2O in ether	% CdI_2	% H_2O in ether	% CdI_2	% H_2O in ether	% CdI_2
0.0	0.143	0.50	3.36	1.00	7.30
0.10	0.78	0.70	4.77	1.10	8.27
0.30	2.07	0.90	6.46	1.14	8.68

(Tyrer, Proc. Chem. Soc. 27. 142.)

Solubility in benzene at $16^\circ = 0.01\%$

" $35^\circ = 0.02\%$

Solubility in ethyl ether at $0^\circ = 0.03\%$

" $15.5^\circ = 0.04\%$

" $20.3^\circ = 0.05\%$

(Linebarger, Am. J. Sci. 1895, (3) 49. 52.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

1 g. CdI_2 is sol. in 4 g. acetone at 18° . Sp. gr. of sat. solution $18^\circ/4^\circ = 0.994$. (Nau-
mann, B. 1904, 37. 4338.)

Sp. gr. of $\text{CdI}_2 + \text{acetone}$.

% CdI_2	Sp. gr. $20^\circ/20^\circ$.
0	0.7998
12.02	0.8929

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 284.)

Sol. in chinolin. (Beckmann and Gabel, Z. anorg. 1906, **51**. 236.)

100 g. benzonitrile dissolve 1.6295 g. CdI_2 at 18° . (Naumann, B. 1914, **47**. 1370.)

Insol. in methylene iodide. (Retgers, Z. anorg. **3**. 343.)

Sl. sol. in ethylamine. (Shinn, J. phys. Chem. 1907, **11**. 538.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1849, **6**. 257.)

Solubility in methyl acetate = 0.7–1.5%; 2.1% at bpt. (Schröder and Steiner, J. pr. 1909, (2) **79**. 49.)

Sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

1 pt. is sol. in 54.3 pts. ethyl acetate at 18° . The sat. solution has $D_{18^\circ/4^\circ} = 0.9145$. (Naumann, B. 1910, **43**. 318.)

Insol. in mustard oil. (Mathews, J. phys. Chem. 1905, **9**. 647.)

Mol. weight determined in piperidine, pyridine, methyl and ethyl sulphide. (Werner, Z. anorg. 1897, **15**. 17.)

Cadmium hydrogen iodide, $\text{CdI}_2 \cdot \text{HI} + 3\text{H}_2\text{O}$.

Decomp. in air. (Dobroserdow, C. C. **1900**, II. 527.)

Cadmium caesium iodide, $\text{CdI}_2 \cdot \text{CsI} + \text{H}_2\text{O}$.

Sol. in H_2O without decomp. (Wells and Walden, Z. anorg. **5**. 271.)

$\text{CdI}_2 \cdot 2\text{CsI}$. As above.

$\text{CdI}_2 \cdot 3\text{CsI}$. Decomp. by H_2O into the above salt.

Cadmium hydrazine iodide, $\text{CdI}_2 \cdot 2\text{N}_2\text{H}_4\text{HI}$.

Sol. in H_2O . (Ferratini, C. A. **1912**. 1612.)

Cadmium mercuric iodide.

Very sol. in H_2O . (Berthemot, J. Pharm. **14**. 613.)

$\text{CdI}_2 \cdot 3\text{HgI}_2$. Sol. in H_2O . Can be recrystallized in alcohol. (Clarke and Kebler, Am. Ch. J. **5**. 235.)

Cadmium potassium iodide, $\text{CdI}_2 \cdot \text{KI} + \text{H}_2\text{O}$.

Sol. in 0.94 pt. H_2O at 15° . (Eder, Dingl. **221**. 89.)

$\text{CdI}_2 \cdot 2\text{KI} + 2\text{H}_2\text{O}$. Deliquescent. Extremely sol. in H_2O . Sol. at 15° in 0.73 pt. H_2O . Sl. sol. in alcohol and wood spirit, but less than CdI_2 . (Croft.)

Sol. at 15° in 1.4 pts. absolute alcohol, 24.5 pts. ether (0.729 sp. gr.), and 4.5 pts. alcohol-ether (1 : 1). (Eder, l. c.)

Sp. gr. of $\text{K}_2\text{CdI}_4 + \text{Aq}$ at 18° .

% K_2CdI_4	1	5	10	15	20
Sp. gr.	1.0065	1.0384	1.0808	1.1269	1.1770

% K_2CdI_4	25	30	35	40	45
Sp. gr.	1.2313	1.2890	1.3557	1.4282	1.5065

(Grotrian, W. Ann. 1883, **18**. 193.)

Sp. gr. of $\text{K}_2\text{CdI}_4 + \text{Aq}$.

% K_2CdI_4	t°	Sp. gr. at t°	Sp. gr. at 18°
0.0328	18		0.99895
0.0596	18		0.99921
0.0804	18		0.99938
0.100	17.12	0.99962	0.99945
	21.82	0.99872	
0.250	18		1.0007
0.500	18		1.0027
1.003	17.32	1.0068	1.0067
	20.63	1.0061	

(Wershofen, Z. phys. Ch. 1890, **5**. 493.)

Sol. in ethyl acetate. (Naumann, B. 1904, **37**. 3601.)

Cadmium sodium iodide, $\text{CdI}_2 \cdot 2\text{NaI} + 6\text{H}_2\text{O}$.

Deliquescent. (Croft.)

Sol. at 15° in 0.63 pt. H_2O , 0.86 pt. abs. alcohol, and 10.1 pts. ether (sp. gr. 0.729). (Eder, Dingl. **221**. 89.)

Cadmium strontium iodide, $\text{CdI}_2 \cdot \text{SrI}_2 + 8\text{H}_2\text{O}$.

Deliquesces in moist, effloresces in dry air; sol. in H_2O . (Croft.)

Cadmium iodide ammonia, $\text{CdI}_2 \cdot 2\text{NH}_3$.

Decomp. by H_2O . (Rammelsberg.)

$\text{CdI}_2 \cdot 4\text{NH}_3$. (Dawson and McCrae, Chem. Soc. 1900, **77**. 1246.)

$\text{CdI}_2 \cdot 6\text{NH}_3$. Decomp. by H_2O ; sol. in warm, less sol. in cold $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg.)

Cadmium iodide hydrazine, $\text{CdI}_2 \cdot 2\text{N}_2\text{H}_4$.

Easily sol. in warm $\text{NH}_4\text{OH} + \text{Aq}$. (Franzen, Z. anorg. 1908, **60**. 281.)

Cadmium iodide hydroxylamine, $\text{CdI}_2 \cdot 3\text{NH}_2\text{OH}$.

Sol. in H_2O and alcohol. Insol. in ether. (Adams, Am. Ch. J. 1902, **28**. 218.)

Cadmium iodide selenide, $\text{CdI}_2 \cdot 3\text{CdSe}$.

Easily decomp. (Fonzes-Diacon, C. R. 1900, **131**. 897.)

Cadmium iodosulphide, $\text{CdI}_2 \cdot 2\text{CdS}$.

Ppt. (Naumann, B. 1904, **37**. 4338.)

Cadmium suboxide, Cd_2O .

Decomp. by H_2O , acids and $\text{NH}_4\text{OH} + \text{Aq}$. (Tanatar, Z. anorg. 1901, **27**. 433.)

Cd_2O . Properties as cadmous hydroxide. (Morse and Jones.)

Cadmium oxide, CdO .

Insol. in H_2O . Sol. in acids. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Easily sol. in $\text{NH}_4\text{Cl} + \text{Aq}$, less in $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett, **1837**.)

in KOH, NaOH, K_2CO_3 , and Na_2CO_3 .

Cadmium hydroxide.

solubility in (calcium succrate+sugar)+

solution containing 418.6 g. sugar and H_2O dissolves 0.22 g. CdO .

solution containing 174.4 g. sugar and H_2O dissolves 0.48 g. CdO .

Wodenbender, J. B. 1865. 600.)

insol. in acetone. (Naumann, B. 1904, 37.)

insol. in methyl acetate. (Naumann, B. 3790.)

insol. in ethyl acetate. (Naumann, B. 3601.)

peroxide, Cd_2O_2 or $Cd_2O_3(?)$.

$Cd(OH)_2$. (Kouriloff, A. ch. (6) 23.)

stable towards H_2O . Insol. in NH_4OH (Faas, B. 1884, 17. 2253.)

$Cd(OH)_2$. Ppt. Insol. in NaOH + kmann, C. C. 1905, I. 1629.)

$CdO + 3H_2O$. Ppt. (Teletow, C. A.)

oxybromide, CdO , $CdBr_2 + H_2O$.

p. by H_2O . (Tassily, C. R. 1897,)

Stable in dry air; insol. in H_2O . (C. R. 1897, 124. 1022.)

Slowly decomp. by H_2O . (Tassily, 1897, 124. 1022.)

(Mailhe, C. R. 1901, 132. 1561.)

oxychloride, $CdCl_2$, $CdO + H_2O$.

insol. in hot H_2O . (Habermann, M. Ch.)

(Mailhe, Bull. Soc. 1901, (3) 25.)

$CdCl_2$. Insol. in H_2O , but slowly thereby. (Canzoneri, Gazz. ch. it. (2) 486.)

oxyiodide, CdO , $CdI_2 + H_2O$.

p. by H_2O . (Tassily, C. R. 1897,)

Stable in dry air; insol. in H_2O . (C. R. 1897, 124. 1022.)

phosphide, Cd_3P_2 .

$HCl + Aq$ with evolution of PH_3 . (Ber.)

Sol. in conc. $HCl + Aq$. (Emmerling, 12. 152.)

decomp. by acids. (Kulisch, A. 231.)

Decomp. by boiling conc. $HCl + Aq$. (C. R. 76. 283.)

selenide, $CdSe$.

$HCl + Aq$. (Uelsmann, A. 116. 122.)
decomp. by acids. (Fonzes-Diacon, 00, 131. 897.)

Cadmium sulphide, CdS .

Insol. in H_2O .

Solubility in H_2O at $16-18^\circ = 6.6 \times 10^{-4}$ mols. per l. (Biltz, Z. phys. Ch. 1907, 58. 291.)

1 l. H_2O dissolves 9.00×10^{-4} mols. CdS (artificial greenockite) at 18° .

1 l. H_2O dissolves 8.86×10^{-4} mols. pptd. CdS at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

Difficultly sol. in hot dil. $HCl + Aq$. Easily sol. in cold conc. $HCl + Aq$. (Stromeyer.) Sol. in $HNO_3 + Aq$ (Meissner), and boiling dil. $H_2SO_4 + Aq$ (1 : 6). (A. W. Hoffmann, A. 115. 286.) Very sl. sol. in $NH_4OH + Aq$. (Wackenroder, Repert. 46. 226.) Insol. in KOH, or $(NH_4)_2S + Aq$. Appreciably sol. in an acid solution of NH_4Cl . (Baxter and Hines, Z. anorg. 1905, 44. 160.)

Much more sol. in $(NH_4)_2S + Aq$ than usually supposed. (Ditte, C. R. 85. 402.) Solubility increases by warming, and at 68° is twice that at ordinary temperatures. A sat. solution of $(NH_4)_2S$ dissolves about 2 g. CdS to a litre. Alkali sulphides dissolve much less. (Ditte.)

Fresenius (Z. anal. 20. 236) could not confirm the above. According to Fresenius, CdS is not appreciably sol. in $(NH_4)_2S + Aq$.

Insol. in Na_2SO_3 or $KCN + Aq$. (Fresenius.)

Insol. in NH_4Cl or $NH_4NO_3 + Aq$. (Brett.)

Sol. in alkali sulpho-molybdates, -tungstates, -vanadates, -arsenates, -antimonates, -stannates + Aq . (Storch, B. 16. 2015.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Min. Greenockite. Sol. in $HCl + Aq$.

Colloidal.—Solution of 4 g. colloidal C in a litre H_2O remains transparent several days. If it contains 11 g. CdS in a litre, it is completely coagulated in 24 hours. Solutions of salts of the following concentration cause an immediate coagulation in an aqueous solution of CdS containing 3.62 g. in a litre.

KCl	.	.	.	1 : 1615
KBr	.	.	.	1 : 727
KI	.	.	.	1 : 57
KCN	.	.	.	1 : 166
KClO ₃	.	.	.	1 : 1666
KNO ₃	.	.	.	1 : 1000
K ₂ S ₂ O ₈	.	.	.	1 : 5000
K ₂ SO ₄	.	.	.	1 : 833
K ₂ Fe(CN) ₆	.	.	.	1 : 166
K ₄ Fe(CN) ₆	.	.	.	< 1 : 100
K ₂ CrO ₄	.	.	.	1 : 400
K ₂ Cr ₂ O ₇	.	.	.	1 : 3571
NaCl	.	.	.	1 : 2666
Na ₂ S ₂ O ₃	.	.	.	1 : 98
NaHCO ₃	.	.	.	1 : 333
Na ₂ CO ₃	.	.	.	1 : 166
Na ₂ HPO ₄	.	.	.	1 : 202

NaC ₂ H ₃ O ₂	1 : 2451
Na benzoate	1 : 10,000
(NH ₄) ₂ C ₂ O ₄	1 : 588
BaCl ₂	1 : 11,764
Ba(NO ₃) ₂	1 : 8032
BaS ₂ O ₆	1 : 5617
MgSO ₄	1 : 41,666
MnSO ₄	1 : 22,222
CdSO ₄	1 : 250,000
Cd(NO ₃) ₂	1 : 285,714
Pb(ClO ₃) ₂	1 : 209
Pb(C ₂ H ₃ O ₂) ₂	1 : 147,058
Hg(CN) ₂	<1 : 20
Al ₂ (SO ₄) ₃	1 : 232,558
Alum	1 : 192,377
Chrome alum	1 : 42,555
HCl	1 : 4807
H ₂ SO ₄	1 : 8000
HC ₂ H ₃ O ₂	1 : 15
H ₂ C ₂ O ₄	1 : 23,255
Succinic acid	<1 : 100
Tartaric acid	1 : 333

(Prost, Belg. Acad. Bull. (3) 14. 312; J. B. 1887. 537.)

Cadmium pentasulphide, CdS₅.

Insol. in H₂O. (Schiff, A. 115. 74.)

Mixture of CdS and S. (Follenius, Z. anal. 13. 412.)

Cadmium potassium sulphide, K₂Cd₂S₄.

(Milbauer, Z. anorg. 1904, 42. 439.)

Cadmium sodium sulphide, 3CdS, Na₂S.

Decomp. by H₂O. (Schneider, J. pr. (2) 8. 29.)

Cadmium sulphioidide.

See Cadmium iodosulphide.

Cadmium telluride, CdTe.

Not attacked by dil. acids. Attacked in the cold only by HNO₃. (Tibbals, J. Am. Chem. Soc. 1909, 31. 908.)

Cadmic acid.

Potassium cadmate.

Insol. in H₂O, but gradually decomp. when in contact therewith. (Meunier, C. R. 63. 330.)

Cæsium, Cs.

Decomp. H₂O with great violence. (Setterberg, A. 211. 100.)

Very sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

Cæsium acetylde acetylene, Cs₂C₂, C₂H₂.

Insol. in C₆H₆ and in CHCl₃. (Moissan, C. R. 1903, 136. 1218.)

Cæsium amide, CsNH₂.

Decomp. by H₂O. Very sol. in liquid NH₃. (Rengade, C. R. 1905, 140. 1185.)

Cæsium ammonia, CsNH₂.

Sol. in liquid NH₃. (Moissan, C. R. 1903, 136. 1177.)

Cæsium azoimide, CsN₃.

Deliquescent. Stable in aq. solution.

224.2 pts. sol. in 100 pts.	H ₂ O	at 0°
307.4 " " " 100 "	H ₂ O	" 16°
1.0366 " " " 100 "	abs. alcohol	" 16°

Insol. in pure ether. (Curtius, J. pr. 1898, (2) 58. 283.)

Cæsium bromide, CsBr.

Ppt. (Chabrié, C. R. 1901, 132. 679.)

Sat. CsBr + Aq at 25° contains 55.23% CsBr. (Foote, Am. Ch. J. 1907, 37. 125.)

Cæsium tribromide, CsBr₃.

Sol. in H₂O; decomp. by alcohols. (Wells, Sill. Am. J. 143. 17.)

Cæsium pentabromide, CsBr₅.

Very unstable. (Wells and Wheeler, Sill. Am. J. 144. 42.)

Cæsium cobalt bromide, Cs₂CoBr₄.

Decomp. by H₂O. (Campbell, Z. anorg. 1894, 8. 126.)

Decomp. by H₂O and by alcohol. (Campbell, Am. J. Sci. 1894, (3) 48. 418.)

Cs₂CoBr₄. Decomp. by H₂O. (Campbell, Z. anorg. 1894, 8. 126.)

Decomp. by H₂O and by alcohol. (Campbell, Am. J. Sci. 1894, (3) 48. 418.)

Cæsium copper bromide, CsBr, CuBr₂.

Sol. in H₂O without decomp. (Wells and Walden, Z. anorg. 5. 304.)

2 CsBr, CuBr₂. (W. and W.)

Cæsium iridium bromide.

See Bromiridate, cæsium.

Cæsium iron (ferric) bromide, CsFeBr₄.

Sol. in H₂O. (Walden, Z. anorg. 1894, 7. 332.)

Cs₂FeBr₄ + H₂O. (Walden, Z. anorg. 1894, 7. 332.)

Cæsium lead bromide, CsBr, 2PbBr₂.

Nearly stable in aqueous solution. (Walden, Sill. Am. J. 145. 127.)

CsBr, PbBr₂. Decomp. by H₂O. (Walden.)

4CsBr, PbBr₂. As above.

Solubility determinations show that the double salts formed by cæsium and lead bromides at 25° are CsPb₂Br₆, CsPbBr₃ and Cs₂PbBr₆. (Foote, Am. Ch. J. 1907, 37. 125.)

Cæsium magnesium bromide, CsBr, MgBr₂ + 6H₂O.

Sol. in H₂O. (Wheeler and Campbell, Z. anorg. 5. 275.)

Cæsium mercuric bromide, $\text{CsBr}, 2\text{HgBr}_2$.

Not decomp. by H_2O . 100 pts. solution sat. at 16° contain 0.807 pt. $\text{CsBr}, 2\text{HgBr}_2$. Sol. in hot strong alcohol, from which $\text{CsBr}, \text{HgBr}_2$ separates on cooling. (Wells, Sill. Am. J. 144. 221.)

$\text{CsBr}, \text{HgBr}_2$. Decomp. by H_2O into above salt. Sol. in alcohol without decomp. (Wells.)

$2\text{CsBr}, \text{HgBr}_2$. Decomp. by H_2O into $\text{CsBr}, 2\text{HgBr}_2$.

$3\text{CsBr}, \text{HgBr}_2$. As above.

Cæsium molybdenyl bromide, $2\text{CsBr}, \text{MoOBr}_2$.

(Weinland and Knöll, Z. anorg. 1905, 44. 107.)

Cæsium nickel bromide, CsNiBr_2 .

Decomp. by H_2O . (Campbell, Z. anorg. 1894, 8. 126.)

Decomp. by H_2O and by alcohol. (Campbell, Am. J. Sci. 1894, (3) 48. 418.)

Cæsium osmium bromide.

See Bromosmate, cæsium.

Cæsium palladium bromide.

See Bromopalladate, cæsium and bromopalladite, cæsium.

Cæsium platinum bromide.

See Bromoplatinate, cæsium.

Cæsium ruthenium bromide.

See Bromoruthenite, cæsium.

Cæsium selenium bromide.

See Bromoselenate, cæsium.

Cæsium tellurium bromide.

See Bromotellurate, cæsium.

Cæsium thallic bromide, $\text{CsBr}, \text{TlBr}_3$.

Sol. in H_2O with decomp. (Pratt, Z. anorg. 1895, 9. 19.)

By recryst. from H_2O , forms $3\text{CsBr}, 2\text{TlBr}_3$. (Pratt, Am. J. Sci. 1895, (3) 49. 403.)

$3\text{CsBr}, 2\text{TlBr}_3$. Can be recryst. unchanged from H_2O . (Pratt, Am. J. Sci. 1895, (3) 49. 402.)

Cæsium tin (stannic) bromide.

See Bromostannate, cæsium.

Cæsium zinc bromide, $3\text{CsBr}, \text{ZnBr}_2$.

Sol. in H_2O . (Wells and Campbell, Z. anorg. 5. 275.)

$2\text{CsBr}, \text{ZnBr}_2$. As above.

Cæsium bromide columbium oxybromide, $2\text{CsBr}, \text{CbOBr}_2$.

Unstable in moist air. Decomp. by H_2O . (Weinland, B. 1906, 39. 3059.)

Cæsium bromochloride, CsBr_2Cl .

Properties as CsBr_2 . (Wells.)

CsBrCl_2 . As above. (Wells.)

Cæsium mercuric bromochloride, $\text{Cs}_2\text{HgCl}_2\text{Br}_2$.

Decomp. by H_2O finally to HgBr_2 . (Wells, Sill. Am. J. 144. 121.)

$\text{Cs}_2\text{HgCl}_2\text{Br}$. As above.

CsHgClBr_2 . As above.

$\text{CsHg}_2\text{ClBr}_4$. As above.

$\text{CsHg}_3\text{ClBr}_{10}$. As above.

Cæsium bromochloroiodide, CsBrClI .

More sol. in H_2O than in alcohol. Not decomp. at once by ether. (Wells.)

Cæsium bromoiodide, CsBrI_2 .

Decomp. by H_2O . Sol. in alcohol. Decomp. by ether with residue of CsBr . (Wells, Sill. Am. J. 143. 17.)

CsBr_2I . More sol. in H_2O than in alcohol. Not decomp. by ether.

$\text{CsBr}_2\text{I} + \text{Aq}$ sat. at 20° contains about 4.45% CsBr_2I . (Wells.)

Cæsium carbide, Cs_2C_2 .

Decomp. by cold H_2O . (Moissan, C. R. 1903, 136. 1221.)

Cæsium chloride, CsCl .

Very deliquescent; sol. in H_2O and alcohol.

Solubility of CsCl at t° .

t°	Pts. by wt. of CsCl in 100 pts. solution
0.3	61.9
10	63.5
20	64.9
30	66.3
40	67.4

(Hinrichsen, Z. phys. Ch. 1904, 50. 99.)

Solubility of CsCl at t° .

t°	% CsCl	t°	% CsCl
0	61.7	60	69.7
10	63.6	70	70
20	65.1	80	71.4
30	66.4	90	72.2
40	67.5	100	73.0
50	68.0	119.4	74.4

(Berkeley, Trans. Roy. Soc. 1904, 203. A. 208.)

A normal solution of CsCl has sp. gr. at $25^\circ = 1.1076$. (Wagner, Z. phys. Ch. 1890, 5. 36.)

Sp. gr. at $20^\circ/4^\circ$ of a normal solution of $\text{CsCl} = 1.125815$. (Haigh, J. Am. Chem. Soc. 1912, 34. 1151.)

Sp. gr. of CsCl+Aq.			
G. equiv. CsCl per l. at 18°	Sp. gr. at 6°/6°	Sp. gr. at 18°/18°	Sp. gr. at 30°/30°
0.504	1.06556	1.06483	1.06452
1.602	1.12962	1.12825	1.12750
2.007	1.25705	1.25452	1.25307
3.994	1.50514	1.50100	1.49859

(Clausen, W. Ann. 1914, (4) 44. 1071.)

Solubility of CsCl+FeCl₃ in H₂O at 21°.

Substance added		Pts. by weight in 100 pts. of solution	
FeCl ₃ grams	CsCl grams	FeCl ₃	CsCl
0	65	0	65.0
0.6	11.6	0.45	55.18
1.4	10.2	2.1	52.38
2.2	8.8	5.24	51.44
2.0	7.4	7.8	47.70
3.8	6.0	8.93	41.15
4.6	4.6	15.34	25.25
5.4	2.8	21.65	14.96
6.2	1.4	27.96	8.42
35	0.2	48.71	0.94
35	0	83.89	0

(Hinrichsen, Z. phys. Ch. 1904, 50. 96.)

Solubility of CsCl+HgCl₂ in H₂O at 25°.

Solution contains		Solid phase
% CsCl	% HgCl ₂	
65.61	0.00	CsCl
65.78	0.215	CsCl+Cs ₂ HgCl ₃
62.36	0.32	Cs ₂ HgCl ₃
57.01	0.64	"
52.35	1.23	"
51.08	1.44	Cs ₂ HgCl ₃ +Cs ₂ HgCl ₄
49.30	1.49	Cs ₂ HgCl ₄
45.95	1.69	"
45.23	1.73	Cs ₂ HgCl ₄ +CsHgCl ₃
38.63	1.32	CsHgCl ₃
17.03	0.51	"
1.53	0.42	"
0.61	2.64	CsHgCl ₃ +CsHg ₂ Cl ₅
0.49	2.91	CsHg ₂ Cl ₅
0.40	3.78	CsHg ₂ Cl ₅ +CsHg ₃ Cl ₁₁
0.44	4.63	"
0.41	4.68	CsHg ₃ Cl ₁₁
0.25	5.65	"
0.18	7.09	CsHg ₃ Cl ₁₁ +HgCl ₂
0.00	6.90	HgCl ₂

(Foote, Am. Ch. J. 1903, 30. 340.)

Solubility of CsCl+HgCl₂ in acetone

Solution contains		Solid phase
% HgCl ₂	% CsCl	
57.74	0.00	HgCl ₂
57.79	0.13	HgCl ₂ +CsHg ₃ Cl ₁₁
57.74	0.20	CsHg ₃ Cl ₁₁
52.54	0.22	"
49.83	0.32	"
44.32	0.50	CsHg ₃ Cl ₁₁ +CsH
44.46	0.44	"
39.65	0.48	CsHg ₂ Cl ₅
28.48	0.48	"
26.96	0.52	CsHg ₂ Cl ₅ +CsH
27.32	0.61	"
21.50	0.46	CsHgCl ₃
13.08	0.45	"
0.16	0.19	Mixtures of sa
0.17	0.25	"
0.02	0.11	"
0.00	0.032	CsCl

(Foote and Haigh, J. Am. Ch. Soc. 1911, 33. 461.)

Insol. in methyl acetate. (Naumann, Z. anorg. 1909, 42. 3790.)

Solubility in glycol at ord. temp. 10.8%. (de Coninck, Belg. Acad. Bul. 1909, 359.)

Insol. in anhydrous pyridine and pyridine+Aq. Sl. sol. in 95% pyridine and in 93% pyridine+Aq. (Kahlen, Am. Chem. Soc. 1908, 30. 1107.)

Cæsium chromium chloride, 2CsCl, CrCl₃·6H₂O.

Stable in the air. Sol. in H₂O. (Vogel, Z. anorg. 1895, 10. 182.)

2CsCl, CrCl₃+4H₂O; hygroscopic; v. sol. in H₂O. (Wells, l. c.)

Cæsium tetra-aquochromium chloride, CrCl₂(OH₂)₄·Cl, 2CsCl.

Ppt. (Werner, B. 1901, 34. 1602.)

Cæsium cobalt chloride, CsCoCl₃+2H₂O.

Decomp. by H₂O and alcohol. (Clausen, Z. anorg. 1894, 8. 126.)

Cs₂CoCl₄. Decomp. by H₂O and alcohol. (Campbell, Z. anorg. 1894, 8. 126.)

Cs₃CoCl₅. Decomp. by H₂O and alcohol. (Campbell, Z. anorg. 1894, 8. 126.)

Cæsium cuprous chloride, CsCl, Cu₂Cl₂.

Decomp. by H₂O into CuCl₂, CsCl. (Z. anorg. 5. 306.)

3CsCl, Cu₂Cl₂. (Wells.)

6CsCl, Cu₂Cl₂. (Wells.)

Cæsium cupric chloride, 2CsCl, CuCl₂.

Easily sol. in H₂O and dil. HCl. Insol. in conc. HCl+Aq. (Godeffroy, Z. anorg. 8. 9.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, 11. 1014.)

100 g. solution in acetone sat. at 25° contain 0.032 g. CsCl. (Foote and Haigh, J. Am. Chem. Soc. 1911, 33. 461.)

Sol. in small amount H_2O without decomp. (Wells and Dupee, Z. anorg. 5. 300.)

+ $2H_2O$. Efflorescent. (W. and D.)

$3CsCl, 2CuCl_2 + 2H_2O$.

$CsCl, CuCl_2$. Sol. in H_2O without decomp. (W. and D.)

Cesium gold chloride.

See Chloraurate, caesium.

Cesium iridium tetrachloride.

See Chloriridate, caesium.

Cesium iron (ferric) chloride, $CsFeCl_4 + \frac{1}{2}H_2O$.

Sol. in H_2O . Decomp. in the air. (Walden, Z. anorg. 1894, 7. 332.)

$Cs_2FeCl_6 + H_2O$. Sol. in H_2O . (Walden.)

$Cs_3FeCl_8 + H_2O$. Sol. in H_2O . (Walden.)

Cesium lanthanum chloride, $Cs_3LaCl_6 + 4H_2O$.

Very hygroscopic. Easily sol. in H_2O . (R. J. Meyer, Z. anorg. 1914, 86. 273.)

Cesium lead chloride, $CsCl, 2PbCl_2$.

Nearly stable in aqueous solution. (Campbell, Sill. Am. J. 145. 126.)

$CsCl, PbCl_2$. Decomp. by H_2O . (Campbell.)

$4CsCl, PbCl_2$. As above. (Campbell.)

Cesium lead tetrachloride.

See Chloroplumbate, caesium.

Cesium magnesium chloride, $CsCl, MgCl_2 + 6H_2O$.

Sol. in H_2O . (Wells and Campbell, Z. anorg. 5. 275.)

Cesium manganous chloride, $CsCl, MnCl_2 + 2H_2O$.

Not deliquescent; sol. in H_2O . (Saunders, Am. Ch. J. 14. 143.)

$2CsCl, MnCl_2$. (Godeffroy.)

+ $2\frac{1}{2}H_2O$. (Godeffroy.)

+ $3H_2O$. Sol. in H_2O . Conc. $HCl + Aq$ precipitates anhydrous salt from aqueous solution. (Godeffroy, B. 8. 9.)

The only salt which exists contains $2H_2O$. (Saunders, Am. Ch. J. 14. 143.)

Cesium manganic chloride, $2CsCl, MnCl_2$.

Easily decomp. (Meyer and Best, Z. anorg. 1899, 22. 187.)

Cesium mercuric chloride, $CsCl, HgCl_2$.

100 pts. solution sat. at 17° contain 1.406 pts. $CsHgCl_3$. Not decomp. by H_2O . Insol. in absolute alcohol, but sol. on diluting with $\frac{1}{2}$ vol. H_2O . (Wells, Sill. Am. J. 144. 221.)

$2CsCl, HgCl_2$. Easily sol. in H_2O and dil. $HCl + Aq$; insol. in conc. $HCl + Aq$. (Godeffroy.)

$3CsCl, HgCl_2$. Decomp. by H_2O ; on recrystallizing from H_2O , $CsCl, HgCl_2$ is finally formed. (Wells, Sill. Am. J. 144. 221.)

$CsCl, 5HgCl_2$. Decomp. by H_2O . (Wells.)

Solubility determinations show that the only double salts of $CsCl$ and $HgCl_2$ which exist at 25° are Cs_2HgCl_5 , Cs_3HgCl_6 , $CsHgCl_3$, $CsHg_2Cl_4$, $CsHg_3Cl_5$. (Foote, Am. Ch. J. 1903, 30. 340.)

Cesium molybdenum chloride, $Cs_2MoCl_6 + H_2O$.

Sol. in H_2O . Nearly insol. in alcohol and ether. (Chilesotti, C. C. 1903, II. 652.)

Cesium molybdenyl chloride, $CsCl, MoO_2Cl_2 + H_2O$.

Hygroscopic. Decomp. by H_2O . (Weinland and Knöll, Z. anorg. 1905, 44. 93.)

$2CsCl, MoO_2Cl_2$. Hygroscopic. Decomp. by H_2O . (Weinland and Knöll, Z. anorg. 1905, 44. 92.)

$2CsCl, 6MoO_2Cl_2 + 22H_2O$. Very hygroscopic. Decomp. by H_2O . (Weinland and Knöll, Z. anorg. 1905, 44. 94.)

$2CsCl, MoOCl_3$. Only sl. sol. in H_2O . (Nordenskjöld, B. 1901, 34. 1573.)

Cesium neodymium chloride, $Cs_3NdCl_6 + 5H_2O$.

Very hygroscopic. Easily sol. in H_2O . (R. J. Meyer, Z. anorg. 1914, 86. 273.)

Cesium nickel chloride, $2CsCl, NiCl_2$.

As the corresponding Cu salt.

$CsNiCl_3$. Decomp. by H_2O and by alcohol. (Campbell, Am. J. Sci. 1894, (3) 48. 418.)

Cesium palladium dichloride.

See Chloropalladite, caesium.

Cesium palladium tetrachloride.

See Chloropalladate, caesium.

Cesium praseodymium chloride, $Cs_3PrCl_6 + 5H_2O$.

Very hygroscopic. Easily sol. in H_2O . (R. J. Meyer, Z. anorg. 1914, 86. 273.)

Cesium rhodium chloride.

See Chlororhodite, caesium.

Cesium ruthenium chloride.

See Chlororuthenite, caesium and chlororuthenate, caesium.

Cesium oxyruthenium chloride, $Cs_2RuO_2Cl_4$.

Ppt.; decomp. by H_2O ; sol. in cold HCl . (Howe, J. Am. Chem. Soc. 1901, 23. 779.)

Cesium samarium chloride, $Cs_3SmCl_6 + 5H_2O$.

Very hygroscopic. Easily sol. in H_2O . (R. J. Meyer, Z. anorg. 1914, 86. 273.)

Cæsium silver chloride, 2CsCl , AgCl .

Easily decomp. by H_2O . (Wells and Wheeler, *Sill. Am. J.* **144**. 155.)

Cæsium tellurium chloride.

See Chlorotellurate, cæsium.

Cæsium thallic chloride, 2CsCl , TiCl_3 .

By recryst. from H_2O forms 3CsCl , 2TiCl_3 . (Pratt, *Am. J. Sci.* 1895, (3) **49**. 398.)

+ H_2O . Readily sol. in hot H_2O but 3CsCl , 2TiCl_3 cryst. from the solution. (Pratt, *Am. J. Sci.* 1895, (3) **49**. 399.)

3CsCl , 2TiCl_3 . Can be recryst. from H_2O without change. (Pratt, *Am. J. Sci.* 1895, (3) **49**. 401.)

3CsCl , TiCl_3 + $2\text{H}_2\text{O}$. Sol. in 36.4 pts. H_2O at 17° and 3 pts. at 100° . (Godeffroy, *Zeitsch. d. allgem. österr. Apothekerv.* **1880**. No. 9.)

Cæsium tin (stannic) chloride.

See Chlorostannate, cæsium.

Cæsium titanium chloride, TiCl_3 , 2CsCl + H_2O .

Difficultly sol. in H_2O . (Stähler, *B.* 1904, **37**. 4409.)

Cæsium tungsten chloride, $\text{Cs}_3\text{W}_2\text{Cl}_9$.

Nearly insol. in cold H_2O .

Sol. in a hot mixture of equal pts. H_2O and conc. HCl .

Nearly insol. in conc. HCl .

Sol. in very dil. NaOH + Aq.

Nearly insol. in most organic solvents. (Olsson, *B.* 1913, **46**. 574.)

Cæsium uranous chloride, Cs_2UCl_6 .

As K salt. (Aloy, *Bull. Soc.* 1899, (3) **21**. 264.)

Cæsium uranyl chloride, 2CsCl , $(\text{UO}_2)\text{Cl}_2$.

Sol. in H_2O . (Wells, *Z. anorg.* 1895, **10**. 183.)

100 pts. of the solution contain at 29.75° , 56.07 pts. UO_2Cl_2 , 2CsCl . (Rimbach, *B.* 1904, **37**. 468.)

Pptd. from aq. solution by gaseous HCl . (Wells, *Am. J. Sci.* 1894, (3) **50**. 251.)

Cæsium vanadium chloride, Cs_2VdCl_5 + H_2O .

Difficultly sol. in H_2O and alcohol. (Stähler, *B.* 1904, **37**. 4412.)

Cæsium zinc chloride, 3CsCl , ZnCl_2 .

Sol. in H_2O . (Wells and Campbell, *Z. anorg.* **5**. 275.)

2CsCl , ZnCl_2 . Easily sol. in H_2O and dil. HCl + Aq. Insol. in conc. HCl + Aq. (Godeffroy.)

Cæsium chloride chromic oxychloride, 2CsCl , CrOCl_2 .

Decomp. in the air.

Sol. in conc. HCl without decomp. (Weinland, *B.* 1906, **39**. 4045.)

Cæsium chloride columbium oxychloride, 2CsCl , CbOCl_2 .

Decomp. by H_2O . (Weinland, *B.* 1906, **39**. 3057.)

Cæsium chloriodide, CsCl_2I .

Properties as CsBrClI . (Wells.)

CsCl_2I . Sl. sol. in H_2O , from which it can be recrystallized without decomp. (Wells and Wheeler.)

Cæsium mercuric chloriodide, $\text{Cs}_2\text{HgCl}_2\text{I}_2$.

Decomp. instantly by H_2O to HgI_2 . (Wells.)

Cæsium fluoride, CsF .

Ppt. (Chabrié, *C. R.* 1901, **132**. 680.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. 100 g. H_2O dissolve 366.5 g CsF at 15° . (de Forcrand, *C. R.* 1911, **152**. 1210.)

Cæsium hydrogen fluoride, CsHF_2 .

Ppt. (Chabrié, *C. R.* 1901, **132**. 680.)

Cæsium tantalum fluoride.

See Fluotantalate, cæsium.

Cæsium tellurium fluoride, CsF , TeF_4 .

Decomp. by H_2O . (Wells, *Am. J. Sci.* 1901, (4) **12**. 190.)

Cæsium titanium fluoride.

See Fluotitanate, cæsium.

Cæsium zirconium fluoride.

See Fluozirconate, cæsium.

Cæsium hydride, CsH .

Decomp. by H_2O with evolution of H_2 . (Moissan, *C. R.* 1903, **136**. 589.)

Cæsium hydroxide, CsOH .

Very deliquescent, and sol. in H_2O . Sol. in alcohol.

79.41% CsOH is contained in a sat. aq. solution at 15° . (de Forcrand, *C. R.* 1909, **149**. 1344.)

75.08% CsOH is contained in sat. aq. solution at 30° . (Schreinemakers, *C. C.* **1900**, I. 11.)

Cæsium iodide, CsI .

Sol. in H_2O .

100 pts. H_2O dissolve 44 pts. CsI at 0° ; 66.3 pts. at 14.5° ; 160 pts. at 61° .

Sp. gr. of CsI + Aq sat. at 14° = 1.393. (Betekoff, *Bull. Soc. Petersb.* (4) **2**. 197.)

a periodide.

Solubility determinations show that CsI , CsI_2 are the only periodides of caesium existing between -4° and $+73^\circ$. (Foote, *Am. J.* 1903, **29**. 203.)

a triiodide, CsI_3 .

mp. sat. $\text{CsI} + \text{Aq}$ dissolves 0.0097 g. and sp. gr. of solution is 1.154. Only soluble by solution in H_2O . Much more soluble in alcohol than in H_2O . Not immediately soluble by ether. (Wells, *Sill. Am. J.* **143**.)

a penta iodide, CsI_5 .

a cobalt iodide, Cs_2CoI_4 .

mp. by H_2O . (Campbell, *Z. anorg.* **12**.)
Deliquescent; decomp. by H_2O and by H_2O_2 . (Campbell, *Am. J. Sci.* 1894, (3) **1**.)

a lead iodide, CsPbI_2 .

sol. in hot $\text{CsI} + \text{Aq}$. (Wheeler, *Sill. Am.* **129**.)

a mercuric iodide, CsI , 2HgI_2 .

mp. by H_2O finally into HgI_2 . (Wells, *Am. J.* **144**. 221.)

3HgI_2 . Decomp. by H_2O finally into

HgI_2 . As above.

HgI_2 . Decomp. by H_2O ; insol. in

HgI_2 . As above.

a silver iodide, CsI , AgI .

field, *Z. anorg.* **1**. 100.)

2AgI . More sol. in hot than in cold. (Marsh, *Chem. Soc.* 1913, **103**.)

a tellurium iodide.

iodotellurate, caesium.

a thallic iodide, CsI , TlI_3 .

mp. by H_2O . (Pratt, *Am. J. Sci.* **1903**, **49**. 403.)

a zinc iodide, 3CsI , ZnI_2 .

sol. in H_2O . (Wells and Campbell, *Z. anorg.* **12**. 275.)

ZnI_2 . As above.

a oxide, Cs_2O .

abs H_2O and CO_2 from the air. Decomp. by H_2O and by liquid NH_3 . (Rengade, *C. R.* 1906, **143**. 593.)

a dioxide, Cs_2O_2 .

mp. by H_2O . (Rengade, *C. R.* 1905, **140**. 1537.)

Caesium trioxide, Cs_2O_3 .

Decomp. by H_2O . (Rengade, *C. R.* 1905, **140**. 1537.)

Caesium tetroxide, Cs_2O_4 .

Decomp. by H_2O . (Rengade, *C. R.* 1905, **140**. 1538.)

Caesium sulphide, $\text{Cs}_2\text{S} + 4\text{H}_2\text{O}$.

Deliquescent; very sol. in H_2O . (Biltz, *Z. anorg.* 1906, **48**. 300.)

Caesium disulphide, Cs_2S_2 .

Anhydrous. Sol. in H_2O . Hydroscopic. (Biltz, *Z. anorg.* 1906, **50**. 72.)

$+ \text{H}_2\text{O}$. From $\text{Cs}_2\text{S}_2 + \text{Aq}$. Hydroscopic. (Biltz, *Z. anorg.* 1906, **50**. 72.)

Caesium trisulphide, Cs_2S_3 .

Anhydrous. Sol. in H_2O . Not hydroscopic. (Biltz, *Z. anorg.* 1906, **50**. 75.)

$+ \text{H}_2\text{O}$. From $\text{Cs}_2\text{S}_3 + \text{Aq}$. (Biltz, *Z. anorg.* 1906, **50**. 76.)

Caesium tetrasulphide, Cs_2S_4 .

Sol. in H_2O . Insol. in abs. alcohol. (Biltz, *Z. anorg.* 1906, **48**. 305.)

Caesium pentasulphide, Cs_2S_5 .

Mpt. 2° . Not hydroscopic. Very sol. in cold 70% alcohol. (Biltz, *B.* 1905, **38**. 129.)

Caesium hydrogen sulphide, CsHS .

Deliquescent; very sol. in H_2O . (Biltz, *Z. anorg.* 1906, **48**. 300.)

Caesium copper tetrasulphide, CsCuS_4 .

Sl. sol. in cold H_2O .

Decomp. by conc. and dil. HCl , H_2SO_4 and HNO_3 .

Sl. sol. in alcohol. (Biltz, *B.* 1907, **40**. 978.)

Calcium, Ca .

Decomp. H_2O violently. Slowly attacked by cold H_2SO_4 . Dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ or $\text{HCl} + \text{Aq}$ attack violently and dissolve. Dil. $\text{HNO}_3 + \text{Aq}$ oxidizes, but fuming HNO_3 scarcely attacks even on boiling. (Bunsen and Matthiessen.) Not attacked by anhydrous alcohol. (Lies-Bodart and Jobin, *A. ch.* (3) **54**. 364.)

Pure Ca is only very slowly decomp. by H_2O at ordinary temp.; sol. in HCl , HNO_3 , H_2SO_4 . (Moissan, *C. R.* 1898, **129**. 589.)

Insol. in liquid NH_3 . (Gore, *Am. Ch. J.* 1898, **20**. 827.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0334 g. Ca in 6 days. (Gates, *J. phys. Chem.* 1911, **15**. 143.)

Calcium amalgam, Ca_2Hg_4 .

Decomp. H_2O readily. (Féré, *C. R.* 1898, **127**. 619.)

CaHg_4 . Rapidly decomp. in moist air. (Schürger, *Z. anorg.* 1900, **25**. 425.)

Calcium amide, $\text{Ca}(\text{NH}_2)_2$.

(Moissan, A. ch. 1899, (7) 18. 326.)

Calcium ammonia, $\text{Ca}, 4\text{NH}_3$.Decomp. at ordinary temp.; takes fire in contact with the air; sl. sol. in liquid NH_3 . (Moissan, C. R. 1898, 127. 691.) $\text{Ca}, 6\text{NH}_3$. (Kraus, J. Am. Chem. Soc. 1908, 30. 665.)**Calcium arsenide, Ca_3As_2 .**Decomp. by cold H_2O ; insol. in cold fuming HNO_3 ; very sol. in hot HNO_3 . (Lebeau, C. R. 1899, 128. 98.)**Calcium azoimide, $\text{Ca}(\text{N}_3)_2$.**

Hydroscopic; explosive.

38.1	pts. sol. in 100 pts.	H_2O	at 0°
45.0	" " " 100 "	H_2O	" 15.2
0.211	" " " 100. "	abs. alcohol	" 16.

Sol. in H_2O ; decomp. when heated and on standing in the air. (Dennis, Z. anorg. 1898, 17. 21.)

Insol. in pure ether. (Curtius, J. pr. 1898, (2) 58. 286.)

Calcium boride, CaB_4 .Not decomp. by H_2O at 250°; sol. in fused oxidizing agents.Insol. in aq. acids; sl. sol. in conc. H_2SO_4 ; sol. in dil. or conc. HNO_3 . (Moissan, C. R. 1897, 125. 631-32.)**Calcium bromide, CaBr_2 .**

Very deliquescent. 100 pts. H_2O dissolve—

at 0°	20°	40°	60°	105°
125	143	213	278	312 pts. CaBr_2 .

(Kremers, Pogg. 103. 65.)

Sat. $\text{CaBr}_2 + \text{Aq}$ contains at:

—22°	—22°	—14°	—7°	—5°
50.5	50.2	52.5	52.6	52.6% CaBr_2

+8°	9°	11°	20°	50°
53.1	55.1	55.7	57.1	62.6% CaBr_2

(Étard, A. ch. 1894, (7) 2. 540.)

Sp. gr. of $\text{CaBr}_2 + \text{Aq}$ at 19.5° containing:

5	10	15	20	25 % CaBr_2 ,
1.044	1.089	1.139	1.194	1.252

30	35	40	45	50 % CaBr_2 .
1.315	1.385	1.461	1.549	1.641

(Kremers, Pogg. 99. 444, calculated by Gerlach, Z. anal. 8. 285.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Very sol. in alcohol. (Henry.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4328.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

+4 H_2O . (Kuznetzov, C. A. 1911. 842.)
+6 H_2O .**Calcium manganous bromide, $\text{CaMnBr}_4 + 4\text{H}_2\text{O}$.**

Sl. hydroscopic. Unstable. (Ephraim, Z. anorg. 1910, 67. 377.)

Calcium mercuric bromide.Decomp. by H_2O . (v. Bonsdorff.)**Calcium molybdenyl bromide, $\text{CaBr}_2, 2\text{MoOBr}_3 + 7\text{H}_2\text{O}$.**

(Weinland and Knöll, Z. anorg. 1905, 44. 112.)

Calcium stannic bromide.

See Bromostannate, calcium.

Calcium bromide ammonia, $\text{CaBr}_2, 6\text{NH}_3$.Sol. in H_2O . (Rammelsberg, Pogg. 55. 239.)**Calcium bromide hydrazine, $\text{CaBr}_2, 3\text{N}_2\text{H}_4$.**Easily sol. in H_2O . (Franzen, Z. anorg. 1908, 60. 288.)**Calcium bromofluoride, $\text{CaBr}_2, \text{CaF}_2$.**Decomp. by H_2O . (Defacqz, A. ch. 1904, (8) 1. 357.)**Calcium carbide, CaC_2 .**Sp. gr. 2.22 at 18°. Insol. in fuming HNO_3 and conc. H_2SO_4 , but readily decomp. by dil. acids and H_2O . (Moissan, Bull. Soc. 1894, (3) 11. 1005.)Insol. in HCl in the cold, but decomp. at red heat. Strong min. acids do not act in the cold; sol. in glacial acetic in the cold; sol. in fused alkali. (Venable, J. Am. Chem. Soc. 1895, 17. 307-310.)**Calcium chloride, CaCl_2 .**Very deliquescent. Very sol. in H_2O with evolution of heat.Anhydrous CaCl_2 is sol. in 1.459 pts. H_2O . (Gerlach.)Anhydrous CaCl_2 is sol. in 1.58 pts. H_2O at 10.3°. (Kremers, Pogg. 103. 65.)Anhydrous CaCl_2 is sol. in 1.35 pts. H_2O at 20°; 0.83 pt. H_2O at 40°; 0.72 pt. H_2O at 60°. $\text{CaCl}_2 + 6\text{H}_2\text{O}$ is sol. in 0.5 pt. H_2O at 0°, and 2.05 pt. at 16°. (Gmelin.) CaCl_2 is sol. in 1.5 pts. cold, and 0.8 pt. boiling H_2O . (Fourcroy.) $\text{CaCl}_2 + \text{Aq}$ sat. in the cold contains 40.7% CaCl_2 . (Fourcroy.) $\text{CaCl}_2 + \text{Aq}$ sat. at 12.5° contains 53.8% CaCl_2 . (Hassensfratz.)100 pts. H_2O dissolve 165.7 pts. $\text{CaCl}_2 + 6\text{H}_2\text{O}$ at 0°; 7141 pts. at 40°. (Tilden, Chem. Soc. 45. 409.)100 pts. H_2O dissolve 60.3 pts. CaCl_2 from $\text{CaCl}_2 + 6\text{H}_2\text{O}$ at 0°, and solution has sp. gr. = 1.367. (Engel, Bull. Soc. (2) 47. 318.)

Solubility of $\text{CaCl}_2 + 6\text{H}_2\text{O}$ in H_2O at t° .

t°	Sat. solution contains % CaCl_2	Sat. solution contains % $\text{CaCl}_2 + 6\text{H}_2\text{O}$
-22	32.24	63.61
0	36.91	72.82
+ 7.39	38.77	76.49
13.86	41.03	80.95
19.35	42.50	83.85
23.46	44.15	87.11
24.47	45.33	89.44
27.71	46.30	91.35
29.53	50.67	99.97

(Hammerl, W.A.B. 72, 2. 287.)

Solubility in 100 pts. H_2O at t° .

t°	Pts. CaCl_2	t°	Pts. CaCl_2
0	59.39	13.86	69.49
5	64.83	19.35	73.91
7.88	66.20	21.89	79.77

(Hammerl, calculated by Bakhuis Roozeboom, R. t. c. 8. 5.)

Solubility in 100 pts. H_2O at t° .

t°	Pts. CaCl_2	t°	Pts. CaCl_2	t°	Pts. CaCl_2
0	49.6	19	72	38	108
1	50	20	74	39	109
2	51	21	75	40	110
3	52	22	77	41	111
4	53	23	79	42	112
5	54	24	80	43	113
6	55	25	82	44	114
7	56	26	84	45	115
8	57	27	87	46	116
9	58	28	89	47	117
10	60	29	91	48	118
11	61	30	93	49	119
12	62	31	96	50	120
13	63	32	98	51	121
14	65	33	100	52	122
15	66	34	103	53	123
16	68	35	104	54	124
17	69	36	105	55	125
18	71	37	107	56	126
57	127	72	137	87	145
58	128	73	138	88	146
59	129	74	138	89	147
60	129	75	139	90	147
61	130	76	139	91	148
62	131	77	140	92	149
63	131	78	141	93	150
64	132	79	141	94	150
65	133	80	142	95	151
66	133	81	142	96	152
67	134	82	143	97	152
68	135	83	143	98	153
69	135	84	144	99	154
70	136	85	144	179.5	325
71	136	86	145

(Mulder, Scheik. Verhandel. 1864. 107.)

If solubility S = pts. anhydrous CaCl_2 in 100 pts. solution, $S = 32 + 0.2148t$ from -18° to $+6^\circ$; $S = 54.5 + 0.0755t$ from 50° to 120° . (Étard, C. R. 98. 1432.)

According to Bakhuis Roozeboom, the solubility of CaCl_2 varies according to the hydrate employed, and the following data were obtained as the result of very exact experiments.

Solubility of $\text{CaCl}_2 + 6\text{H}_2\text{O}$ in 100 pts. H_2O at t° .

t°	Pts. CaCl_2	t°	Pts. CaCl_2	t°	Pts. CaCl_2
20.4	75.1	28.0	88.8	29.5	96.07
25.05	81.67	28.9	92.05	30.2	102.7

There are two modifications of $\text{CaCl}_2 + 4\text{H}_2\text{O}$, α and β .

Solubility of $\text{CaCl}_2 + 4\text{H}_2\text{O}\beta$ in 100 pts. H_2O at t° .

t°	Pts. CaCl_2	t°	Pts. CaCl_2
18.4	103.3	35.0	122.74
25.0	108.8	38.4	127.50
30.0	114.1

Solubility of $\text{CaCl}_2 + 4\text{H}_2\text{O}\alpha$ in 100 pts. H_2O at t° .

t°	Pts. CaCl_2	t°	Pts. CaCl_2
22.0	92.67	35.95	107.21
24.7	95.59	40.00	115.3
29.8	100.6	45.00	129.9

Solubility of $\text{CaCl}_2 + 2\text{H}_2\text{O}$ in 100 pts. H_2O at t° .

t°	Pts. CaCl_2	t°	Pts. CaCl_2	t°	Pts. CaCl_2
40	128.1	95.8	156.5	139	191.0
45	129.9	115	169.5	155	214.3
50	132.3	124	176.0	165	236.2
59.5	136.5	137	187.6	174	275.7
80.5	145.3

Solubility of $\text{CaCl}_2 + \text{H}_2\text{O}$ in 100 pts. H_2O at t° .

t°	Pts. CaCl_2
191	306
235	331

(Bakhuis Roozeboom, R. t. c. 8.1.)

Sp. gr. of CaCl ₂ + Aq.					
% CaCl ₂	Sp. gr.	% CaCl ₂	Sp. gr.	% CaCl ₂	Sp. gr.
3.95	1.03	20.85	1.18	34.57	1.33
7.66	1.06	23.93	1.21	36.49	1.36
11.23	1.09	26.86	1.24	38.31	1.39
14.42	1.12	29.67	1.27	40.43	1.42
17.60	1.15	32.35	1.30	41.91	1.45

(Richter.)

Sp. gr. of CaCl₂ + Aq at 19.5° containing pts. CaCl₂ to 100 pts. H₂O.

Pts. CaCl ₂	Sp. gr.	Pts. CaCl ₂	Sp. gr.
6.97	1.0545	36.33	1.2469
12.58	1.0954	50.67	1.3234
23.33	1.1681	62.90	1.3806

(Kremers, Pogg. 99. 444.)

Sp. gr. of CaCl₂ + Aq. G = sp. gr. at 15° if % is CaCl₂, according to Gerlach; S = sp. gr. at 18.3° if % is CaCl₂ + 6H₂O, according to Schiff.

%	G	S	%	G	S
1	1.00852	1.0039	36	1.35610	1.1575
2	1.01704	1.0079	37	1.36790	1.1622
3	1.02555	1.0119	38	1.37970	1.1671
4	1.03407	1.0159	39	1.39150	1.1719
5	1.04259	1.0200	40	1.40330	1.1768
6	1.05146	1.0241	41	...	1.1816
7	1.06033	1.0282	42	...	1.1865
8	1.06921	1.0323	43	...	1.1914
9	1.07808	1.0365	44	...	1.1963
10	1.08695	1.0407	45	...	1.2012
11	1.09628	1.0449	46	...	1.2062
12	1.00561	1.0491	47	...	1.2112
13	1.10494	1.0534	48	...	1.2162
14	1.12427	1.0577	49	...	1.2212
15	1.13360	1.0619	50	...	1.2262
16	1.14332	1.0663	51	...	1.2312
17	1.15305	1.0706	52	...	1.2363
18	1.16277	1.0750	53	...	1.2414
19	1.17250	1.0794	54	...	1.2465
20	1.18222	1.0838	55	...	1.2516
21	1.19251	1.0882	56	...	1.2567
22	1.20279	1.0927	57	...	1.2618
23	1.21308	1.0972	58	...	1.2669
24	1.22336	1.1017	59	...	1.2721
25	1.23365	1.1062	60	...	1.2773
26	1.24450	1.1107	61	...	1.2825
27	1.25535	1.1153	62	...	1.2877
28	1.26619	1.1199	63	...	1.2929
29	1.27704	1.1246	64	...	1.2981
30	1.28789	1.1292	65	...	1.3034
31	1.29917	1.1339	66	...	1.3087
32	1.31045	1.1386	67	...	1.3140
33	1.32174	1.1433	68	...	1.3193
34	1.33602	1.1480	69	...	1.3246
35	1.34430	1.1527	70	...	1.3300

(Calculated by Gerlach, Z. anal. 8, 283.)

Sp. gr. of CaCl₂ + Aq : a = no. of half molecule in grammes dissolved in 1000 g. H₂O; b = sp. gr. at 24.3° when a = CaCl₂ 6H₂O (½ mol. = 109.5 g.); c = sp. gr. 24.3° when a = CaCl₂ (½ mol. = 55.5 g)

a	b	c	a	b	c
1	1.041	1.043	7	1.198	1.25
2	1.076	1.084	8	1.214	...
3	1.106	1.122	9	1.229	...
4	1.133	1.159	10	1.242	...
5	1.157	1.193	11	1.255	...
6	1.179	1.227

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of CaCl₂ + Aq at 18°.

% CaCl ₂	Sp. gr.	% CaCl ₂	Sp. gr.
5	1.0409	25	1.2305
10	1.0852	30	1.2841
15	1.1311	35	1.3420
20	1.1794

(Kohlrausch, W. Ann. 1879. 1.)

CaCl₂ + Aq sat. at 0° has sp. gr. = 1.3 (Engel, Bull. Soc. 1887, (2) 47. 318.)

Sp. gr. of CaCl₂ + Aq at 9.5°C.

Mass of salt per unit mass of solution	Density of solution (g. per cc.)
0.00191	1.00168
0.00381	1.00317
0.00570	1.00465
0.00759	1.00615
0.00947	1.00765
0.01320	1.01050

(McGregor, C. N. 1887, 55. 6.)

Sp. gr. of CaCl₂ + Aq at 25°.

Concentration of CaCl ₂ + Aq.	Sp. gr.
1-normal	1.0446
1/2- "	1.0218
1/4- "	1.0105
1/8- "	1.0050

(Wagner, Z. phys. Ch. 1890, 5. 36.)

Sp. gr. at 16°/4° of CaCl₂ + Aq contain 12.1638% CaCl₂ = 1.10489. (Schönrock, phys. Ch. 1893, 11. 768.)

Sp. gr. of CaCl ₂ +Aq at 17.925°C.					
Sp. gr.	% CaCl ₂	Sp. gr.	% CaCl ₂	Sp. gr.	
0.99869	13	1.11206	33	1.31562	
0.99954	14	1.12130	34	1.32689	
1.00037	15	1.13067	35	1.33821	
1.00116	16	1.14016	36	1.34956	
1.00201	17	1.14969	37	1.36100	
1.00371	18	1.15926	38	1.37242	
1.00539	19	1.16920	39	1.38400	
1.00703	20	1.17910	40	1.39489	
1.01127	21	1.18897	41	1.40641	
1.01548	22	1.19901	42	1.41770	
1.02386	23	1.20901	43	1.42882	
1.03238	24	1.21918	44	1.44007	
1.04089	25	1.22941	45	1.45124	
1.04951	26	1.23969	46	1.46238	
1.05822	27	1.25030	47	1.47329	
1.06680	28	1.26092	48	1.48450	
1.07569	29	1.27182	49	1.49573	
1.08467	30	1.28271	50	1.50676	
1.09373	31	1.29360	51	1.51778	
1.10288	32	1.30461			

(Pickering, B. 1894, 27. 1385.)

Sp. gr. of CaCl ₂ +Aq at t°.	
Concentration of CaCl ₂ +Aq	Sp. gr.
1 pt. CaCl ₂ in 7.1045 pts. H ₂ O	1.1062
1 " " " 164.25 " "	1.0032

(Hittorf, Z. phys. Ch. 1902, 39. 628.)

Sp. gr. of CaCl ₂ +Aq at 20°.	
g. mols. CaCl ₂ per l.	Sp. gr.
0.010	1.000982
0.025	1.002539
0.050	1.004874
0.075	1.006814
0.10	1.008971
0.25	1.02267
0.50	1.04451
0.75	1.06641
1.00	1.08744

nes and Pearce, Am. Ch. J. 1907, 38. 606.)

at. CaCl₂+Aq forms a crust at 150°, and
tains 178 pts. CaCl₂ to 100 pts. H₂O.
rlach.)
at. CaCl₂+Aq boils at 180°. (Rüdorff.)

B.-pt. of CaCl₂+Aq containing pts. CaCl₂ to
100 pts. H₂O. G=according to Gerlach
(Z. anal. 26. 440); L=according to Le-
grand (A. ch. (2) 39. 43).

B.-pt	G	L	B.-pt.	G	L
101°	6.0	10	134°	...	117.2
102	11.5	16.5	135	119	...
103	16.5	21.6	136	...	123.5
104	21.0	25.8	138	...	129.9
105	25.0	29.4	140	137.5	136.3
106	29.0	32.6	142	...	142.8
107	32.5	35.6	144	...	149.4
108	35.5	38.5	145	157	...
109	38.5	41.3	146	...	156.2
110	41.5	44.0	148	...	163.2
111	46.8	150	178	170.5
112	49.7	152	...	178.1
113	52.6	154	...	186.0
114	55.6	155	200	...
115	55.0	58.6	156	...	194.3
116	61.6	158	...	203.0
117	64.6	160	222	212.1
118	67.6	162	...	221.6
119	70.6	164	...	231.5
120	69.0	73.6	165	245	...
121	76.7	166	...	241.9
122	79.8	168	...	252.8
123	82.9	170	268	264.2
124	86.0	172	...	276.1
125	89.1	174	...	285.5
126	92.2	175	292	...
128	98.4	176	...	301.4
130	101	104.6	178	305	314.8
130.4	102.67	...	179.5	...	325.0
132	110.9

B.-pt. of CaCl₂+Aq.

% CaCl ₂	B.-pt.	% CaCl ₂	B.-pt.
5.6	101°	17.5	104°
10.3	102	20.0	105
14.5	103

(Skinner, Chem. Soc. 61. 340.)

Less sol. in HCl+Aq than in H₂O. HCl+
Aq sat. at 12° dissolves 27% CaCl₂, which
crystallizes out with 2H₂O. (Ditte, C. R. 92.
242.)

Solubility of CaCl₂ in HCl+Aq at 0°.

Sp. gr. of solutions	g. per 100 cc. solution	
	CaCl ₂	HCl
1.367	51.45	0.0
1.344	46.45	3.32
1.326	42.80	5.83
1.310	36.77	10.66
1.283	29.84	15.84
1.250	20.12	23.05
1.238	11.29	34.62

(Engel, C. R. 1887, 104, 434.)

CaCl₂+CaO₂H₂. Solubility of CaCl₂+CaO₂H₂ in H₂O at 25°.

% CaCl ₂	% CaO ₂ H ₂	Solid phase
5.02	0.101	CaO ₂ H ₂
10.00	0.115	"
12.94	0.128	"
15.14	0.140	"
17.20	0.145	"
18.15	0.148	CaO ₂ H ₂ +CaCl ₂ , 4CaO.14H ₂ O
18.01	0.152	CaCl ₂ , 4CaO.14H ₂ O
21.02	0.147	"
23.80	0.146	"
24.33	0.147	"
28.37	0.170	"
29.54	0.180	"
32.67	0.225	CaO ₂ H ₂ (?)
33.21	0.245	CaCl ₂ , 4CaO.14H ₂ O
33.72	0.254	CaCl ₂ , 4CaO.14H ₂ O+CaCl ₂ , CaO.2H ₂ O
34.36	0.173	CaCl ₂ , CaO.2H ₂ O
38.61	0.060	"
41.32	0.048	"
44.30	0.030	"
44.60	0.029	CaCl ₂ , 6H ₂ O+CaCl ₂ , CaO.2H ₂ O
44.77	...	CaCl ₂ .6H ₂ O

(Schreinemakers and Figuee, Chem. Weekbl. 1911, 8. 685.)

See also under Calcium hydroxide

CaCl₂+KCl. 100 pts. H₂O dissolve 56 pts. CaCl₂ at 7°; 100 pts. H₂O dissolve 31 pts. KCl at 7°; 100 pts. H₂O dissolve 63.5 pts. CaCl₂+4.9 pts. KCl at 7°. (Mulder, J. B. 1866. 67.)

CaCl₂+NaCl. 100 pts. H₂O dissolve 53 pts. CaCl₂ at 4°, and 56 pts. at 7°; 100 pts. H₂O dissolve 35.7 pts. NaCl at 4°, and 35.7 pts. at 7°; 100 pts. H₂O dissolve 57.6 pts. CaCl₂+2.4 pts. NaCl at 4°; 100 pts. H₂O dissolve 59.5 pts. CaCl₂+4.6 pts. NaCl at 7°. (Mulder, l. c.)

100 g. H₂O dissolve 72.6 g. CaCl₂+16.0 g. NaCl at 15°. (Rüdorff.)

Sol. in sat. KNO₃+Aq. (Fourcroy.)

Insol. in liquid CO₂. (Büchner, Z. phys. Ch. 1906, 54. 674.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

Sol. in 1 pt. strong boiling alcohol. (Wenzel.)

Sol. in 8 pts. alcohol at 15°, and in 1 pt. spirits of wine. (Bergman.)

Sol. in 0.7 pt. boiling absolute alcohol. (Otto.)

Sol. in 1.43 pts. boiling absolute alcohol at 78.3°. (Graham.)

Solubility of CaCl₂ in methyl alcohol.

CaCl₂ forms with methyl alcohol two complexes: CaCl₂.4CH₃OH and CaCl₂.3CH₃OH.

Solubility of CaCl₂, 4CH₃OH in CH₃OH.

t°	% by weight of CaCl ₂ . 4CH ₃ OH
0	33.3
10	37.6
20	42.2
30	47.0
40	52.0
50	57.3
55	60.0
56	61.3

Solubility of CaCl₂, 3CH₃OH in CH₃OH.

t°	% by weight of CaCl ₂ . 3CH ₃ OH
55	60.5
75	63.1
95	66.3
115	70.3
135	75.2
155	81.8
165	86.2
170	89.5
174	93.5
177 (mpt.)	100

(Menschutkin, Z. anorg. 1907, 52. 21.)

Solubility of CaCl₂ in ethyl alcohol.

CaCl₂ forms with ethyl alcohol a complex, CaCl₂.3C₂H₅OH.

Solubility of CaCl₂.3C₂H₅OH in C₂H₅OH at t°.

t°	% by weight of CaCl ₂ .3C ₂ H ₅ OH	t°	% by weight of CaCl ₂ .3C ₂ H ₅ OH
0	34.8	80	86.8
20	46.0	85	89.2
40	58.7	90	91.9
60	73.0	95	96.2
70	80.8	97 mpt.	100

(Menschutkin, Z. anorg. 1907, 52. 23.)

Sp. gr. at 16°/4° of CaCl₂+alcohol containing 5.668% CaCl₂=0.83636. (Schönrock, Z. phys. Ch. 1893, 11. 768.)

B.-pt. of an alcoholic solution of CaCl₂.

% CaCl ₂	B.-pt.
2.4	78.43° + 0.70°
5.39	78.43° + 2.15
8.01	78.32 + 4.18
9.93	78.43 + 5.55
15.94	78.43 + 11.75

(Skinner, Chem. Soc. 61. 340.)

Sl. sol. in propyl alcohol. (Berthelot.)

100 g. propyl alcohol dissolve 10.75 g. CaCl₂. (Schlamp, Z. phys. Ch. 1894, 14, 276.)

Sl. sol. in amyl alcohol. (Bouis.)

Pptd. from alcoholic solution by ether. (Döbbereiner.)

sol. in wood-spirit; sol. in lignone (Liebig);
 sol. in lignone. (Gmelin.)
 insol. in acetone; sol. in butyl alcohol.
 (urtz.)
 Very sl. sol. in acetone. (Krug and
 Elroy, J. Anal. Ch. 6. 184.)
 Solubility in acetone + Aq at 20°.

Cl₂ will salt out acetone from aqueous
 solution. The table shows the composi-
 tion of the solutions at the points at
 which inhomogeneous solutions of CaCl₂,
 acetone and H₂O just become homogen-
 eous at 20°. 100 g. of the solution contain:

g. CaCl ₂	g. H ₂ O	g. acetone
13.03	52.49	34.48
8.5	45.37	46.15
6.38	39.51	54.11
5.35	35.95	58.70
4.11	31.8	64.09
3.58	29.88	66.54
3.31	28.59	68.10
3.04	27.03	69.93
2.77	26.67	70.56
13.90	53.47	32.63
10.12	48.86	41.02
8.47	45.59	45.94
6.92	41.24	51.84
6.31	39.15	54.54
5.28	36.09	58.63
4.94	34.72	60.34
4.37	33.8	61.83
1.99	23.38	74.63
1.6	21.4	77.00
1.35	19.92	78.73
18.787	55.301	25.913
12.443	52.153	35.404
10.70	49.61	39.69
9.59	47.75	42.66
8.82	46.04	45.14
7.48	42.75	49.77
7.07	41.54	51.39
6.72	40.48	52.8
30.04	49.39	20.57
18.23	55.01	26.76
15.49	54.00	30.51
13.18	52.52	34.3
11.40	50.20	38.40
28.09	51.71	20.20
26.81	52.01	21.18
22.67	55.66	21.67
18.189	56.21	25.60
31.21	48.00	20.81
2.23	24.93	72.84
1.82	22.27	75.89
0.68	15.87	83.44
0.58	14.93	84.49
0.45	13.55	86.00
0.48	14.49	85.13
0.27	12.31	87.42
0.20	9.95	89.85
0.15	9.05	90.81

Rankforter, J. Am. Chem. Soc. 1914, 36.
 1125.)

Sol. in many compound ethers, as ethyl
 acetate (Liebig), ethyl lactate (Strecker).
 Sol. in considerable quantity in amyl sul-
 phocyanide. (Medlock, Chem. Soc. 1. 374.)
 Sol. in valyl. (Kolbe.)
 Very sol. in conc. HC₂H₃O₂. (Liebig.)
 Solubility of CaCl₂ in acetic acid.
 CaCl₂ forms with acetic acid a complex,
 CaCl₂, 4CH₃COOH.

Solubility of CaCl₂, 4CH₃COOH in
 CH₃COOH at t°.

t°	% by wt. CaCl ₂ , 4CH ₃ COOH
11.1	42.0
30	47.6
35	50.0
40	54.7
45	63.0
50	69.5
60	79.5
65	84.5
70	91.2
73	100.0

(Menschutkin, Z. anorg. 1907, 54. 95.)

Insol. in benzonitrile. (Naumann, B. 1914,
 47. 1370.)

Insol. in ethyl acetate. (Naumann, B.
 1910, 43. 314.)

Sl. sol. in anhydrous pyridine.

Sol. in 97%, 95% and 93% pyridine + Aq.
 (Kahlenberg, J. Am. Chem. Soc. 1908, 30.
 1107.)

100 g. sat. solution of CaCl₂ in sat. sugar +
 Aq at 31.25° contain 42.84 g. sugar + 25.25 g.
 CaCl₂, or 100 g. H₂O dissolve 135.1 g. sugar
 + 79.9 g. CaCl₂ at 31.25°. (Köhler, Z. Ver.
 Zuckerind, 1907, 47. 447.)

+H₂O. (Bakhuis Roozeboom.) See above.

+2H₂O. (Bakhuis Roozeboom.) See above.

+4H₂O. Two modifications. (Bakhuis
 Roozeboom.) See above.

α and β modifications (α = stable form.)
 (Kuznetsov, C. A. 1911, 842.)

+6H₂O. Very deliquescent. Sol. in H₂O
 with absorption of much heat.

250 pts. CaCl₂ + 6H₂O with 100 pts. H₂O
 at 10.8° lower the temp. 23.2°. (Rüdorff, B.
 2. 68.)

Melts in crystal H₂O at 28° (Tilden, Chem.
 Soc. 45. 409); at 30.2° (Bakhuis Roozeboom.)

Sat. solution in H₂O contains at:

—22°	—17°	—5°	—5°	+4°
31.5	32.4	35.1	35.2	36.5% salt,
8°	22°	29°	35°	49°
37.9	42.1	46.1	49.0	55.1% salt,
63°	80°	104°	115°	
55.9	57.5	58.5	58.6% salt.	

(Étard, A. ch. 1894, (7) 2. 532.)

Sat. solution of CaCl₂ + 6H₂O contains

44.77 g. CaCl_2 at 25° . (Schreinemakers and Figuee, Chem. Weekbl. 1911, 8. 685.)

See also above.

Solubility of $\text{CaCl}_2 + 6\text{H}_2\text{O}$ in ethyl alcohol + Aq under addition of increasing amounts of CaCl_2 .

Per cent of alcohol by volume	G. CaCl_2 added	Grams CaCl_2 in 5 cc. of solution
92.3		1.430
97.3		1.409
99.3		1.429
"	1	1.529
"	2	1.561
"	3	1.590
"	4	1.641
"	5	1.709

(Bödtker, Z. phys. Ch. 1897, 22. 510.)

Calcium hydroxylamine chloride, $\text{CaCl}_2 \cdot 3\text{NH}_2\text{OH} \cdot \text{HCl}$.

(Antonow, J. Russ. Phys. Chem. Soc. 1905, 37. 479.)

Calcium iodine trichloride, $2\text{ICl}_3 \cdot \text{CaCl}_2 + 8\text{H}_2\text{O}$.

Hydroscopic. (Weinland, Z. anorg. 1902, 30. 142.)

Calcium mercuric chloride, $\text{CaCl}_2 \cdot 5\text{HgCl}_2 + 8\text{H}_2\text{O}$.

Decomp. by cold H_2O , which dissolves out CaCl_2 , but all dissolves on heating. (v. Bonsdorff, 1829.)

$\text{CaCl}_2 \cdot 2\text{HgCl}_2 + 6\text{H}_2\text{O}$. Deliquescent. Very sol. in H_2O . (v. Bonsdorff.)

$\text{CaCl}_2 \cdot 6\text{HgCl}_2 + 6\text{H}_2\text{O}$. Very deliquescent. Decomp. by H_2O . (Strömholm, J. pr. 1902, (2) 66. 521.)

Calcium lead chloride, basic.

See Calcium lead oxychloride.

Calcium magnesium chloride, $\text{CaCl}_2 \cdot 2\text{MgCl}_2 + 12\text{H}_2\text{O}$.

Min. *Tachhydrite*. Deliquescent.

100 pts. H_2O dissolve 160.3 pts. at 18.75° . By dissolving 20 pts. in 80 pts. H_2O the temp. is raised 7.75° . (Bischof.)

Calcium mercuric chloride, basic, $\text{CaCl}_2 \cdot 2\text{HgO} + 4\text{H}_2\text{O}$.

See Calcium mercuric oxychloride.

Calcium thallic chloride, $2\text{TlCl}_3 \cdot \text{CaCl}_2 + 6\text{H}_2\text{O}$.

Can be cryst. from H_2O . (Gewecke A. 1909, 366. 222.)

Calcium tin (stannic) chloride.

See Chlorostannate, calcium.

Calcium uranium chloride, $\text{CaCl}_2 \cdot \text{UCl}_4$.

Decomp. by H_2O . (Aloy, Bull. Soc. 1899, (3) 21. 265.)

Calcium zinc chloride.

$\text{CaZnCl}_4 + 5\frac{1}{2}\text{H}_2\text{O}$, and $\text{Ca}_2\text{ZnCl}_6 + 6\text{H}_2\text{O}$. Very hygroscopic. (Ephraim, Z. anorg. 1910, 67. 379.)

Calcium chloride ammonia, $\text{CaCl}_2 \cdot 8\text{NH}_3$.

Sol. in H_2O with decomp. (Faraday.)

Calcium chloride hydrazine, $\text{CaCl}_2 \cdot 2\text{N}_2\text{H}_4(?)$.

Ppt. (Franzen, Z. anorg. 1908, 60. 288.)

Calcium chloride hydroxylamine, $\text{CaCl}_2 \cdot \text{NH}_2\text{OH} + 5\text{H}_2\text{O}$.

Not hygroscopic. (Antonow, J. Russ. Phys. Chem. Soc. 1905, 37. 479.)

$\text{CaCl}_2 \cdot 2\text{NH}_2\text{OH}$.

+ H_2O . Aqueous solution sat. at 20° contains 56.6 pts. salt.

+ $2\text{H}_2\text{O}$. (Antonow, l. c.)

$2\text{CaCl}_2 \cdot 3\text{NH}_2\text{OH} + 6\text{H}_2\text{O}$. (Antonow, l. c.)

$2\text{CaCl}_2 \cdot 5\text{NH}_2\text{OH} + 4\text{H}_2\text{O}$. (Antonow, l. c.)

Calcium chloride lead oxide, $\text{CaCl}_2 \cdot 3\text{PbO} + 3\text{H}_2\text{O}$.

See Calcium lead oxychloride.

Calcium chloroferrite, $\text{CaO} \cdot \text{CaCl}_2 \cdot \text{Fe}_2\text{O}_3$.

Insol. in H_2O . (le Chatelier, C. R. 99. 276.)

Calcium chlorofluoride, $\text{CaF}_2 \cdot \text{CaCl}_2$.

Decomp. by H_2O , by very dil. HCl , HNO_3 , or acetic acid, by hot dil. or conc. H_2SO_4 . Sol. in conc. HCl or HNO_3 . Insol. in, and not decomp. by cold or boiling alcohol. (Defacqz, A. ch. 1904, (8) 1. 355.)

Calcium cyanamide, basic, $\text{CN}_2(\text{CaOH})_2 + 6\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Meyer, J. pr. 1878, (2) 18. 425.)

Calcium cyanamide, CaCN_2 .

Decomp. by H_2O . (Meyer, J. pr. 1878, (2) 18. 425.)

Calcium subfluoride, CaF .

Decomp. by H_2O .

Sol. in hot dil. HCl and somewhat sol. in dil. acetic acid.

Somewhat sol. in boiling absolute alcohol. (Wöhler, Z. anorg. 1909, 61. 81.)

Calcium fluoride, CaF_2 .

Sol. in 26,923 pts. H_2O at 15.5° . (Wilson, Ch. Gaz. 1850. 366.)

1 l. H_2O dissolves 16 mg. CaF_2 at 18° . (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)

16.3 mg. in 1 l. of sat. solution at 18° . (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

When pptd. not completely insol. in H_2O ; scarcely sol. in dil., more sol. in conc. $HCl + Aq$; decomp. by conc. H_2SO_4 ; not decomp. by dil. alkaline solutions. (Fresenius.)

Not decomp. by conc. H_2SO_4 below 40° , but forms a transparent syrup. CaF_2 is pptd. from this solution by addition of H_2O .

Sol. in conc. HCl , and $HNO_3 + Aq$ in the same way, but the liquid is not viscid. Very sl. sol. in HF . Boiling $HCl + Aq$ dissolves slightly. Decomp. by boiling $HNO_3 + Aq$.

Sol. in NH_4 salts + Aq . (Rose.)

Partly decomp. by boiling K_2CO_3 , and $Na_2CO_3 + Aq$. (Dulong, A. ch. 82. 278.)

Insol. in liquid HF . (Franklin, Z. anorg. 1905, 46. 2.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Min. *Fluorite* (*Fluorspar*). Calculated from electrical conductivity of $CaF_2 + Aq$, 1 l. H_2O dissolves 14 mg. CaF_2 at 18° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

Calcium hydrogen fluoride, $CaH_2F_4 + 6H_2O$.

Decomp. by boiling H_2O . Sol. in $HF + Aq$. (Freymy, A. ch. (3) 47. 35.)

Calcium tantalum fluoride.

See *Fluotantalate, calcium*.

Calcium stannic fluoride.

See *Fluostannate, calcium*.

Calcium titanium fluoride.

See *Fluotitanate, calcium*.

Calcium fluoiodide, CaF_2, CaI_2 .

Very deliquescent. Decomp. by cold H_2O , more rapidly by hot H_2O , by dil. HCl , HNO_3 , H_2SO_4 , conc. H_2SO_4 , and by alcohol and by ether if these reagents are not absolute. (Defacqz, A. ch. 1904, (8) 1. 358.)

Calcium hydride, CaH_2 .

Decomp. by $HCl + Aq$. (Winkler, B. 24. 1975. (Moldenhauer, Z. anorg. 1913, 82. 136.)

CaH_2 . Readily decomp. by H_2O and dil. acids, almost insol. in conc. acids. Insol. in benzene, turpentine and alkyl haloids. (Moissan, C. R. 1898, 127. 30-31.)

Decomp. H_2O and ether; sol. in dil. H_2SO_4 and HNO_3 ; almost insol. in conc. H_2SO_4 and HNO_3 . (von Lengyll, C. C. 1898, II. 262.)

Insol. in CCl_4 , CS_2 , alcohols and ethers. No known solvent. (Moissan, C. C. 1903, I. 863.)

Calcium hydrosulphide, CaS_2H_2 .

Cryst. with $6H_2O$. Extremely sol. in H_2O and alcohol. $\frac{1}{4}$ of its weight of H_2O at or-

dinary temp. more than suffices to hold it in solution. (Divers and Shimidzu, Chem. Soc. 45. 271.)

Sp. gr. of aqueous solution containing 32% anhydrous CaS_2H_2 , (64% $CaS_2H_2 + 6H_2O$) = 1.255; 37.5% CaS_2H_2 , (75.5% $CaS_2H_2 + 6H_2O$) = 1.310. (Divers and Shimidzu.)

Calcium hydroxide, CaO_2H_2 .

See also *Calcium oxide*.

Sl. sol. in cold, and less in hot H_2O .

1 pt. CaO dissolves at t° in pts. H_2O .

t°	Pts. H_2O	Authority
20	450	Davy.
0	656	Phillips (A. Phil. 17. 107)
	700	Bergman (Essays, etc.).
13	785	Pavesi and Rotondi (B. 7. 817)
18	780	Bineau (A. ch. (3) 51. 290).
19.5	806	P. and R. (l. c.).
23	814	P. and R. (l. c.).
18.75	960	Abl.
54.4	972	Dalton (Syst. 2. 231).
15.6	778	Dalton (l. c.).
15.6	752	Phillips (l. c.).
15.6	731	Wittstein (Repert. Pharm. 1. 182).
15.6	741	Tichborne (Bull. Soc. (2) 17. 24).
100	1270	Dalton (l. c.).
100	1280	Phillips (l. c.).
100	1330	Wittstein (l. c.).
100	1340	Tichborne (l. c.).
100	1500	Bineau (l. c.).
100	1758	Tichborne (l. c.).

Solubility in H_2O . 1000 pts. $CaO_2H_2 + Aq$ sat. at t° contain pts. CaO .

t°	Pts. CaO		
	From Nitrate	Marble	Hydrate
0	1.362	1.381	1.430
10	1.311	1.342	1.384
15	1.277	1.299	1.344
30	1.142	1.162	1.195
45	0.996	1.005	1.033
60	0.884	0.868	0.885
100	0.562	0.576	0.584

(Lamy, C. R. 86. 333.)

Solubility of CaO_2H_2 in H_2O at t° .

t°	Pts. H_2O to 1 pt. CaO	Pts. CaO in 100 pts. H_2O	t°	Pts. H_2O to 1 pt. CaO	Pts. CaO in 100 pts. H_2O
0	759	0.131	60	1136	0.088
10	770	0.129	70	1235	0.080
20	791	0.126	80	1362	0.073
30	862	0.116	90	1579	0.063
40	932	0.107	100	1650	0.060
50	1019	0.098

(Maben, Pharm. J. Trans. (3) 14. 505.)

1 pt. CaO_2H_2 is sol. in 640 pts. H_2O at 19° , and 3081 pts. at 150° . (Shenstone and Cundall, Chem. Soc. 53. 550.)

1000 g. H_2O dissolve 1.251 g. CaO . (Carles, Arch. Pharm. (3) 4. 558.)

Solubility of CaO_2H_2 in H_2O . 100 pts. H_2O dissolve pts. CaO at t° .

t°	Pts. CaO	t°	Pts. CaO
20	0.1374	80	0.0845
40	0.1162	100	0.0664
60	0.1026

(Zahorsky, Z. anorg. 3. 34.)

1 pt. CaO is sol. in pts. H_2O at t° .
 t° 15° 20° 25° 30° 35° 40° 45°
pts. H_2O 776 813 848 885 924 962 1004

t° 50° 55° 60° 65° 70° 75° 80°
pts. H_2O 1044 1108 1158 1244 1330 1410 1482

(Herzfeld, C. C. 1897, I, 932.)

100 g. sat. $\text{CaO}_2\text{H}_2 + \text{Aq}$ contain g. CaO at t° :

t° 5 10 15 20 25
g. CaO 0.135 0.1342 0.132 0.1293 0.1254

t° 30 35 40 50 60
g. CaO 0.1219 0.1161 0.1119 0.0981 0.0879

t° 70 80 90 100
g. CaO 0.0781 0.074 0.0696 0.0597

(Guthrie, J. Soc. Chem. Ind. 1901, 20. 223.)

Solubility in H_2O at high temp.
1 litre of the solution contains at:
120° 150° 190°
0.305 0.169 0.084 g. CaO .

(Herold, Z. elektrochem. 1905, 11. 421.)

Solubility in H_2O at t° .

t°	1 g. CaO is sol. in g. H_2O at t°
2	768.5
10	786.8
15	804.3
20	826.4
25	868.7
30	908.2
40	988.1
50	1083.0
60	1179.0
70	1274.8
80	1368.1

(Moody, Chem. Soc. 1908, 93. 1772.)

Sat. $\text{CaO}_2\text{H}_2 + \text{Aq}$ contains at:
95° 76°
0.0580 0.0705% by wt. CaO .

(Tschugaeff, Z. anorg. 1914, 86. 159.)

100 g. sat. solution of CaO_2H_2 in H_2O at 25° contain 0.117 g. CaO_2H_2 . (Cameron and Potter, J. phys. Ch. 1911, 15. 70.)
Readily sol. in most acids.

Sol. in $\text{H}_3\text{BO}_3 + \text{Aq}$ at 30°. (Sborgi, Real. Ac. Linc. 1913, (5) 22. I, 715 and 798.)
Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. Much more sol. in $\text{NaCl} + \text{Aq}$ than in H_2O . (Rose.)

Solubility of CaO_2H_2 in $\text{NH}_4\text{Cl} + \text{Aq}$ at 25°.

Concentration of $\text{NH}_4\text{Cl} + \text{Aq}$ in millimols per liter	Solubility of CaO_2H_2 in millimols per liter
0.00	20.22
21.76	29.08
43.52	39.23
87.03	59.68

(Noyes and Chapin, Z. phys. Ch. 1899, 22. 520.)

Solubility of CaO_2H_2 in $\text{CaCl}_2 + \text{Aq}$. 100 pts. $\text{CaCl}_2 + \text{Aq}$ of given strength dissolve pts. CaO at t° .

t°	$\text{CaCl}_2 + \text{Aq}$ 5% CaCl_2	$\text{CaCl}_2 + \text{Aq}$ 10% CaCl_2	$\text{CaCl}_2 + \text{Aq}$ 15% CaCl_2	$\text{CaCl}_2 + \text{Aq}$ 20% CaCl_2	$\text{CaCl}_2 + \text{Aq}$ 25% CaCl_2	$\text{CaCl}_2 + \text{Aq}$ 30% CaCl_2
20	0.1370	0.1661	0.1993	0.1857*	0.1661*	0.1630*
40	0.1160	0.1419	0.1781	0.2249	0.3030*	0.3684*
60	0.1020	0.1313	0.1706	0.2204	0.2989	0.3664
80	0.0936	0.1328	0.1736	0.2295	0.3261	0.4122
100	0.0906	0.1389	0.1842	0.2325	0.3710	0.4922

* In these cases, ppts. of 3CaO , $\text{CaCl}_2 + 15\text{H}_2\text{O}$ were formed.
(Zahorsky, Z. anorg. 3. 34.)

See also $\text{CaCl}_2 + \text{CaO}_2\text{H}_2$ under Calcium chloride.

Solubility in $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 25°.

Sp. gr. 25°/25°	g. CaO as $\text{Ca}(\text{OH})_2$ in 100 g. H_2O	g. $\text{Ca}(\text{NO}_3)_2$ in 100 g. H_2O	Solid phase
1.0249	0.096	3.38	$\text{Ca}(\text{OH})_2$
1.0484	0.109	8.52	
1.0940	0.125	13.42	
1.1383	0.181	20.73	
1.1840	0.187	28.98	$\text{Ca}(\text{OH})_2$ and solid solution $\text{CaO}, x\text{N}_2\text{O}_5, y\text{H}_2\text{O}$
1.2101	0.198	32.84	
1.2287	0.212	36.83	
1.2290	0.213	37.55	
1.2541	0.224	40.25	Solid solution $\text{CaO}, x\text{N}_2\text{O}_5, y\text{H}_2\text{O}$
1.2581	0.230	41.98	
1.2826	0.260	47.00	
1.2905	0.263	47.16	
1.3337	0.332	58.67	Solid solution $\text{CaO}, x\text{N}_2\text{O}_5, y\text{H}_2\text{O}$ and $\text{Ca}(\text{NO}_3)_2 \cdot 3\frac{1}{2}\text{H}_2\text{O}$
1.3735	0.429	69.40	
1.4195	0.545	83.03	
1.4840	0.449	99.70	
1.5330	0.371	115.50	$\text{Ca}(\text{NO}_3)_2 \cdot 3\frac{1}{2}\text{H}_2\text{O}$ and $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
1.5809	0.303	135.30	
1.5842	0.000	139.30	

(Cameron and Robinson, J. phys. Chem. 1907, 11. 275.)

solubility of CaO_2H_2 in $\text{Ca}(\text{NO}_3)_2 + \text{Aq.}$
Temp. = 25°.

per 100 g. sat. solution		Solid phase
$\text{Ca}(\text{NO}_3)_2$		
0	0	CaO_2H_2
3	4.84	"
4	9.36	"
3	13.77	"
4	22.46	"
0	27.83	"
1	32.94	"
9	40.66	"
0	44.44	"
2	45.28	$\text{Ca}_2\text{N}_2\text{O}_7 \cdot 3\text{H}_2\text{O}$
4	47.79	"
4	51.07	"
9	53.20	"
6	55.25	"
6	57.72	$\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
	57.98	"

Temp. = 100°.

1	0	CaO_2H_2
0	2.42	"
4	4.91	"
0	15.39	"
0	16.10	"
	21.86	"
	33.03	"
	42.26	"
	50.94	"
	53.75	"
	55.40	"
	55.43	"
	55.65	"
	56.89	$\text{CaO}_2\text{H} + \text{Ca}_2\text{N}_2\text{O}_7 \cdot 2\text{H}_2\text{O}$
	57.03	$\text{Ca}_2\text{N}_2\text{O}_7 \cdot 2\text{H}_2\text{O}$
	57.91	"
	58.67	"
	60.44	"
	62.82	"
	66.44	"
	69.12	"
	70.60	$\text{Ca}_2\text{N}_2\text{O}_7 \cdot 2\text{H}_2\text{O} + \text{Ca}_2\text{N}_2\text{O}_7 \cdot \frac{1}{2}\text{H}_2\text{O}$
	70.40	$\text{Ca}_2\text{N}_2\text{O}_7 \cdot \frac{1}{2}\text{H}_2\text{O}$
	71.44	"
	73.85	"
	75.74	"
	76.94	"
	77.62	$\text{Ca}(\text{NO}_3)_2$
	77.74	"
	78.43	"

ett and Taylor, Chem. Soc. 1914, 105.
1926.)

solubility of CaO in KCl and $\text{NaCl} + \text{Aq.}$
values are given which show that the solu-
of lime in solutions of either NaCl or

KCl is a maximum for all temps. when the solution contains about 60 g. of salt per l. It is a minimum at any fixed temp. when the solution is sat., the solubility then being much less than in pure H_2O of the same temp. A solution of NaCl dissolves more lime at all temps. and concentrations than a corresponding solution of KCl . In all cases the maximum solubility of lime occurs when the temp. is lowest. With solutions of all concentrations the solubility decreases regularly as the temp. increases. (Cabot, J. Soc. Chem. Ind. 1897, 16. 417-419.)

Solubility in $\text{KCl} + \text{Aq}$ increases with increased quantities of KCl and then diminishes, becoming less than the solubility in H_2O alone. (Kernot, Gazz. ch. it. 1908, 38. (1) 532.)

KOH or $\text{NaOH} + \text{Aq}$ containing 1 pt. KOH or NaOH in 100 pts. H_2O do not dissolve more than $\frac{1}{10000}$ pt. CaO_2H_2 , but it is sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Pelouze, A. ch. (3) 33. 11.)

Solubility in $\text{NaOH} + \text{Aq}$ at t°.

G. NaOH per l	Solubility of CaO in g. per liter at			
	20°	50°	70°	100°
0	1.17	0.88	0.75	0.54
0.400	0.94	0.65	0.53	0.35
1.600	0.57	0.35	0.225	0.14
2.666	0.39	0.20	0.11	0.05
5.000	0.18	0.06	0.04	0.01
8.000	0.11	0.02	0.01	traces
20.000	0.02	traces	0	0

(d'Anselme, Bull. Soc. 1903, (3) 29. 936.)

Solubility of CaO in $\text{NaCl} + \text{NaOH} + \text{Aq.}$

G. NaCl per l.	G. CaO per l. of solution containing		
	No NaOH	0.89 g. NaOH per l.	4.09 g. NaOH per l.
0	1.3	0.8	0.22
5	1.4	0.9
10	1.6	1.0
25	1.7	1.1
50	1.8	1.25
75	1.9	1.4	0.55
100	1.85	1.4
150	1.65	1.25	0.44
175	1.6	1.2
182	1.6	1.2
225	1.4	1.0
250	1.3	0.9
300	1.1	0.7	0.22

(Maigret, Bull. Soc. 1905, (3) 33. 631.)

Solubility of CaO_2H_2 in $\text{CaSO}_4 + \text{Aq}$ at 25° .

G. per 100 cc. sat. solution		Solid phase
CaSO_4	CaO	
0	0.1166	CaO_2H_2
0.0391	0.1141	"
0.0666	0.1150	"
0.0955	0.1215	"
0.1214	0.1242	"
0.1588	0.1222	$\text{CaO}_2\text{H}_2 + \text{CaSO}_4 \cdot 2\text{H}_2\text{O}$
0.1634	0.0939	$\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$
0.1722	0.0611	"
0.1853	0.0349	"
0.1918	0.0176	"
0.2030	0.0062	"
0.2126	0	"

(Cameron and Bell, J. Am. Chem. Soc. 1906, 28. 1220.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Alcohol dissolves traces.

Methyl alcohol forms colloidal solution containing 1.125 g. per l. (Neuberg and Rewald, Biochem. Z. 1908, 9. 545.)

Insol. in ether.

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Much more sol. in glycerine, or sugar + Aq than in H_2O .

Solubility of CaO in glycerine.

Wt. of glycerine in 100 ccm. of solution	Wt. CaO contained in 100 ccm. of liquid sat. with CaO	Relation of CaO to glycerine	
		CaO	Glycerine
10.00	0.370	3.6	96.4
5.00	0.240	4.6	95.4
2.86	0.196	6.4	93.6
2.50	0.192	7.1	92.9
2.00	0.186	8.5	91.5
1.00	0.165	14.2	85.8

(Berthelot, A. ch. (3) 46. 176.)

1000 g. H_2O dissolve 1.251 g. CaO ; 1000 g. $\text{H}_2\text{O} + 50$ g. glycerine dissolve 1.865 g. CaO ; 1000 g. $\text{H}_2\text{O} + 100$ g. glycerine dissolve 2.583 g. CaO ; 1000 g. $\text{H}_2\text{O} + 200$ g. glycerine dissolve 4.040 g. CaO ; 1000 g. $\text{H}_2\text{O} + 400$ g. glycerine dissolve 6.569 g. CaO . (Carles, Arch. Pharm. (3) 4. 558.)

Insol. in pure glycerine.

Solubility of CaO_2H_2 in glycerine + Aq at 25° .

G = g. glycerine in 100 g. glycerine + Aq.
 $\frac{1}{2}\text{Ca}(\text{OH})_2$ = millimols sol. in 100 cc. glycerine + Aq.

G	$\frac{1}{2}\text{CaO}_2\text{H}_2$	Sp. gr.
0	4.3	1.0003
7.15	8.13	1.0244
20.44	14.9	1.0537
31.55	22.5	1.0842
40.95	40.1	1.1137
48.7	44.0	1.1356
69.2	95.8	1.2027

(Herz and Knoch, Z. anorg. 1905, 46. 193.)

Solubility in glycerine + Aq at 25° .

Solution contains			Sp. gr.
% $\text{Ca}(\text{OH})_2$	% glycerine	% H_2O	
0.117	0	96.32	1.008
0.178	3.50	80.28	1.042
0.413	15.59	81.68	1.088
0.48	17.84	64.80	1.149
0.88	34.32	43.62	
1.34	55.04		

Solid phase in this system is CaO_2H_2 .
(Cameron and Patten, J. phys. Chem. 1911, 15. 71.)

100 pts. sugar dissolved in H_2O dissolve 55.6 pts. CaO (Osann); 50 pts. CaO (Ure); 49.6 pts. CaO (Daniell); 29–30.6 pts. CaO (Hunton); 23 pts. CaO . (Soubeiran.)

Sugar solution at 100° takes up $\frac{1}{4}$ mol. CaO for each mol. sugar; at 0° , if it contains not less than 25% of sugar, it takes up 2 mols. CaO to 1 mol. sugar. (Debrunfaut.)

Amount dissolved is proportional to the density and temperature of the solutions.

Solubility of CaO in sugar + Aq.

Pts. sugar dissolved in 100 pts. H_2O	Relation of CaO to sugar	
	CaO	Sugar
40	21.0	79.8
37.5	20.8	79.2
35.0	20.5	79.5
32.5	20.3	79.7
30.0	20.1	79.9
27.5	19.9	80.1
25.0	19.8	80.2
22.5	19.3	80.7
20.0	18.8	81.2
17.5	18.7	81.3
15.0	18.5	81.5
12.5	18.3	81.7
10.0	18.1	81.9
7.5	16.9	83.1
5.0	15.3	84.7
2.5	13.8	86.2

(Peligot, C. R. 33. 335.)

100 g. solution of sugar sat. with CaO between 10° and 54.4° contain 22.5 to 23.5% CaO . (Hunton, 1897.)

Solubility of CaO in dil. sugar solutions.

Wt. of sugar in 100 ccm. solution	Wt. of CaO contained in 100 ccm of liquid sat. with CaO	Relation of CaO to sugar	
		CaO	Sugar
850	1.031	17.5	82.5
401	0.484	16.8	83.2
000	0.433	17.8	82.2
660	0.364	18.0	82.0
386	0.326	19.0	81.0
200	0.316	20.8	79.2
058	0.281	21.0	79.0
960	0.264	21.6	78.4
400	0.194	32.7	67.3
191	0.172	47.4	52.6
096	0.154	61.6	78.4
000	0.148		

(Berthelot, A. ch. (3) 46. 176.)

Solubility in sugar + Aq at t°.

t°	G. sugar in 100 ccm. of solution	G. CaO dissolved per 100 g. sugar
	0.7814	37.9
	0.9120	32.3
	1.4000	30.5
	1.6930	28.9
	4.754	27.7
	5.730	27.1
	10.159	27.5
	11.200	27.2
	12.500	27.3
	13.930	27.9
	14.487	27.5
	16.410	28.0
	0.625	71.6
	0.964	53.4
	2.084	36.0
	3.028	32.3
	3.451	31.7
	4.168	30.2
	4.880	28.7
	5.73	28.3
	6.12	27.4
	6.25	27.7
	6.51	27.5
	7.55	27.9
	8.20	27.3

enaberg, Bull. Soc. 1899, (3) 21. 775.)

Solubility in sugar + Aq at 25°.

Solution contains			Sp. gr.
% Ca(OH) ₂	% sugar	% H ₂ O	
0.117	0		0.983
0.188	0.62	99.19	1.000
0.730	4.82	94.50	1.021
1.355	7.50	91.12	1.037
2.31	9.87	87.85	1.051
3.21	11.90	84.89	1.067
4.57	15.10	80.33	1.092
5.38	17.42	76.93	1.109
6.07	19.86	73.07	1.123

The solid phase in this system consists of a series of solid solutions with Ca(OH)₂; a limiting case.

(Cameron and Patten, J. phys. Chem. 1911, 15. 70.)

Solubility of CaO in sugar + Aq at 80°.

% sugar	% CaO	% Sugar	% CaO
4.90	0.117	19.50	0.358
9.90	0.189	24.60	0.458
14.75	0.230	29.70	1.017

Solid phase, CaO.H₂.

(von Ginneken, Proc. Kon. Akad. v. Wetensch., Amsterdam, 1911, 14. 457.)

Solubility of CaO in mannite + Aq

Wt. of mannite in 100 ccm. of solution	Wt. of CaO contained in 100 ccm. of liquid sat. with CaO	Relation of CaO to mannite	
		CaO	Mannite
9.60	0.753	7.3	92.7
4.80	0.372	7.2	92.8
2.40	0.255	9.6	90.4
1.92	0.225	10.5	89.5
1.60	0.207	11.4	88.6
1.37	0.194	12.5	87.5
1.20	0.193	13.9	86.1
1.07	0.190	15.1	84.9
0.96	0.186	16.2	86.8
0.192	0.155	44.6	55.4
0.096	0.154	61.6	38.4
0.000	0.148		

(Berthelot, A. ch. (3) 46. 176.)

Solutions of CaO in sugar, mannite, or glycerine afford an abundant ppt. on being heated, but this redissolves on cooling. (Berthelot.)

Sol. in sorbite + Aq (Pelouze); sl. sol. in quercite + Aq. Sol. in monobasic Ca saccharate + Aq (Peligot.) Much more sol. in gelatine + Aq than in pure H₂O.

Calcium hydroxyhydrosulphide, Ca(OH)SH + 3H₂O.

Easily sol. in H₂O with almost immediate decomposition. Insol. in alcohol, but slowly

decomp thereby. (Divers and Shimidzu, Chem Soc. 45. 270.)

Calcium subiodide, CaI .

Decomp. by moisture. (Wöhler, Z. anorg. 1909, 61. 76.)

Calcium iodide, CaI_2 .

Deliquescent. 100 pts. H_2O dissolve—
 at 0° 20° 40° 43° 92°
 192 204 228 286 435 pts. CaI_2 .
 (Kremers, Pogg. 103. 65.)

Sp. gr. of $\text{CaI}_2 + \text{Aq}$ at 19.5° containing:
 5 10 15 20 25 30% CaI_2 ,
 1.044 1.09 1.14 1.198 1.26 1.321

35 40 45 50 55 60% CaI_2 .
 1.398 1.477 1.567 1.665 1.78 1.91
 (Kremers, calculated by Gerlach, Z. anal. 8. 285.)

Sol in absolute alcohol. (Gay-Lussac, A. ch. 91. 57.)

Sol in acetone. (Naumann, B. 1904, 37. 4328; Eidmann, C. C. 1899, II. 1014.)

Sol in ethyl acetate. (Naumann, B. 1910, 43. 314.)

+4 H_2O . (Kuznetsov, C. A. 1911. 842.)

+6 H_2O . Sat. aq. solution contains at:

—22° +7° 10° 19°
 61.6 65.0 65.1 66.3% salt.

51° 64° 130° 248°
 69.4 75.9 81.3 87.1% salt.

(Étard, A. ch. 1894, (7) 2. 543.)

+7 H_2O . (Kuznetsov.)

Calcium periodide, $\text{CaI}_2 + 15\text{H}_2\text{O}$.

(Moanier, A. ch. 1897, (7) 12. 401.)

CaI_2 . (Herz and Bulla, Z. anorg. 1911, 71. 255.)

Calcium mercuric iodide, $\text{CaI}_2 \cdot \text{HgI}_2 + 8\text{H}_2\text{O}$

Very deliquescent. Sol. in H_2O , alcohols, allyl iodide, aldehyde, acetic acid, ethyl oxalate and aniline. Sl. sol. in nitrobenzene. Completely insol. in CHCl_3 , CCl_4 , ethyl iodide, ethylene bromide, C_6H_6 , monochlorobenzene and toluene. (Duboin, C. R. 1906, 142. 573.)

3 CaI_2 , 4 $\text{HgI}_2 + 24\text{H}_2\text{O}$. Sol. in H_2O with pptn. of red HgI_2 .

Very sol. in alcohols, glycerine, ethyl acetate, methyl and isobutyl propionate, allyl iodide, aldehyde, acetone, aniline and ethyl oxalate. Insol. or sl. sol. in nitrobenzene. Insol. in CHCl_3 , C_6H_6 , ethyl iodide, monochlorobenzene, etc. (Duboin, C. R. 1906, 142. 397.)

$\text{CaI}_2 \cdot 2\text{HgI}_2$.

Decomp. by H_2O . (Boullay.)

CaI_2 , 5 $\text{HgI}_2 + 8\text{H}_2\text{O}$. Decomp. by H_2O ,

alcohols, glycerine, aldehyde, and acetic acid; slowly by nitrobenzene and ethyl oxalate. Insol. in monochlorobenzene, toluene, CHCl_3 and ethylene bromide. (Duboin, l. c.)

Calcium silver iodide, CdI_2 , 2 $\text{AgI} + 6\text{H}_2\text{O}$.

Immediately decomp. by H_2O . (Simpson, Roy. Soc. Proc. 27. 120.)

Calcium zinc iodide, CaI_2 , $\text{ZnI}_2 + 8\text{H}_2\text{O}$.

Very hygroscopic. (Ephraim, Z. anorg. 1910, 67. 384.)

Calcium iodide ammonia, CaI_2 , 6 NH_3 .

(Isambert, C. R. 66. 1259.)

Calcium nitride, Ca_3N_2 .

Sol. in dil. acids; insol. in conc. (water free) acids. (Moissan, C. R. 1898, 127. 499.)

Calcium oxide, CaO .

Decomp. by H_2O , with evolution of much heat, to form CaO_2H_2 , which see for solubility in H_2C , etc.

Calcium peroxide, CaO_2 .

Very sl. sol. in H_2O ; easily sol. in acids, and NH_4 salts + Aq . Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Conroy, Chem. Soc. (2) 11, 808.)

+2 H_2O . True composition is $\text{CaO}_2\text{H}_2 + \text{H}_2\text{O}_2$. (de Forerand, C. R. 1900, 130. 1390.)

+8 H_2O . Efflorescent. Difficultly sol. in H_2O with gradual decomp. Insol. in alcohol or ether. (Gay-Lussac and Thénard, A. ch. (2) 8. 313.)

Calcium oxybromide, 3 CaO , $\text{CaBr}_2 + 16\text{H}_2\text{O}$.

Decomp. by H_2O and alcohol. Very easily sol. in hydracids and dil. HNO_3 . (Tassily, C. R. 1894, 119. 372.)

Calcium oxychloride, $\text{Ca}_2\text{O}_2\text{Cl}_2 + 15\text{H}_2\text{O} = 3\text{CaO}$, $\text{CaCl}_2 + 15\text{H}_2\text{O}$.

Decomp. by H_2O or alcohol. (Rose.)

Formula is $\text{Ca}_2\text{HO}_2\text{Cl} + 7\text{H}_2\text{O}$. (Grimshaw, C. N. 30. 280.)

+16 H_2O . Decomp. by H_2O into CaO_2H_2 and CaCl_2 until a maximum of 85 g. CaCl_2 are dissolved per litre. (Ditte, C. R. 91. 576.)

4 CaO , $\text{CaCl}_2 + 14\text{H}_2\text{O}$. (Schreinemakers and Figeo, Chem. Weekbl. 1911, 8. 685.)

CaO , CaCl_2 . (Schreinemakers and Figeo.)

Calcium lead oxychloride, CaCl_2 , CaO , 2 $\text{PbO} + 4\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (André, C. R. 104. 359.)

CaCl_2 , 3 $\text{PbO} + 3\text{H}_2\text{O}$. (André.)

Calcium mercuric oxychloride, CaCl_2 , 2 $\text{HgO} + 4\text{H}_2\text{O}$.

Decomp. immediately by H_2O . (Klinger, B. 16. 997.)

oxyiodide, 3CaO , $\text{CaI}_2 + 16\text{H}_2\text{O}$.

p. by H_2O , alcohol, and acids. Sol. in dil. HNO_3 . (Tassily, *Bull. Soc. Chim.* 1904, 119. 372.)

oxysulphide, $\text{Ca}_4\text{O}_3\text{S}_4 + 12\text{H}_2\text{O} = 4\text{CaO}$, $\text{CaS}_4 + 12\text{H}_2\text{O}$.

p. by H_2O . Not acted on by alcohol. (Schöne, *Pogg.* 117. 77.) According to Geuther (*A.* 224. 178) $= \text{CaS}_2$, or $11\text{H}_2\text{O}$. Sol. in dil. $\text{HCl} + \text{Aq}$ with evolution of S.

$4 + 18\text{H}_2\text{O} = 4\text{CaO}$, $\text{CaS}_4 + 18\text{H}_2\text{O}$.

p. by H_2O , but not acted on by alcohol. (Schöne, *Pogg.* 117. 82.)

According to Geuther (*A.* 224. 178) $= \text{CaS}_2$, or $15\text{H}_2\text{O}$.

$4 + 20\text{H}_2\text{O} = 5\text{CaO}$, $\text{CaS}_4 + 20\text{H}_2\text{O}$.

(*Pogg.* 55. 433.)

400 pts. cold, decomp. by boiling water; sl. sol. in cold, much more in hot, but it is not deposited on cooling. Aqueous solution sat. at $6^\circ - 7.2^\circ$ has sp. gr. = 1.01 (Mierschell); sol. in alcohol (Gay-Lussac). Insol. in alcohol (Gmelin).

phosphide, CaP .

Decomp. in moist air or water. Not attacked by conc. HNO_3 , but by dil. $\text{HNO}_3 + \text{Aq}$. (Thénard, *A.* 14.)

Crystallized.

p. by H_2O .

Not attacked by conc. H_2SO_4 . Violently attacked by dil. H_2SO_4 .

Not attacked by abs. alcohol, ether, benzene, or oil of turpentine. (Moissan, *C. R.* 1892, 792.)

Insol. in liquid CO_2 . (Büchner, *Ch.* 1906, 54. 674.)

selenide, CaSe .

Insol. in H_2O . Very easily decomp. (R. 102. 1469.)

silicide, CaSi_2 .

decomp. by H_2O ; sol. in conc. H_2SO_4 , HNO_3 with evolution of H_2 . With HCl it gives H_2 , Si and silicon hydride; with HCl , H_2 and a yellow substance. (Moissan, *C. R.* 1902, 134. 505.)

Modifications:

1. Sol. in HNO_3 ; decomp. H_2O to sol. ppt. on addition of HCl .

2. Sol. in HNO_3 and acetic acid; with HCl to give a ppt. which is sol. in H_2O . (de Chalmot, *Am. Ch. J.* 1896, 14. 104.)

Slowly decomp. by H_2O , rapidly by acetic acid or by $\text{H}_2\text{SO}_4 + \text{Aq}$ without evolution of spontaneously inflammable gas. (Mierschell, *M.* 1909, 30. 497.)

3. p. by dil. min. acids, with evolution

of spontaneously inflammable gas. (Hackspill, *Bull. Soc.* 1908, (4) 3. 619.)

$\text{Ca}_2\text{Si}_{10}$. Insol. in all solvents. Decomp. by boiling H_2O , by conc. HCl and by acetic acid. Sol. in dil. alkali and alkali carbonates + Aq . Hardly attacked by conc. H_2SO_4 or HNO_3 . (Kolb, *Z. anorg.* 1909, 64. 349.)

$\text{Ca}_{11}\text{Si}_{10}$. Easily decomp. by boiling with H_2O . Decomp. by dil. acetic acid, dil. or conc. HCl . (Kolb, *Z. anorg.* 1909, 64. 349 and 356.)

Calcium siliconitride, CaSi_2N_2 .

(Kolb, *Z. anorg.* 1909, 64. 363.)

$\text{Ca}_2\text{Si}_2\text{N}_4$. Slowly decomp. by boiling with H_2O , somewhat more rapidly with dil. $\text{NaOH} + \text{Aq}$. Slowly decomp. by conc. HCl . (Kolb, *l. c.*)

$\text{Ca}_{11}\text{Si}_{10}\text{N}_{10}$. Completely decomp. by HCl . (Kolb, *l. c.*)

Calcium sulphide, CaS .

500 pts. H_2O dissolve 1 pt. CaS completely; less H_2O dissolves out CaS_2H_2 and leaves CaO_2H_2 . Very much H_2O decomposes completely into CaO_2H_2 and H_2S . (Béchamp, *A. ch.* (4) 16. 222.)

Not decomp. by H_2O , and only sl. sol. therein at ordinary temp. (Pelouze.)

After 48 hours contact with CaS , 1 l. H_2O contains at:

10°	18°	40°	60°	90°
0.15	0.23	0.30	0.48	0.33 g. CaS .

After boiling for 2 hours, 0.27 g. CaS is dissolved; addition of NaCl diminishes solubility, but Na_2SO_4 increases it. Lime-water dissolves at 14° 0.18 g. CaS , the same amount which H_2O dissolves at 60° . Milk of lime dissolves 0.55 g. at 60° . H_2O containing 3 to 79 g. Na_2O per litre dissolves only traces of CaS at 10° , but at $40-60^\circ$, or by boiling, a large amount of Na_2S is formed. (Kolb, *A. ch.* (4) 7. 126.)

Sol. in 12,500 pts. H_2O at 12.6° . (Scheurer-Kestner, *Répert. chim. appl.* 1862. 331.)

Sat. $\text{Na}_2\text{CO}_3 + \text{Aq}$ has scarcely any action on CaS , but a dilute solution has more action. (Kolb.)

Sol. in H_2O and sulphur, forming CaS_4 .

Insol. in liquid NH_3 . (Franklin, *Am. Ch. J.* 1898, 20. 827.)

Insol. in methyl acetate. (Naumann, *B.* 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, *B.* 1904, 37. 3601.)

Insol. in methylal. (Eidmann, *C. C.* 1899, II. 1014.)

Sol. in 10 pts. glycerine. (Cap and Garot, *J. Pharm.* (3) 26. 81.)

Sol. in sugar + Aq . (Stolle, *C. C.* 1900, I. 1044.)

Calcium tetrasulphide, CaS_4 .

Known only in solution.

Calcium pentasulphide, CaS_5 .
Sol. in H_2O and alcohol. (Berzelius.)
Exists only in aqueous solution. (Schöne, Pogg. 117. 73.)

Calcium hydroxyl sulphide, $\text{Ca(OH)SH} + 3\text{H}_2\text{O}$.
Easily sol. in H_2O with immediate decomp. and separation of Ca(OH)_2 . Insol. in alcohol, but slowly decomp. thereby. (Divers and Shimidzu, Chem. Soc. 45. 270.)

Calcium stannic sulphide.
See Sulphostannate, calcium.

Calomel.
See Mercurous chloride.

Carbamic acid.

Ammonium carbamate acid carbonate (commercial carbonate of ammonia).
See Carbonate carbamate, ammonium hydrogen.
— (salts of hartshorn), $2\text{NH}_4\text{HCO}_3$, NH_4CONH_2 .
See Carbonate carbamate, ammonium hydrogen.

Carbazote silicon, C_2SiN .
Insol. in acids, even HF ; also in boiling $\text{KOH} + \text{Aq.}$ (Schützenberger and Colson, C. R. 92. 1508.)

Carbon, C.
Insol. in all solvents.
Diamond is unacted upon by $\text{KClO}_3 + \text{fum. HNO}_3$; graphite forms graphitic acid by $\text{KClO}_3 + \text{fum. HNO}_3$; amorphous carbon is sol. in $\text{KClO}_3 + \text{fum. HNO}_3$. (Berthelot, A. ch. (4) 19. 399.)
Diamond is sol. in molten iron at 1160° . Amorphous carbon is insol. in molten iron at 1160° , but becomes sol. therein by heating to 1400° . (Hempel, B. 18. 998.)
Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54. 674.)
Charcoal is insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.)
The quantity of carbon dissolved by iron diminishes by increasing phosphorus, falling by about 0.5% for each additional 2.0% of phosphorus. (Fettweis, Metallurgie, 1906, 3. 60.)
Solubility in iron is reduced by the presence of tin and of sulphur. (Wüst, Metallurgie, 1906, 3. 169.)
The solubility of C in iron is increased by the presence of chromium; 9.2% C dissolved when 62% Cr is present in the mixture. (Goerens, Metallurgie, 1907, 4. 18.)

Carbon boride, CB_4 .
Insol. in boiling $\text{HNO}_3 + \text{Aq.}$ (Joly, C. R. 97. 456.)

Carbon suboxide, C_3O_2 .
B.-pt. $+7^\circ$ at 761 mm.
Sol. in H_2O with formation of malonic acid. Slowly decomp. on standing in a sealed tube. (Diels, B. 1906, 39. 696.)

Carbon monoxide, CO.
Sol. in 50 vols. recently boiled H_2O . (Davy.)
Sol. in 16 vols. H_2O . (de Saussure.)
Sol. in 27 vols. H_2O . (Dalton.)
100 vols. H_2O dissolve 6.2 vols. CO at 18° . (de Saussure.)

Solubility of CO in H_2O : 1 vol. H_2O at t° dissolves V vols. CO reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	0.03287	7	0.02796	14	0.02406
1	0.03207	8	0.02739	15	0.02432
2	0.03131	9	0.02686	16	0.02402
3	0.03057	10	0.02635	17	0.02374
4	0.02987	11	0.02588	18	0.02350
5	0.02920	12	0.02544	19	0.02329
6	0.02857	13	0.02504	20	0.02312

(Bunsen's Gasometry, pp. 287, 128, 146.)

Coefficient of absorption = $0.032874 - 0.00081632t + 0.000016421t^2$. (Bunsen and Pauli, A. 93. 16.)

Solubility of CO in H_2O .
 β = Vol. CO absorbed by 1 vol. H_2O at a partial pressure of 760 mm.
 β^1 = Vol. CO (reduced to 0° and 760 mm.) absorbed by 1 vol. of H_2O under a total pressure of 760 mm.
 q = g. CO dissolved by 100 g. H_2O at a total pressure of 760 mm.

t°	β	β^1	q
0	0.03537	0.03516	0.0044
5	0.03149	0.03122	0.0039
10	0.02816	0.02782	0.0035
15	0.02543	0.02501	0.0031
20	0.02319	0.02266	0.0028
25	0.02142	0.02076	0.0026
30	0.01998	0.01915	0.0024
40	0.01775	0.01647	0.0021
50	0.01615	0.01420	0.0018
60	0.01488	0.01197	0.0015
70	0.01440	0.00998	0.0013
80	0.01430	0.00762	0.0010
90	0.01420	0.00438	0.0006
100	0.01410	0.00000	0.0000

(Winkler, B. 1901, 34. 1416.)

solubility in H₂O at various pressures.
Volume of the absorbing liquid.
Fig pressure in metres.
Coefficient of solubility.

	t°	P	λ
ccm.	17.7	0.9202	0.02791
		1.1438	0.02787
		1.4624	0.02786
		1.7986	0.02783
		2.3659	0.02782
		2.8390	0.02776
		3.2622	0.02771
		4.0114	0.02770
		4.6017	0.02763
		5.1953	0.02761
		5.8717	0.02756
		6.5462	0.02744
		7.0983	0.02738
ccm.	19.0	8.0184	0.02715
		0.9176	0.02716
		1.1506	0.02717
		1.3897	0.02715
		1.7044	0.02712
		2.1239	0.02708
		2.7173	0.02701
		3.2576	0.02693
		3.9311	0.02689
		4.4584	0.02680
		5.2470	0.02673
		6.0346	0.02665
		6.6303	0.02654
		7.1842	0.02636
		7.9542	0.02617

Massuto, Phys. Zeit. 1904, 5. 236.)

coefficient of absorption of CO in H₂O at various pressures equals 0.0154. (Findlay and Creighton, Trans. Am. J. 1911, 5. 294.)
Cuprous chloride in an hydrochloric acid or ammoniacal solution, and ammoniacal solution of cuprous salts absorb large amounts of CO. (Leblanc, C. R. 30. 488.)
Cuprous chloride dissolved in HCl + Aq absorbs 15-20 vols. CO. (Berthelot, A. ch. (3) 61. 463.)
CO is absorbed by KOH, NaOH, Ba(OH)₂, and Ca(OH)₂ + Aq; more readily by ether, alcohol, wood spirit, with formation of formic acid in HCN. (Böttinger, B. 10. 1122.)
Ethyl alcohol absorbs 0.20443 vols. CO at all temperatures between 0° and 25°. (de Saussure, 1814, A. 94. 135.)

100 vols. alcohol (0.84 sp. gr.) dissolve 14.5 vols. CO at 18°; 100 vols. rectified naphtha (0.784 sp. gr.), 20.0 vols. CO at 18°; 100 vols. oil of lavender (0.88 sp. gr.), 15.2 vols. CO at 18°; 100 vols. olive oil (0.915 sp. gr.), 15.2 vols. CO at 18°; 100 vols. sat. KCl + Aq (1.168 sp. gr.), 5.2 vols. CO at 18°. (de Saussure, 1814.)
Oil of turpentine absorbs 0.16-0.20 vol. CO. (de Saussure.)
CO in ether. (Regnault.)
CO in caoutchine.

Solubility in alcohol + Aq.				
% alcohol by weight	0.00	9.09	16.67	23.08
Solubility	2.41	1.87	1.75	1.68
% alcohol by weight	28.57	33.33	50.00	
Solubility	1.50	1.94	3.20	
(Lubarsch, W. Ann. 1889, 37. 524.)				

Solubility of CO in organic solvents.		
Solvent	Solubility at 20° C.	Solubility at 25° C.
Glycerine	Not measurable	
Water	0.02404	0.02586
Aniline	0.05358	0.05055
Carbon bisulphide	0.08314	0.08112
Nitrobenzene	0.09366	0.09105
Benzene	0.1707	0.1645
Glacial acetic acid	0.1714	0.1689
Amyl alcohol	0.1714	0.1706
Xylene	0.1781	0.1744
Toluene	0.1808	0.1742
Ethyl alcohol (99.8%)	0.1921	0.1901
Chloroform	0.1954	0.1897
Methyl alcohol	0.1955	0.1830
Amyl acetate	0.2140	0.2108
Acetone	0.2225	0.2128
Isobutyl acetate	0.2365	0.2314
Ethyl acetate	0.2516	0.2419

(Just, Z. phys. Ch. 1901, 37. 361.)
Solubility of CO in ether at 0° = 0.3618, and at 10° = 0.3842. (Christoff, Z. phys. Ch. 1912, 79. 459.)

Solubility of CO in organic mixtures. CO in benzene and naphthalene at 25° C.		
Per cent by weight of naphthalene	Per cent by weight of benzene	* Solubility of CO
0	100	0.174
11.52	88.48	0.164
11.65	88.35	0.163
23.98	76.02	0.149
23.60	76.40	0.148
32.35	67.65	0.142
32.74	67.26	0.143
33.79	66.21	0.141

(Skirrow, Z. phys. Ch. 1902, 41. 144.)
* See under Oxygen.

CO in benzene and phenanthrene at 25° C.		
Per cent by weight of phenanthrene	Per cent by weight of benzene	Solubility of CO
0	100	0.174
10.48	89.52	0.144
10.48	89.52	0.144
19.22	80.78	0.132
18.99	81.01	0.133
27.04	72.96	0.128
27.39	72.61	0.127

(Skirrow.)

CO in benzene and α -naphthol at 25° C.		
Per cent by weight of α -naphthol	Per cent by weight of benzene	Solubility of CO
0	100	0.174
3.48	96.52	0.149
6.75	93.25	0.145
6.59	93.41	0.144
12.10	87.90	0.139
11.81	88.19	0.139

(Skirrow.)

CO in benzene and β -naphthol at 25° C.		
Per cent by weight of β naphthol	Per cent by weight of benzene	Solubility of CO
0	100	0.174
2.06	97.94	0.158
4.14	95.86	0.151
4.36	95.64	0.149

(Skirrow.)

CO in benzene and nitrobenzene at 25° C.		
Per cent by weight of nitrobenzene	Per cent by weight of benzene	Solubility of CO
0	100	0.174
14.5	85.5	0.162
14.12	85.88	0.162
28.18	71.82	0.152
28.14	71.86	0.152
40.58	59.42	0.140
40.63	59.37	0.140
54.9	45.1	0.126
54.9	45.1	0.127
83.33	16.67	0.101
83.2	16.8	0.102
100	0	0.093

(Skirrow.)

CO in benzene and aniline at 25° C.		
Per cent by weight of aniline	Per cent by weight of benzene	Solubility of CO
0	100	0.174
12.69	87.31	0.156
12.03	87.97	0.158
19.57	80.43	0.145
19.43	80.57	0.144
28.43	71.57	0.131
28.26	71.74	0.131
57.68	42.32	0.0945
57.38	42.62	0.0953
78.90	21.10	0.0689
78.80	21.20	0.0684
100	0	0.053

(Skirrow.)

CO in toluene and naphthalene at 25° C.		
Per cent by weight of naphthalene	Per cent by weight of toluene	Solubility of CO
0	100	0.182
7.13	92.87	0.169
7.10	92.9	0.171
15.10	84.9	0.161
15.13	84.87	0.161
22.75	77.25	0.153
22.58	77.42	0.154

(Skirrow.)

CO in toluene and phenanthrene at 25° C.		
Per cent by weight of phenanthrene	Per cent by weight of toluene	Solubility of CO
0	100	0.182
5.59	94.41	0.170
5.58	94.42	0.171
11.16	88.84	0.161
11.20	88.8	0.161
21.62	78.38	0.147
21.93	78.07	0.147

(Skirrow.)

CO in toluene and nitrobenzene at 25° C.		
Per cent by weight of nitrobenzene	Per cent by weight of toluene	Solubility of CO
0	100	0.182
8.86	91.14	0.168
8.87	91.13	0.168
18.27	81.73	0.160
18.19	81.81	0.161
26.82	73.18	0.151
26.76	73.24	0.151
49.14	50.86	0.131
49.02	50.98	0.131
76.31	23.69	0.108
76.31	23.69	0.108
100	0	0.093

(Skirrow.)

CO in toluene and aniline at 25° C.		
Per cent by weight of aniline	Per cent by weight of toluene	Solubility of CO
0	100	0.182
6.61	93.39	0.169
6.61	93.39	0.168
13.56	86.44	0.157
13.55	86.45	0.156
19.91	80.09	0.148
19.96	80.04	0.148
44.64	55.36	0.115
44.31	55.69	0.116
74.63	25.37	0.0768
75.03	24.97	0.0753
100	0	0.053

(Skirrow.)

n toluene and α-naphthol at 25° C.

Per cent by weight of α-naphthol	Per cent by weight of toluene	Solubility of CO
	100	0.182
46	95.54	0.171
44	95.56	0.171
75	91.25	0.162
89	91.11	0.163

(Skirrow.)

n acetone and naphthalene at 25° C.

Per cent by weight of naphthalene	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
	100	229.6	0.238
31	86.69	212.4	0.199
40	72.60	196.6	0.187

(Skirrow.)

n acetone and phenanthrene at 25°C.

Per cent by weight of phenanthrene	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
	100	229.6	0.238
77	87.23	218	0.205
04	74.96	207.5	0.183

(Skirrow.)

n acetone and β-naphthol at 25° C.

Per cent by weight of β-naphthol	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
	100	229.6	0.238
95	86.05	213	0.190
88	73.12	195	0.169

(Skirrow.)

n acetone and nitrobenzene at 25° C.

Per cent by weight of nitrobenzene	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
	100	229.6	0.238
.59	78.4	201	0.207
.20	46.8	152	0.157
	0	...	0.093

(Skirrow.)

CO in acetone and aniline at 25° C.

Per cent by weight of aniline	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
	100	229.6	0.238
.83	79.17	192	0.179
.10	44.9	120	0.110
	0	...	0.053

(Skirrow.)

CO in acetic acid and nitrobenzene at 25° C.

Per cent by weight of nitrobenzene	Per cent by weight of acetic acid	Solubility of CO
0	100	0.173
21.65	78.35	0.156
51.03	48.97	0.130
100	0	0.093

(Skirrow.)

CO in acetic acid and aniline at 25° C.

Per cent by weight of aniline	Per cent by weight of acetic acid	Solubility of CO
0	100	0.173
13.5	86.5	0.110
41.64	58.36	0.0699
60.77	39.23	0.0618
82.21	17.79	0.0580
100	0	0.053

(Skirrow.)

CO in methyl alcohol and glycerine at 25°C.

Per cent by weight of glycerine	Per cent by weight of methyl alcohol	Measured vapor pressure	Solubility of CO
0	100	122	0.196
39.6	60.4	106	0.0964
60.5	39.5	91	0.0515
77.1	22.9	63	0.0246
100	0	...	very small

(Skirrow.)

CO in acetone and chloroform at 25° C.

Per cent by weight of chloroform	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
0	100	229.6	0.238
33.38	66.62	202	0.226
53.2	46.8	179	0.219
65.03	34.97	167	0.220
73.46	26.54	162	0.212
79.83	20.17	163	0.204
87.3	12.7	168	0.207
94.4	5.6	178	0.205
100	0	188	0.207

(Skirrow.)

CO in acetone and carbon bisulphide at 25° C.

Per cent by weight of carbon bisulphide	Per cent by weight of acetone	Measured vapor pressure	Solubility of CO
0	100	229.6	0.238
8.18	91.82	306	0.236
18.02	81.98	367	0.236
49.46	50.54	443	0.227
62.6	37.4	457	0.210
74.05	25.95	457	0.187
85.51	14.49	433	0.144
96.42	3.58	382	0.114
100	0	356	0.0959

(Skirrow.)

CO in benzene and ethyl alcohol at 25° C.

Per cent by weight of acetic acid	Per cent by weight of chloroform	Measured vapor pressure	Solubility of CO
0	100	95.9	0.174
15.43	84.56	125	0.179
52.34	47.66	119	0.181
100	0	59	0.192

(Skirrow.)

CO in chloroform and methyl alcohol at 25° C.

Per cent by weight of alcohol	Per cent by weight of chloroform	Measured vapor pressure	Solubility of CO
0	100	188	0.207
13	87	233	0.202
100	0	122	0.196

(Skirrow.)

CO in acetic acid and benzene at 25° C.

Percent by weight of acetic acid	Percent by weight of benzene	Measured vapor pressure	Solubility of CO
0	100	95.9	0.174
19.17	80.83	87.5	0.190
33.54	66.46	82	0.198
67.51	32.49	64.5	0.199
100	0	14	0.172

(Skirrow.)

CO in acetic acid and toluene at 25° C.

Per cent by weight of acetic acid	Per cent by weight of toluene	Measured vapor pressure	Solubility of CO
0	100	9	0.182
20.48	79.52	31.6	0.190
56.89	43.11	28	0.195
74.71	25.29	25.6	0.191
100	0	14	0.172

(Skirrow.)

CO in acetic acid and chloroform at 25° C.

Per cent by weight of acetic acid	Per cent by weight of chloroform	Measured vapor pressure	Solubility of CO
0	100	188	0.206
26.67	73.33	144.5	0.207
56.46	43.54	88.5	0.196
100	0	14	0.172

(Skirrow.)

CO in carbon bisulphide and ethylene dichloride at 25° C.

Per cent by volume of carbon bisulphide	Per cent by volume of ethylene dichloride	Measured vapor pressure	Solubility of CO
0	100	77	0.147
25	75	231	0.159
49	51	294	0.160
81.6	18.4	338	0.140
100	0	356.5	0.083

(Skirrow.)

Coefficient of absorption for petroleum = 0.123 at 20°, and 0.134 at 10°. (Gniewas and Walfisz, Zeit. phys. Ch. 1. 70.)

Carbon dioxide, CO₂.

Gas.—

H₂O dissolves about its own vol. CO₂ at the ordinary temperature (the solution obtained being of 1.0018 sp. gr.) and pressure, and an additional vol. for the pressure of each additional atmosphere to which it is subjected.

The power of H₂O to absorb CO₂ does not increase in precisely the same ratio as the pressure. (Soubeiran.)

5 vols. CO₂ dissolve in 1 vol. H₂O at 7 atmos. pressure, and much greater pressure is necessary in order to increase the amount of gas dissolved; but up to 4 or 5 atmospheres the amount of gas dissolved is very nearly proportional to the pressure. (Courbe, J. Pharm. 24. 121.)

100 vols. H₂O at 12.78° absorb 116 vols. CO₂ (Cavendish); at 29.44°, 84 vols. CO₂ (Henry); at 15.56°, 106 vols. CO₂ (Saussure); at 15.56°, 106 vols. CO₂ (Henry); at 15.56°, 100 vols. CO₂ (Dalton).

100 vols. H₂O at t° C. absorb V vols. of CO₂ gas reduced to 60° F. and 30 in. pressure.

t°	V	t°	V
0	175.72	26.7	68.60
4.4	147.94	32.2	57.50
10	122.27	37.8	50.39
15.6	100.50	65.6	11.40
21.1	83.86	100	trace

(Rogers, Am. J. Sci. (2) 6. 107.)

1 vol. H₂O at 5° absorbs somewhat more than 1 vol. CO₂; at 10° scarcely 1 vol., and still less at higher temp. CO₂ + Aq sat. at 2° has 1.0015 sp. gr.; most of the CO₂ escapes upon exposing the solution to the air, the more quickly the higher the temperature. But as CO₂ diminishes, the remainder is more obstinately held, so that boiling for ½ hour is necessary to expel it completely. (Bergman.)

Solubility of CO₂ in H₂O. 1 vol. H₂O at t° and 760 mm. dissolves V vols. CO₂ gas reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	1.7967	7	1.3339	14	1.0321
1	1.7207	8	1.2809	15	1.0020
2	1.6481	9	1.2311	16	0.9753
3	1.5787	10	1.1847	17	0.9519
4	1.5126	11	1.1416	18	0.9318
5	1.4497	12	1.1018	19	0.9150
6	1.3901	13	1.0653	20	0.9014

(Bunsen's Gasometry, pp. 287, 128, 152.)

Coefficient of absorption = 1.7967—0.07761t + 0.001642t². (Bunsen.)

bility in H₂O at various pressures: P = pressure in atmospheres.

Vol. gas. in 1 ccm. H ₂ O		P	Vol. gas in 1 ccm. H ₂ O	
at 0°	at 12.43°		at 0°	at 12.43°
1.797	1.086	20	26.65	17.11
8.65	5.15	25	30.55	20.31
16.03	9.65	30	33.74	23.35
21.95	13.63	..	—	...

(Wroblewski, C. R. 94. 1355.)

orption of CO₂ in H₂O at various pressures: P = pressure in mm.; V = vols. CO₂, reduced to 0° and 760° mm., absorbed by vol. H₂O.

P	V	P	V
7.71	0.9441	2188.65	3.1764
9.03	1.1619	2369.02	3.4857
9.41	1.8647	2554.00	3.7152
9.95	2.1623	2738.33	4.0031
2.06	2.9067	3109.51	4.5006

hanikoff and Longuinine, A. ch. (4) 11. 412.)

C = coefficient of absorption in H₂O at t° and 760 mm.

t°	C	t°	C	t°	C
5.2	1.009	18.38	0.896	21	0.838
7.6	0.930	18.3	0.885	23	0.798

Setchenow, Mém. Acad. St. Petersb. 22. Nos. 6, 7.)

Absorption coefficient of CO₂ in H₂O at 0° = 1.7306. (Prytz and Holst, W. Ann. 1895, 136.)

Absorption of CO₂ by H₂O at t°. a = coefficient of absorption.

t°	a	t°	a
0	1.713	19	0.902
1	1.646	20	0.878
2	1.584	21	0.854
3	1.527	22	0.829
4	1.473	23	0.804
5	1.424	24	0.781
6	1.377	25	0.759
7	1.331	26	0.738
8	1.282	27	0.718
9	1.237	28	0.699
10	1.194	29	0.682
11	1.154	30	0.665
12	1.117	35	0.592
13	1.083	40	0.530
14	1.050	45	0.479
15	1.019	50	0.436
16	0.985	55	0.394
17	0.956	60	0.359
18	0.928

(Bohr, W. Ann. 1899, 68. 504.)

Solubility in H₂O at 25° = 0.8255; at 15° = 1.070. (Geffcken, Z. phys. Ch. 1904, 49. 273.)

75 cc. H₂O absorb 0.1381 g. CO₂ at 15.5° and 720 mm. (Christoff, Z. phys. Ch. 1905, 53. 329.)

Absorption-coefficient of CO₂ in H₂O at 20° = 0.877, or 1000 g. H₂O dissolve 878 cc. CO₂. (Usher, Chem. Soc. 1910, 97. 72.)

Solubility of CO₂ in H₂O = 1.158 at 12° and 0.825 at 25°. (Findlay and Shenn, Chem. Soc. 1911, 99. 1315.)

Absorption of CO₂ by H₂O at high pressure.

Amount of H₂O used { a = 0.210 ccm.
b = 0.102 ccm.

V = ccm. of CO₂ absorbed by H₂O at t°, reduced to a pressure of 1 kg. per sq. cm.

V₁ = ccm. of CO₂ absorbed by 1 ccm. of H₂O.

Pressure kg/sq. cm.	t°	V ₁	
		a	b
25	20°	17.77
30		19.77
40		21.52
50		28.09
55		29.75
30	35°	11.77	13.57
40		14.82	20.00
50		18.96	24.64
60		22.90	22.50
70		27.18	27.62
80		32.85
40	60°	10.88	9.798
50		12.24	13.72
60		14.46	15.28
70		16.80	17.46
80		19.74	22.67
90		22.74	21.16
100		26.21	27.85
110		28.92	28.79
120		30.20	33.90
60	100°	8.965
70		10.11	6.395
80		11.05	9.591
90		12.63	10.85
100		13.63	12.40
110		14.88	16.31
120		16.40	15.78
130		17.93	16.89
140		19.56	17.71
150		20.58	17.49
160		22.07
170		22.78

(Sander, Z. phys. Ch. 1912, 78. 537.)

Solubility of carbon dioxide in water at 25°.

P. = Pressure in mm. Hg.

S. = Solubility calculated according to formula for which see the original article. (Findlay, Chem. Soc. 1910, 97. 538.)

P	S	P	S
743	0.816	1059	0.817
752	0.817	1064	0.819
800	0.815	1153	0.818
841	0.817	1243	0.819
955	0.816	1351	0.820
955	0.817	1351	0.820

(Findlay and Creighton, Chem. Soc. 1910, 97. 538.)

Solubility of carbon dioxide in water at 25°.

P. = Pressure in mm. Hg.

S. = Solubility. See above.

P	S	P	S
755	0.826	1069	0.823
759	0.825	1084	0.825
836	0.825	1210	0.825
841	0.826	1211	0.825
927	0.826	1350	0.824
934	0.824	1350	0.826

(Findlay and Creighton, Chem. Soc. 1912, 101. 1460.)

Solubility of carbon dioxide in water at 25°.

P = Pressure in mm. Hg.

S = Solubility. See above.

P	S	P	S
263	0.817	495	0.816
271	0.816	651	0.816
382	0.814	667	0.817
392	0.811	752	0.818
479	0.816	768	0.817

(Findlay and Creighton, Chem. Soc. 1913, 103. 638.)

Sl. sol. in HCl + Aq.

100 vols. H₂SO₄ of 1.840 sp. gr. absorb 45 vols. CO₂. (de Saussure.)

H₂SO₄ of ordinary density at 15.56° and common pressure absorbs 94% of its vol. of CO₂; fuming H₂SO₄, 125%; the absorption for pure H₂O under the same conditions being 98%. (Rogers, Am. J. Sci. (2) 8. 115.)

H₂SO₄ absorbs 7-10% CO₂. (Hlasiwetz, W. A. B. 20. 193.)

Coefficient of absorption by conc. H₂SO₄ = 0.932, which is the same as that by H₂O; but this diminishes on diluting, and is at its lowest limit 0.666, when the composition of the solution is H₂SO₄, H₂O; upon further dilution the coefficient of solubility gradually increases, and when 58 H₂O are present to 1 H₂SO₄, the

coefficient of absorption is 0.857. (Schenow, J. B. 1876. 46.)

Absorption of CO₂ by H₂SO₄ + Aq.

Solution	Grams CO ₂ absorbed by 75 cc. at 15.5° and 720 mm.
1/2-N H ₂ SO ₄	0.1273
1-N H ₂ SO ₄	0.1179
2-N H ₂ SO ₄	0.1092
4-N H ₂ SO ₄	0.1003

(Christoff, Z. phys. Ch. 1905, 53. 329.)

Solution	Grams CO ₂ absorbed by 75 cc. at 15.5° and 720 mm.
2.5% H ₂ SO ₄	0.1282
5% "	0.1179
10% "	0.0833
20% "	0.0755
30% "	0.0751
40% "	0.0713
45% "	0.0725
70% "	0.0918
90% "	0.1433

(Christoff, l. c.)

Coefficient of absorption for 96% H₂SO₄, 0.926 at 20.2°. (Bohr, Z. phys. Ch. 1910, 71. 48.)

Absorption of CO₂ by acids.

M = Content in gram-equivalents per liter
S = Solubility (see under oxygen).

Absorption of CO₂ by HNO₃ + Aq.

M	S _{25°}	S _{15°}
0.472	0.8382	1.073
0.475	0.8366	1.075
0.557	0.8387	1.069
0.704	0.8447	1.060
1.382	0.8620	1.093
1.387	0.8622	1.093
1.860	0.8752	1.105
2.519	0.8839	1.109
2.539	0.8865	1.111

(Geffcken, Z. phys. Ch. 1904, 49. 273.)

Absorption of CO by HCl + Aq.

M	S _{25°}	S _{15°}
0.499	0.8047	1.041
0.511	0.8074	1.042
1.212	0.7973	1.020
1.249	0.7984	1.023
2.080	0.7951	0.9864
2.180	0.7951	1.009

(Geffcken.)

absorption of CO₂ by $\frac{H_2SO_4}{2} + Aq.$

	S_{25}°	S_{15}°
12	0.7723	1.016
17	0.7936	1.016
35	0.7693	0.9772
39	0.7685	0.9775
37	0.7672	0.9756
56	0.7302	0.9175
58	0.7273	0.9143
90	0.6736	0.8354
10	0.6747	0.8385

(Geffcken.)

eting CO₂ gas in pneumatic operations, a solution of common salt is better than H₂O the trough. This solution will only absorb of the amount of CO₂ absorbed by pure H₂O. (re, l. c.)

of the following solutions at 18° and ordinary absorb vols. CO₂—

	Sp. gr.	Vols. CO ₂
+Aq (containing 29% of NaCl)	1.212	32.9
+Aq (containing 27.53% of NaCl)	1.078	76
+Aq (containing 26% of KCl)	1.168	61
+Aq (containing 40.2% of NaCl)	1.402	26.1
+Aq (containing 9.42% of NaCl)	1.077	62
+Aq (containing 11.14% of NaCl)	1.105	58
(SO ₄) ₂ + Aq (containing 9.14% of NaCl)	1.047	70
+Aq (containing 20.6% of NaCl)	1.139	57
O ₂ + Aq (containing 26.4% of NaCl)	1.206	45
H ₂ O ₂ + Aq (containing 53.37% of NaCl)	1.288	41

causure. Gilbert's Ann. Phys. 47. 167)

half as sol. in NaCl + Aq (15% in H₂O.

more sol. in Na₂HPO₄ + Aq or +Aq than in H₂O, the quantity decreasing with the amount of salt in ion. The solubility in these solutions on the coefficient of solubility in H₂O product of a constant coefficient d by the amount of salt in the solution; constant equals 0.069 for Na₂HPO₄, 3 for Na₂CO₃. (Fernet, A. ch. (3) 47.

's determinations are not accurate. (er, A. Suppl. 2. 157.)

. Na₂HPO₄ in dil. Na₂HPO₄ + Aq 2 mols. CO₂. (Settschenow.)

ms of salts of similar constitution are it in regard to their power of absorption, when they contain the same per- of crystal water. Experiments were th solutions of alum, MgSO₄, 7H₂O, IO₄, 7H₂O, containing 10% of the the MgSO₄ solution absorbed the proportional amount of CO₂, and the least. The further rule was de- at with salts of similar constitution same amount of crystal water, the

absorptiometric equivalents are identical with the chemical equivalents. (Settschenow, B. 8. 1461.)

Salts can be divided into two classes, according as CO₂ has chemical action on the salt or not. In the first case, i. e., when there is chemical combination or action of CO₂ on the salt in solution, the amount of CO₂ absorbed increases with increasing concentration of the solution; in the second case, however, the amount of CO₂ decreases with the strength of the solution. Several salts can be arranged in a series as regards their power of absorption, beginning with that which has the greatest, as follows: Na₂CO₃, Na₂B₄O₇, Na₂HPO₄, NaC₂H₃O₂, Na₂C₂H₃O₇, Na₂C₂O₄, NaC₂H₃O₂, MnO₂, MCl, M₂SO₄. The division between the two classes occurs in this series at Na₂C₂O₄.

The matter is discussed at length in the original papers. (Settschenow, Mémoires Acad. St. Petersb. 22. No. 8. Also further, Settschenow, ib. 34. No. 3, and 35. No. 7. See also Ostwald, Allgemeine Chemie, 2^{te} Aufl. vol. 1, p. 629.)

Solubility of CO₂ in salts + Aq at 15.2°.

CO₂ = cc. CO₂ (at 0° and 760 mm.) dissolved per cc. of salt solution.

Salt	g. salt per l.	CO ₂
NH ₄ Cl	1	1.005
"	10	0.985
"	51.6	0.941
"	172	0.819
"	258	0.770
NH ₄ NO ₃	2.8	1.013
"	11.2	1.002
"	55	0.989
"	101	0.962
"	202.1	0.911
"	404.3	0.807
"	810.4	0.612
(NH ₄) ₂ SO ₄	72.2	0.712
"	144.4	0.575
Ba(NO ₃) ₂	62.7	0.922
Ca(NO ₃) ₂	41	0.923
LiCl	16.72	1.035
"	50.15	0.808
"	125.4	0.596
"	250.8	0.497
"	501.5	0.120
MgSO ₄	26.5	0.901
"	79.5	0.669
"	159	0.441
"	318	0.188
KBr	83.9	0.906
"	167.7	0.819
"	251.5	0.748
"	503.1	0.579
KI	319.1	0.777
"	478.6	0.688
"	957.3	0.506
KSCN	326	0.691
"	489	0.580

Solubility of CO₂ in salts + Aq at 15.2°—Cont.

Salt	g. salt per l.	CO ₂
KSCN	978	0.387
KNO ₃	58.8	0.959
"	117.5	0.890
"	235.1	0.781
NaCl	12.9	0.978
"	64	0.760
"	128	0.580
"	192	0.466
NaBr	115.1	0.775
"	460.3	0.364
"	690.4	0.221
NaNO ₂	89.3	0.835
"	125	0.762
"	208.4	0.621
"	416.8	0.385
"	625.2	0.244
NaClO ₃	233.3	0.625
"	349.9	0.506
"	699.8	0.257
Na ₂ SO ₄	14.2	0.950
"	94.8	0.620
"	284.4	0.234
ZnSO ₄	38.3	0.903
"	76.7	0.783
"	230	0.474
"	460	0.209

(Setschenow, A. ch. 1892, (6) 25. 226.)

CO₂ is not disengaged at ordinary temp. from H₂O, in which 1/1000 pt. of CaCO₃ or MgCO₃ is held in solution thereby. These solutions have a great power of retaining CO₂, even at a boiling temp. or with diminished pressure, and they also absorb CO₂ from the air in much larger quantity than pure H₂O. (Bineau.)

BaCO₃ in H₂O also retains CO₂, even after long boiling. (Storer.)

CO₂ is also absorbed from the air by Na₂CO₃, or K₂CO₃ + Aq, especially if dilute.

Absorption of CO₂ by NaCl + Aq at t°.

α = Coefficient of absorption for a 6.52% NaCl solution.

α₁ = Coefficient of absorption for a 17.62% NaCl solution.

t°	α	α ₁
0	1.234	0.678
5	1.024	0.577
10	0.875	0.503
15	0.755	0.442
20	0.664	0.393
25	0.583	0.352
30	0.517	0.319
35	0.460	0.288
40	0.414	0.263
45	0.370	0.235
50	0.335	0.215
55	0.305	0.198
60	...	0.183

(Bohr, W. Ann. 1899, 68. 504.)

Absorption of CO₂ by CsCl + Aq.

M = Content in g. equiv. per l.

S = Solubility. (See under Oxygen.)

M	S _{15°}	S _{11°}
0.552	0.7771	1.001
0.554	0.7769	0.9995

(Geffcken, Z. phys. Ch. 1904, 49. 273.)

Absorption of CO₂ by KNO₃ + Aq.

M	S _{15°}	S _{11°}
0.536	0.7832	1.002
0.537	0.7818	0.9997
1.022	0.7452	0.9439
1.033	0.7447	0.9421

(Geffcken.)

Absorption of CO₂ by KI + Aq.

M	S _{15°}	S _{11°}
0.559	0.7678	0.9809
0.573	0.7676	0.9835
1.043	0.7236	0.9144
1.119	0.7166	0.9090

(Geffcken.)

Absorption of CO₂ by RbCl + Aq.

M	S _{15°}	S _{11°}
0.479	0.7705	0.9908
0.481	0.7698	0.9910
1.007	0.7190	0.9210
1.012	0.7157	0.9200

(Geffcken.)

Absorption of CO₂ by KBr + Aq.

M	S _{15°}	S _{11°}
0.550	0.7621	0.9783
0.565	0.7619	0.9766
1.056	0.7030	0.9100
1.064	0.7068	0.9065

(Geffcken.)

Absorption of CO₂ by KCl + Aq.

M	S _{15°}	S _{11°}
0.423	0.7695	0.9892
0.432	0.7667	0.9865
1.045	0.6920	0.8875
1.058	0.6961	0.8910

(Geffcken.)

Absorption of CO₂ by salts + Aq.

Salt	Grams CO ₂ absorbed by 75 cc. of salt solution at 15.5° and 720 mm
Kr	0.1280
CO ₂	0.1231
Cl	0.1213
Cl	0.1204
Cl	0.1087
Cl	0.1050
H ₂ SO ₄	0.1093
H ₂ SO ₄	
) ₂ + 24H ₂ O	0.0991
SO ₄	0.1002
SO ₄	
) ₂ + 24H ₂ O	0.1054
SO ₄	0.1140
SO ₄	0.1209
SO ₄	0.1047
SO ₄	0.0656
SO ₄	0.0527
SO ₄	0.0751
SO ₄	0.0720
ISO ₄	0.1017
ISO ₄	0.0999
(AsO ₄) ₃	0.0808
(PO ₄) ₃	0.0852
(AsO ₄) ₃	0.1111
(PO ₄) ₃	0.4989
B ₂ O ₃	0.2205
B ₂ O ₃	0.5317
B ₂ O ₃	0.8511
BO ₂	0.8124
(H ₂ B ₂ O ₄) ₂	0.7672
) ₂ + 12H ₂ O	0.5828
O ₂ + 10H ₂ O	0.8457
z	0.2081
	0.2618

ff, Z. phys. Ch. 1905, 53. 338-340.)

ility of CO₂ in KCl + Aq at 25°.tration, 7.45 g. in 100 cc. of solution.
gr. = 1.043.756 850 953 1116 1249 1362
0.694 0.693 0.688 0.700 0.709 0.710tration, 5 g. in 100 cc. of solution,
0.31.756 832 901 1050 1150 1223
0.731 0.727 0.724 0.726 0.735 0.736tration, 2.56 g. in 100 cc. of solution,
0.016.756 852 981 1079 1190 1362
0.767 0.761 0.761 0.762 0.768 0.766
and Creighton, Chem. Soc. 1910,
97. 557.)Solubility of CO₂ in NH₄Cl + Aq at 25°.Concentration (C) denotes number of
grams of solute in 100 cc. of solution.Density (D) equals the specific gravity of
the solution.Solubility (S) calculated by formula given
in the original article.

C	D	S
2 35	1 005	0.791
5 05	1 013	0.754
8 24	1 022	0.732
10 02	1 027	0.712
17 09	1 045	0.665

(Findlay and Shenn, Chem.
Soc. 1912, 101. 1461.)Solubility of CO₂ in KCl + Aq at 25°.

C	D	S
1 84	1 008	0.792
3 05	1 017	0.764
4 58	1 026	0.749
7.46	1 044	0.701

(Findlay and Shenn.)

Solubility of CO₂ in BaCl₂ + Aq at 25°.

C	D	S
2 80	1 018	0.789
5 81	1 040	0.741
8 15	1 054	0.710
9 97	1 070	0.676

(Findlay and Shenn.)

Solubility of CO₂ in (NH₄)₂Fe(SO₄)₂ + Aq at
25°.

C	D	S
9 51	1 052	0.641
10 26	1 057	0.629
22 47	1 124	0.460

(Findlay and Shenn.)

Solubility of CO₂ in solutions of sucrose at
25°.

C	D	S
2 63	1.009	0.813
5.16	1 018	0.798
9.68	1 038	0.767
12 33	1.051	0.744

(Findlay and Shenn.)

Solubility of CO₂ in solutions of chloral hy-
drate at 25°.

C	D	S
5.08	1.019	0.815
10 12	1 041	0.795

(Findlay and Shenn.)

100 vols. alcohol (0.803 sp. gr.) at 18° absorb 260 vols. CO₂.
100 vols. alcohol (0.840 sp. gr.) at 18° absorb 186 vols. CO₂. (de Saussure, l. c.)

Solubility of CO₂ in alcohol. 1 vol. alcohol at t° and 760 mm. dissolves V vols. CO₂ gas reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	4.3295	9	3.5844	18	3.0402
1	4.2368	10	3.5140	19	2.9921
2	4.1466	11	3.4461	20	2.9465
3	4.0589	12	3.3807	21	2.9034
4	3.9736	13	3.3178	22	2.8628
5	3.8908	14	3.2573	23	2.8247
6	3.8105	15	3.1993	24	2.7890
7	3.7327	16	3.1438
8	3.6573	17	3.0908

(Bunsen's Gasometry, pp. 287, 128, 153.)

Coefficient of absorption = 4.32955—
0.09395t + 0.00124t². (Bunsen.)

Much less sol. in 30% alcohol than in pure alcohol or pure H₂O. (Müller, W. Ann. 37. 24.)

Solubility of CO₂ in 99% alcohol at t°.

a = Coefficient of absorption, i. e., the no. of ccm. of CO₂ measured at 0° and 760 mm. which are absorbed at the given temp. and at an absorption pressure of 760 mm. by 1 ccm. alcohol.

a₁ = Coefficient of absorption corrected for increase in the volume of the alcohol used due to absorption of CO₂.

t°	a	a ₁
—65	38.41	35.93
—25	8.75	8.61
—20	7.51	7.41
—15	6.59	6.51
—10	5.75	5.69
—5	5.01	4.96
0	4.44	4.40
+5	3.96	3.93
10	3.57	3.55
15	3.25	3.23
20	2.98	2.96
25	2.76	2.74
30	2.57	2.56
35	2.41	2.39
40	2.20	2.19
45	2.01	2.00

(Bohr, W. Ann. 1900, (4) 1. 249.)

Solubility in 98.7% alcohol at t°.

a = Coefficient of absorption.

a₁ = Coefficient of absorption corrected for increase in volume of the alcohol used due to absorption of CO₂.

t°	a	a ₁
0	4.35	4.31
—10	5.43	5.38
—20	7.25	7.16
—30	9.97	9.79
—40	14.25	13.89
—50	21.28	20.49
—60	31.25	29.59
—65	39.89	37.22
—67	44.07	40.83

(Bohr, W. Ann. 1900, (4) 1. 253.)

Solubility in alcohol + Aq at t°.

t°	% by wt. of alcohol in the solvent	Solubility of CO ₂ in alcohol + Aq	Solubility of CO ₂ in H ₂ O
1.4	6.325	1.5864	1.6916
3.2	4.464	1.4878	1.5652
9.2	7.276	1.1829	1.2216
13.8	2.870	1.0268	1.0385

(Langer, C. C. 1904, I, 1583.)

Solubility of CO₂ in ethyl alcohol at 25°.

Concentration. 2.95 g. alcohol in 100 cc. of solution. Sp. gr. 25°/15° = 0.99308.

Pressure 737 836 929 1073 1213 1338
Solubility 0.812 0.813 0.812 0.811 0.813 0.811

Concentration. 3.01 g. alcohol in 100 cc. of solution. Sp. gr. 25°/15° = 0.99295.

Pressure 745 823 937 1083 1226 1357
Solubility 0.814 0.812 0.815 0.813 0.812 0.812

Concentration. 8.83 g. alcohol in 100 cc. of solution. Sp. gr. 25°/15° = 0.98342.

Pressure 747 846 942 1090 1231 1360
Solubility 0.786 0.786 0.784 0.785 0.786 0.788
(Findlay and Shenn, Chem. Soc. 1911, 99. 1315.)

Solubility of CO₂ in organic solvents at low temperatures.

Solvent. Ethyl alcohol.

t = —78°; sp. gr. = 0.872		
Pressure	Coefficient of absorption	Solubility
100	111.8	68.4
200	115.7	69.5
400	123.8	71.4
700	138.6	74.7

Solubility of CO₂ in organic solvents at low temperatures.—Continued

t = -59°; sp. gr. = 0.856		
Pressure	Coefficient of absorption	Solubility
100	40.85	27.27
200	41.00	27.16
400	42.35	27.65
700	44.15	28.10

Solvent. Methyl alcohol.

t = -78°; Sp. gr. = 0.884		
Pressure	Coefficient of absorption	Solubility
50	194.0	120.5
100	195.0	119.6
200	202.9	120.1
400	221.5	122.2
500	226.4	...
740	260.0	126.8

t° = -59°; sp. gr. = 0.866		
Pressure	Coefficient of absorption	Solubility
100	63.0	42.5
200	64.2	42.7
400	66.3	43.1
700	69.0	43.3

Solvent. Acetone.

t = -78°; sp. gr. = 0.900		
Pressure	Coefficient of absorption	Solubility
50	311	196.6
100	322	198.1
200	344.5	201.5
400	400	208.8
640	487	215.7
700	545.5	...

t = -59°; sp. gr. = 0.879		
Pressure	Coefficient of absorption	Solubility
100	97.8	67.2
200	101.2	68.0
460	106.6	72.8
700	118.8	72.8

Solvent.—Ethyl acetate.

t = -78°; sp. gr. = 1.017		
Pressure	Coefficient of absorption	Solubility
50	250.2	177.5
100	255.6	177.1
200	271.8	179.2
400	310.9	183.2
650	386.9	191.2

Solubility of Co₂ in organic solvents at low temperatures.—Continued

t = -59°; sp. gr. = 0.994		
Pressure	Coefficient of absorption	Solubility
100	85.3	65.6
200	86.3	65.3
400	91.6	66.7
700	101.5	69.7

Solvent. Methyl acetate.

t = -78°; sp. gr. = 1.056		
Pressure	Coefficient of absorption	Solubility
50	304.9	224.1
100	315.0	224.3
200	337.4	223.1
400	389.3	225.6
650	498.1	231.2

t = -59°; sp. gr. = 1.032		
Pressure	Coefficient of absorption	Solubility
100	94.3	75.8
200	98.45	77.1
400	103.6	77.6
700	112.9	79.0

(Stern, Z. phys. Ch. 1912, 81. 468.)

Solubility of CO₂ in ether at 0° = 7.33; at 10° = 6.044; at 15° = 5.46. (Christoff, Z. phys. Ch. 1912, 79. 459.)

Coefficient of absorption in chloroform is 0.20376 at 36.57 mm., and 4.43757 at 762 mm. pressure. (Woukoloff, C. R. 109. 62.)

100 vols. of following liquids absorb vols. CO ₂ at 18°—		
	Sp. gr.	Vols. CO ₂
Ether	0.727	217
Rectified naphtha	0.784	169
Oil of turpentine	0.860	166
Oil of lavender (freshly distilled)	0.880	191
Oil of thyme	0.890	188
Linseed oil	0.940	156
Olive oil	0.915	151
Gum-arabic + Aq (containing 25% of the gum)	1.092	75
Cane-sugar + Aq (containing 25% of sugar)	1.104	72

(de Saussure, l. c.)

1 vol. oil of turpentine absorbs 1.7-1.9 vols. CO₂ (Saussure.)

1 vol. spirit at 10° absorbs 2 vols. CO₂. (de Saussure.)

1 vol. olive oil at 10° absorbs 1 + vol. CO₂. (de Saussure.)

1 vol. oil of turpentine at 10° absorbs 2 vols. CO₂. (Bergman.)

1 vol. caoutchine absorbs 11 vols. CO₂. (Bergman.)

Coefficient of absorption for petroleum is 1.17 at 20° and 1.31 at 10°. (Gniewasz and Walfisz, Zeit. phys. Ch. 1. 70.)

100 vols. petroleum absorb 70 vols. CO₂ at 10°. (Robinet, C. R. 58. 608.)

Solubility of CO_2 in $\frac{N}{2}$ solutions of various organic substances at 20° .

Substance	Sp. gr. of $\frac{N}{2}$ solution	Coeff. of absorption	cc. CO_2 dissolved in 1000 g. H_2O
Dextrose	1.0328	0.792	841
Mannite	1.03031	0.782	833
Glycerine	1.01413	0.843	864
Pyrogallol	1.01718	0.853	894
Hydrochinon	1.00946	0.887	928
Resorcin	1.00958	0.901	945
Pyrocatechin	1.0107	0.868	908
Urethane	1.0037	0.869	907
Carbamide	1.00715	0.864	884
Thiocarbamide	1.00917	0.859	885
Antipyrine	1.01339	0.859	935
Acetamide	1.005	0.879	906
Acetic acid	1.0026	0.868	893
N. Propylic acid	0.9939	0.869	902

(Usher, Chem. Soc. 1910, 97. 73.)

Absorption of CO_2 by ethyl alcohol.

Amount of alcohol used = 0.093 ccm.

V = ccm. of CO_2 absorbed by the solvent at t° , reduced to a pressure of 1 kg. per sq. cm.

V_1 = ccm. of CO_2 absorbed by 1 ccm. of the solvent.

Pressure kg/sq.cm.	t°	Gas volume ccm.	V	V_1
30	20°	57.31	9.462	104.8
40			15.15	149.7
50			23.04	188.8
30	35°	60.05	7.114	77.87
40			10.52	113.1
50			14.73	144.5
60			19.63	173.0
70			27.39	210.8
40	60°	64.44	6.429	72.82
50			9.023	97.09
60			12.27	122.5
70			15.64	145.2
80			19.11	167.9
90			20.64	180.7
100			23.88	195.7
50	100°	72.19	3.809	42.49
60			6.034	66.05
70			8.374	88.67
80			10.76	111.2
90			13.06	129.0
100			14.90	145.7
110			16.22	155.0
120			18.93	174.6
130			20.48	182.6
140			20.61	186.0

(Sander, Z. phys. Ch. 1912, 78. 524.)

Absorption of CO_2 by propyl alcohol.
Amount of alcohol used = 0.103 ccm.
 V and V_1 . See under absorption of by ethyl alcohol.

Pressure kg/sq.cm.	t°	Gas volume ccm.	V	
20	20°	60.59	4.867	5
30			8.472	8
40			13.46	12
50			21.62	17
20	35°	62.96	3.493	4
30			6.307	6
40			9.296	9
50			13.99	12
60			18.90	15
70			35.03	22
80			49.23	26
20	60°	68.08	2.602	2
30			4.722	4
40			6.723	6
50			9.810	8
60			13.05	11
70			17.15	14
80			19.61	15
90			24.75	18
100			30.19	21
40	100°	76.27	2.592	2
50			5.669	5
60			8.025	7
70			10.44	9
80			13.13	10
90			15.72	13
100			17.10	14
110			20.95	16
120			23.55	17

(Sander.)

Absorption of CO_2 by ether.

Amount of ether used = 0.131 ccm.

V and V_1 . See under absorption of by alcohol.

Pressure kg/sq.cm.	t°	Gas volume ccm.	V	
45	35°	62.06	42.62	20
50			46.81	21
60			57.83	24
50	60°	67.11	28.49	17
60			35.24	19
70			42.01	21
80			46.64	22
90			50.72	23
100			56.63	24
60	100°	71.03	12.57	10
70			20.00	13
80			26.34	14
90			32.16	16
100			35.70	17

(Sander.)

Absorption of CO₂ by benzene.

Amount of benzene used = 0.080 ccm.
 d V₁. See under absorption of CO₂ by
 alcohol.

t°	Gas volume ccm.	V	V ₁
20°	55.14	2.728	46.89
		4.845	71.16
		9.618	125.3
		18.70	192.4
		30.10	264.3
35°	58.17	2.225	39.94
		3.373	48.65
		6.879	94.39
		11.56	138.3
		17.09	186.6
		25.73	243.1
		35.80	289.0
60°	61.86	2.140	34.57
		3.880	55.97
		6.699	88.71
		10.28	128.5
		13.57	156.6
		17.71	184.6
		22.50	215.0
		28.09	246.6
		33.76	284.4
100°	73.75	2.822	46.52
		3.981	58.46
		6.440	91.27
		8.398	119.0
		11.96	155.8
		14.57	182.5
		17.79	212.9
		20.60	237.7
		23.98	258.2

(Sander.)

Absorption of CO₂ by chlorbenzene.

Amount of chlorbenzene used = 0.106 ccm.
 d V₁. See under absorption of CO₂ by
 alcohol.

t°	Gas volume ccm.	V	V ₁
20°	61.03	5.813	62.61
		10.25	95.22
		17.17	137.3
		26.59	187.5
35°	64.16	4.650	46.66
		7.705	72.73
		11.81	101.5
		16.83	137.3
		22.82	168.3
		32.83	205.5
60°	69.38	3.685	35.86
		5.510	53.94
		7.982	73.69

Absorption of CO₂ by chlorbenzene.—Cont.

Pressure kg/sq.cm.	t°	Gas volume ccm.	V	V ₁
50			11.16	99.06
60			13.74	118.1
70			16.65	134.5
80			19.50	149.3
90			22.23	165.5
110			31.64	204.4
30	100°	77.73	3.562	33.65
40			5.008	48.16
50			7.106	63.78
60			8.701	77.24
70			10.37	91.02
80			12.05	103.00
90			13.88	121.2
100			14.89	121.5
110			16.35	130.7
120			17.77	140.7
130			18.54	146.8

(Sander.)

Absorption of CO₂ by brombenzene.

Amount of brombenzene used = 0.113 ccm.
 V and V₁. See under absorption of CO₂
 by ethyl alcohol.

Pressure kg/sq.cm.	t°	Gas volume ccm.	V	V ₁
30	20°	60.84	4.531	50.83
40			7.793	83.29
50			12.22	121.1
60			17.37	160.0
20	35°	63.96	3.947	43.38
30			5.782	62.69
40			8.508	90.43
50			11.96	116.4
60			16.00	146.0
70			22.56	184.1
80			41.26	233.9
20	60°	69.16	2.650	30.58
30			3.714	46.15
40			5.971	62.64
50			7.406	77.19
60			9.718	98.73
70			10.27	108.4
80			13.99	131.4
90			16.70	144.3
100			20.06	169.7
110			23.13	190.6
30	100°	77.48	2.970	30.56
40			4.032	41.49
50			5.833	59.64
60			7.239	72.64
70			8.330	82.56
80			9.714	92.86
90			11.14	107.1
100			12.79	118.0
110			13.80	125.3
120			15.50	140.7

(Sander.)

Absorption of CO_2 by nitrobenzene.

Amount of nitrobenzene used = 0.164 ccm.
V and V_1 . See under absorption of CO_2
by ethyl alcohol.

Pressure kg./sq. cm.	t°	Gas volume ccm.	V	V_1
15	20°	57.65	5.459	41.60
20			7.354	57.12
30			12.14	92.50
40			15.93	115.9
50			21.71	155.9
20	35°	59.86	5.644	44.48
30			8.658	68.23
40			11.98	94.39
50			15.59	113.4
60			19.94	145.1
70			25.57	179.6
80			34.95	227.0
20	60°	64.73	3.787	31.38
30			4.519	38.23
40			6.308	52.26
50			7.750	64.21
60			8.887	72.15
70			10.15	82.40
80			10.80	85.03
20	100°	75.52	2.749	24.67
30			4.162	41.00
40			5.393	50.36
50			6.832	63.80
60			7.763	70.85
70			9.048	75.75
80			10.65	86.86

(Sander.)

Absorption of CO_2 by toluene.

Amount of toluene used = 0.114 ccm.
V and V_1 . See under absorption of CO_2
by ethyl alcohol.

Pressure kg./sq. cm.	t°	Gas volume ccm.	V	V_1
20	20°	59.97	7.420	57.91
30			13.31	103.3
40			23.25	155.9
50			45.10	235.8
20	35°	63.05	6.018	49.60
30			10.13	82.63
40			16.03	118.8
50			23.34	155.8
60			31.39	192.1
70			44.17	225.8
30	60°	68.17	6.735	54.67
40			9.885	78.67
50			13.98	104.6
60			18.00	128.1
70			22.66	150.1
80			26.60	171.9
90			31.66	191.5
100			38.86	210.0

Absorption of CO_2 by toluene.—Continued

Pressure kg./sq. cm.	t°	Gas volume ccm.	V	V_1
30	100°	76.37	3.356	28.06
40			5.945	49.25
50			8.703	67.93
60			11.18	85.11
70			13.72	101.7
80			16.30	117.6
90			18.88	132.6
100			21.85	149.0
110			24.86	161.9
120			26.80	171.8
130			28.21	178.7

(Sander.)

Absorption of CO_2 by ethyl acetate.

Amount of ethyl acetate used = 0.155 ccm.
V and V_1 . See under absorption of CO_2
by ethyl alcohol.

Pressure kg./sq. cm.	t°	Gas volume ccm.	V	V_1
25	20°	60.30	29.43	158.6
30			37.91	188.2
40			51.26	227.9
30	35°	63.40	26.54	145.2
40			38.69	188.4
50			48.35	213.9
60			51.88	219.8
30	60°	68.55	18.12	118.0
40			25.67	140.5
50			33.21	165.2
60			40.12	186.7
70			45.47	201.1
80			49.16	223.4
40	100°	76.80	12.76	80.70
50			18.80	110.1
60			24.12	132.0
70			28.99	159.0
80			32.96	162.3
90			36.92	172.1
100			42.75	191.8

(Sander.)

Absorption of CO_2 by $\text{CH}_3\text{COOH} + \text{CCl}_4$.

Solvent	cc. CO_2 absorbed
1 mol. CH_3COOH	58.8
0.8 " $\text{CH}_3\text{COOH} +$	
0.2 " CCl_4	61.0
0.5 " $\text{CH}_3\text{COOH} +$	
0.5 " CCl_4	62.4
0.2 " $\text{CH}_3\text{COOH} +$	
0.8 " CCl_4	60.2
1 " CCl_4	57.6

(Christoff, J. phys. Ch. 1905, 53, 382.)

Absorption of CO₂ by C₂H₄Cl₂ + CS₂.

Solvent	cc. CO ₂ absorbed
1 mol. C ₂ H ₄ Cl ₂	209.7
0.8 " C ₂ H ₄ Cl ₂ +	
0.2 " CS ₂	173.4
0.5 " C ₂ H ₄ Cl ₂ +	
0.5 " CS ₂	140.0
0.2 " C ₂ H ₄ Cl ₂ +	
0.8 " CS ₂	71.9
1 " CS ₂	19.9

(Christoff.)

Solubility of CO₂ in organic solvents

$\frac{ds}{dt}$ = change of solubility for 1° increase in temp.

Solvent	Solubility at 25° C.	Solubility at 20° C.	Solubility at 15° C.	$\frac{ds}{dt}$
Glycerine	0.0302			...
Water	0.8256			
Carbon bisulphide	0.8699	0.8888	0.9448	-0.00747
Iodobenzene	1.301	1.371	1.440	-0.0139
Aniline	1.324	1.434	1.531	-0.0207
n-Toluidine	1.381	1.473	1.539	-0.0158
m-Toluidine	1.438	1.581	1.730	-0.0244
Eugenol	1.539	1.653	1.762	-0.0223
Benzonitrile	1.643			
Cumene	1.782	1.879	1.978	-0.0196
Carbene	1.802	1.921	2.034	-0.0232
Dichlorhydrin	1.810	1.917	2.020	-0.0210
Amyl alcohol	1.831	1.941	2.058	-0.0227
Bromobenzene	1.842	1.964	2.092	-0.0250
Isobutyl alcohol	1.849	1.964	2.088	-0.0239
Benzyl chloride	1.938	2.072	2.180	-0.0242
o-Xylene	2.090	2.216	2.348	-0.0256
Ethylene bromide	2.157	2.294	2.424	-0.0267
Chlorobenzene	2.265	2.420	2.581	-0.0316
Carbon tetrachloride	2.294	2.502	2.603	-0.0309
Propylene bromide	2.301	2.453	2.586	-0.0291
Toluene	2.305	2.426	2.557	-0.0256
Benzene	2.425	2.540	2.710	-0.0285
Amyl bromide	2.455	2.638	2.803	-0.0348
Nitrobenzene	2.456	2.655	2.845	-0.0389
Propyl alcohol	2.498			
Carrol	2.498	2.690	2.914	-0.0416
Ethyl alcohol (97%)	2.706	2.923	3.130	-0.0424
Benzaldehyde	2.841	3.057	3.304	-0.0463
Amyl chloride	2.910	3.127	3.363	-0.0453
Isobutyl chloride	3.105	3.388	3.659	-0.0554
Chloroform	3.430	3.681	3.956	-0.0526
Butyric acid	3.478	3.767	4.084	-0.0606
Ethylene chloride	3.525	3.796	4.061	-0.0536
Pyridine	3.656	3.862	4.291	-0.0635
Methyl alcohol	3.837	4.205	4.606	-0.0769
Amyl formate	4.026	4.329	4.646	-0.0620
Propionic acid	4.078	4.417	4.787	-0.0709
Amyl acetate	4.119	4.411	4.850	-0.0731
Glacial acetic acid	4.679	5.129	5.614	-0.0935
Isobutyl acetate	4.691	4.968		-0.0554
Acetic anhydride	5.206	5.720	6.218	-0.1012
Acetone	6.295	6.921		-0.1252
Methyl acetate	6.494			

(Just, Z. phys. Ch. 1901. 37. 354.)

Absorption of CO₂ by organic substances + Aq at 15°.

P = % of the organic substance in the solvent.

 β_{15}° = Coefficient of absorption at 15°. S_{15}° = Solubility at 15°.

Organic substance used	P	β_{15}°	S_{15}°
Chloral hydrate	0	0.996	
	0	0.992	1.056
	0	1.012	
	17.7	0.885	0.935
	21.8	0.860	0.908
	31.6	0.803	0.848
	37.0	0.790	0.834
	38.3	0.781	0.825
	49.8	0.760	0.802
	51.1	0.769	0.812
	52.6	0.764	0.807
	57.1	0.765	0.808
	61.1	0.780	0.824
	68.8	0.797	0.842
	71.0	0.812	0.857
	74.6	0.848	0.895
	79.4	0.903	0.953
Glycerine	0	1.003	1.064
	0	1.013	
	26.11	0.785	0.829
	27.69	0.800	0.845
	43.72	0.639	0.675
	46.59	0.620	0.655
	62.14	0.511	0.540
	73.36	0.449	0.474
	77.75	0.430	0.454
	87.74	0.422	0.446
	90.75	0.404	0.427
	96.64	0.415	0.438
	99.26	0.410	0.438

(Hammel, Z. phys. Ch. 1915, 90. 123.)

Solubility of carbon dioxide in solutions of aniline at 25°.

I. Concentration, 0.206 g. aniline in 100 c. c. of solution.

P = Pressure.

S = Solubility calc. according to formula given in original article.

P	S	P	S
748	0.865	1053	0.855
808	0.855	1159	0.862
920	0.857	1243	0.860

II. Concentration, 0.425 g. aniline in 100 c. c. of solution.

P	S	P	S
760	0.909	1150	0.897
816	0.897	1236	0.902
921	0.897	1380	0.908

Solubility of carbon dioxide in solutions of aniline at 25°.—*Continued*

III. Concentration, 0.566 g. aniline in 100 c. c. of solution.

P	S	P	S
760	0.935	1082	0.923
823	0.929	1223	0.924
941	0.925	1341	0.930

IV. Concentration, 0.743 g. aniline in 100 c. c. of solution.

P	S	P	S
760	0.953	1063	0.940
895	0.941	1223	0.940
983	0.940	1302	0.942

(Findlay and Creighton, Chem. Soc. 1910, 97. 555.)

Solubility of CO₂ in CS₂ increases approx. proportionally with the pressure. The absorption is greater at lower temp. and less at higher temp. than is required by Dalton's law. (Woukoloff, C. R. 1889, 108. 674.)

Absorption of CO₂ by sugar + Aq.

Sugar + Aq	Grams CO ₂ absorbed by 75 cc. of solution at 15.5° and 720 mm.
1/10-N sugar solution	0.1225
1/5-N " "	0.1089
1-N " "	0.0931

(Christoff, Z. phys. Ch. 1905, 53. 329.)

Absorption of CO₂ in sugar + Aq at 20°.

Conc. of solution	Sp. gr.	Coefficient of absorption
1 mol. per l.	1.01518	0.846
1/4 " " "	1.03125	0.815
1/2 " " "	1.06372	0.756
1 " " "	1.12809	0.649

(Usher, Chem. Soc. 1910, 97. 72.)

Liquid.—Not miscible with H₂O, though slightly sol. therein, or with fatty oils; miscible with alcohol, ether, CS₂, and the essential oils. (Thilorier, Mitchell.)

Unacted upon by H₂O; sol. in alcohol, ethers, petroleum, oil of turpentine, and CS₂. (Mareska and Donny.)

Petroleum dissolves 5 to 6 vols. liquid CO₂. (Cailletet, C. R. 75. 1271.)

Sl. sol. in CS₂. (Cailletet.)

Solid.—When immersed in H₂O, rapidly volatilizes and dissolves. With alcohol or ether it forms a semi-fluid mixture. (Channing, Am. J. Sci. (2) 5. 186.)

Only slightly sol. in anhydrous ether, but may be mixed therewith to a paste. (Thilorier.)

Sol. in methyl chloride below —65° to the point of sat. without decomp. (Villard, C. R. 1895, 120. 1413.)

+6H₂O. (Villard, C. R. 1894, 119. 369.)

Carbon selenide, C₄Se.

Sol. only in hot conc. H₂SO₄. (v. Bartsch, Ch. Z. 1906, 30. 810.)

C₄Se. Insol. in H₂O, CS₂, and ether. Easily sol. in hot conc. H₂SO₄; sol. in conc. NaOH + Aq from which it is pptd. by HCl (v. Bartsch.)

Carbon silicide CSi.

(Carborundum.) Not attacked by any acids, even HF; sl. attacked by caustic alkalies or carbonates. (Acheson, C. N. 68. 179.)

Not attacked by KOH + Aq. (Schützenberger, C. R. 114. 1089.)

Carbon monosulphide, CS.

Insol. in H₂O, alcohol, oil of turpentine, or benzene; somewhat sol. in CS₂ or ether; sol. in warm HNO₃; sol. in conc. KOH + Aq. (Sidot, C. R. 81. 32.)

Readily absorbed by alcohol and aniline. (Deninger, J. pr. 1895, (2) 51. 349.)

Carbon disulphide, CS₂.

Very sl. sol. in H₂O.

1 l. H₂O dissolves 2–3 g. CS₂ (Ckiani, Bull. Soc. 43. 562); 3.5–4.52 g. (Peligot, ib. 43. 563).

30 ccm. CS₂ shaken with 8690 ccm. H₂O at 20–23° for 18 days decreased 11 ccm. in 9 days and 1.4 ccm. in the next 3 days by diffused light, and 0.6 ccm. in the last 5 days (no light). Part of the CS₂ was decomp. and 7.85 ccm. were dissolved, therefore H₂O dissolves 1/100 of its weight CS₂. (Sestini, Gazz. ch. it. 1. 473.)

Solubility of CS₂ in H₂O.

100 pts. H₂O dissolve 0.203 pts. CS₂ at 12–13°

"	"	0.191	"	"	15–16°
"	"	0.168	"	"	25–27°
"	"	0.145	"	"	30–33°

(Page, C. N. 41. 195.)

Solubility of CS₂ in H₂O. a = g. CS₂ in 1000 ccm. solution at t°.

a	t°	a	t°	a	t°
2.04	0	1.79	20	1.11	40
1.99	5	1.69	25	0.70	45
1.94	10	1.55	30	0.14	49
1.87	15	1.37	35		

(Chancel and Parmentier, C. R. 100. 773.)

100 g. H₂O dissolve at t°:

t°	0	10	20	30
	0.258	0.239	0.201	0.195 g. CS ₂ .

(Rex, Z. phys. Ch. 1906, 55. 365.)

on of CS₂ vapor by H₂O at t°.

	Coefficient of absorption
	3.573
	2.189
	1.346
	0.799

data of Chancel and Parmentier, 33.)

r, Z. phys. Ch. 1906, 55. 352.)

CS₂ are most easily absorbed by solution of KOH. Sl. absorbed by and very slowly by CuSO₄, +Aq, conc. H₂SO₄, or CaCl₂ in (Berthelot, A. ch. (3) 51. 74.)

alcohol. S=strength of alcohol at by weight; P=pts. CS₂ which 10 ccm. alcohol at 17°.

P	S	P
∞	91.37	5.00
18.20	84.12	3.00
13.20	76.02	2.00
10.00	48.40	0.20
7.00	47.90	0.00

midt and Follenius, B. 4. 583.)

with absolute alcohol, ether, ethery oils, and liquid CO₂.

disulphide, C₂S₂.

H₂O; easily sol. in alcohol, ether, benzene, and CS₂. The alcoholic solutions decomp. on standing. (26. 2960.)

alcohol with decomp. Sol. in CS₂, benzene. (Stock, B. 1912, 45. 3575.)
 ification. Insol. in H₂O and solvents. Sol. in KOH+Aq.

hoselenide, CSSe.

5°, bpt. +84°.

by light. Not attacked by H₂O. t conc. HNO₃. Decomp. by Br₂ ol. in alcohol with decomp. Mis-S₂. (Stock, B. 1914, 47. 150.)

hotelluride, CSTe.

t°. Very unstable.

with CS₂ and benzene without stock, B. 1914, 47. 142.)

chloroplatindiamine carbon-chloroplatindiamine nitrate.

[₂H₄]₂(CO₃)₂, Cl₂Pt(N₂H₄NO₃)₂. e. (Cleve, J. B. 1867. 321.)

Carbonatonitratoplatindiamine carbonate, CO₃(NO₃)₂[Pt(N₂H₄)₂(CO₃)₂].

Sol. in boiling H₂O. (Cleve.)

Carbonatotetramine cobaltic bromide, Co(NH₃)₄CO₃Br.

Much less sol. than chloride. (Jørgensen, Z. anorg. 2. 279.)

— carbonate, [Co(NH₃)₄CO₃]₂CO₃+3H₂O. Very sol. in H₂O. (Jørgensen.)

— chloraurate, [Co(NH₃)₄CO₃]₂AuCl₄+ $\frac{1}{2}$ H₂O.

Somewhat sol. in H₂O; nearly absolutely insol. in alcohol. (Jørgensen.)

— chloride, Co(NH₃)₄CO₃Cl.

Easily sol. in H₂O; insol. in alcohol. (Jørgensen.)

— chloroplatinate, [Co(NH₃)₄CO₃]₂PtCl₄+2H₂O.

Nearly insol. in H₂O and alcohol. (Jørgensen.)

— chloroplatinite, [Co(NH₃)₄CO₃]₂PtCl₄.

Nearly insol. in H₂O; wholly in alcohol. (Jørgensen.)

— dithionate, [Co(NH₃)₄CO₃]₂S₂O₆.

Ppt. (Jørgensen.)

— iodide, Co(NH₃)₄CO₃I.

Much less sol. than bromide or chloride. (Jørgensen.)

— nitrate, Co(NH₃)₄CO₃NO₃+ $\frac{1}{2}$ H₂O.

Sol. in about 15 pts. cold H₂O; insol. in alcohol. (Jørgensen.)

— sulphate, [Co(NH₃)₄CO₃]₂SO₄+3H₂O.

Considerably less sol. in H₂O than the nitrate. (Jørgensen.)

Carbonic acid, H₂CO₃.

See Carbon dioxide.

Carbonates.

Carbonates of Na, K, Rb, and Cs are easily sol. in H₂O; carbonates of Li and Tl are much less sol.; other carbonates are nearly or quite insol. All carbonates are sol. to some extent in H₂O containing CO₂. All carbonates, except those of NH₄, Rb, and Cs, are insol. in alcohol.

Sol. in those acids which are themselves sol. in H₂O, except HCN and H₃BO₃.

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 824.)

Aluminum carbonate, basic.

$5\text{Al}_2\text{O}_3, 6\text{CO}_2 + 37\text{H}_2\text{O} = 3\text{Al}(\text{OH})_3, \text{Al}_2(\text{CO}_3)_3 + 14\text{H}_2\text{O}$. (Seubert, Z. anorg. 1893, 4. 67.)

$\text{Al}_2\text{O}_3, \text{CO}_2$. (Parkmann, Sill. Am. J. (2) 34. 324.)

$3\text{Al}_2\text{O}_3, 2\text{CO}_2 + 16\text{H}_2\text{O}$. (Muspratt and Danson, A. 72. 120.)

$3\text{Al}_2\text{O}_3, 2\text{CO}_2 + 9\text{H}_2\text{O}$. (Wallace, Chem. Gas. 1858. 410.)

$5\text{Al}_2\text{O}_3, 3\text{CO}_2 + 18\text{H}_2\text{O}$. (Bley, J. pr. 39. 11.)

$2\text{Al}_2\text{O}_3, \text{CO}_2 + 6\text{H}_2\text{O} = 10\text{Al}(\text{OH})_3, \text{Al}_2(\text{CO}_3)_3 + 3\text{H}_2\text{O}$. Sol. in cold dil. acids. (Schlumberger, Bull. Soc. 1895, (3) 13. 46.)

+ $8\text{H}_2\text{O}$. (Urbain and Renoul, J. Pharm. (4) 30. 340.) = $10\text{Al}(\text{OH})_3, \text{Al}_2(\text{CO}_3)_3 + 9\text{H}_2\text{O}$. (Seubert, Z. anorg. 1893, 4. 67.)

$8\text{Al}_2\text{O}_3, 3\text{CO}_2 + 40\text{H}_2\text{O}$. (Langlois, A. ch. (3) 43. 505.)

All are precipitates, insol. in H_2O , sol. in acids, and give off CO_2 at slight heat.

There are no definite carbonates of aluminum. (Cameron, J. phys. Chem. 1908, 12. 572.)

Aluminum ammonium carbonate, $\text{Al}_2\text{O}_3, \text{CO}_2, (\text{NH}_4)_2\text{CO}_3 + 4\text{H}_2\text{O}$.

Precipitate. (Rose, Pogg. 91. 460.)

Aluminum sodium carbonate, $\text{Al}_2\text{O}_3, \text{CO}_2, 2\text{Na}_2\text{CO}_3 + 24\text{H}_2\text{O}$.

Precipitate. Sol. in cold dil. acids. (Bley, J. pr. 39. 22.)

Ammonium carbonate, $(\text{NH}_4)_2\text{CO}_3 + \text{H}_2\text{O}$.

Sol. at 15° in its own weight H_2O . Solution in H_2O gives off gas at $70-75^\circ$ and boils at $75-80^\circ$. Sl. sol. in cold dil. $\text{NH}_4\text{OH} + \text{Aq}$, more sol. at ordinary temp. Insol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Divers, Chem. Soc. (2) 8. 171, 259, and 364.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.)

Insol. in alcohol

Insol. in CS_2 . (Arcetowski, Z. anorg. 1894, 6. 257.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

100 g. pure glycerine dissolve 20 g. $(\text{NH}_4)_2\text{CO}_3$ at 15° . (Ossendowski, Pharm. J. 1907, 79. 575.)

Ammonium hydrogen carbonate, NH_4HCO_3 .

Sol. at 15° in about 8 pts. H_2O . (Berthollet, J. Phys. 66. 108.)

Sol. at 12.8° in about 6 pts. H_2O . (J. Davy, N. Edinb. J. 16. 245.)

Solution decomp. on air or by gentle heat or by addition of the solid salt. (Berthollet.)

100 pts. H_2O dissolve at 0° , 11.9 pts.; at 10° , 15.85 pts.; at 20° , 21 pts.; at 30° , 27 pts. NH_4HCO_3 . (Dibbitts, J. pr. (2) 10. 417.)

Solubility of NH_4HCO_3 in $\text{NH}_4\text{Cl} + \text{Aq}$, sat. with CO_2 , at t° .

t°	g. per 100 g. H_2O		Sp. gr. of sat. solution
	NH_4Cl	NH_4HCO_3	
0°	0 29.06	11.9 3.6	1.077
15°	0 2.99 6.06 8.51 11.68 18.30 26.93 33.25 34.35	18.64 16.29 14.22 12.69 11.68 9.33 7.73 6.64 6.42	1.064 1.063 1.062 1.062 1.065 1.069 1.076 1.085 1.085
30°	0 39.7	27.0 9.1	

(Fedotieff, Z. phys. Ch. 1904, 49. 168.)

Solubility of NH_4HCO_3 in $\text{NaHCO}_3 + \text{Aq}$, sat. with CO_2 , at t° .

t°	g. per 100 g. H_2O		Sp. gr. of sat. solution
	NaHCO_3	NH_4HCO_3	
0°	0 4.82	11.90 10.94	1.072
15°	0 5.92	18.64 17.06	1.064 1.080
30°	0 7.0	27.0 23.0	

(Fedotieff, Z. phys. Ch. 1904, 49. 168.)

Solubility of NH_4HCO_3 in $\text{NH}_4\text{NO}_3 + \text{Aq}$ at t° .

t°	g. per 100 g. H_2O		Sp. gr. of sat. solution
	NH_4NO_3	NH_4HCO_3	
0°	0 118	11.90 4.52	1.2023
15°	0 23.26 49.82 103.4 128.9 166.9	18.64 12.91 10.33 8.25 7.79 7.49	1.064 1.113 1.164 1.242 1.269 1.303
30°	0 231.9	26.96 12.57	..

(Fedotieff and Koltunoff, Z. anorg. 1914, 34. 251.)

Insol. in alcohol. (J. Davy.)

Insol. in acetone. (Eidmann, C. C. 1000, II. 1014; Naumann, B. 1904, 37. 4329.)

ium dihydrogen carbonate,
 $\text{H}_4\text{H}_2(\text{CO}_3)_2 + \text{H}_2\text{O}$.

5 pts. H_2O at 15° ; decomp. by more
 by heat. (Divers, Chem. Soc. (2) 8.
), and 364.)
 l. in alcohol.

ium hydrogen carbonate carbamate,
 $\text{H}_4\text{HCO}_3, \text{NH}_4\text{CONH}_2$. (Salts of harts-
 n.)

salt dissolves at:

13° in 4 pts. H_2O .

16.7° " 3.3 "

32.2° " 2.7 "

40.6° " 2.4 "

49° " 2 "

J. Davy, N. Edinb. J. 16. 245.)

g alcohol dissolves out carbamate, and
 onate remains undissolved.

$\text{HCO}_3, \text{NH}_4\text{CO}_2\text{NH}_2$. (Commercial
 te of ammonia.)

at 15° in 4 pts. H_2O , at 65° in $1\frac{1}{2}$ pts.
 Divers.)

s. salt + 100 pts. H_2O lower temp. from
 3.2° . (Rüdorff, B. 2. 68.)

1.667 pts. cold, and 0.833 pt. hot H_2O . (Four-

100 pts. H_2O at 13° dissolve 25 pts.

" " 17° " 30 "

" " 37° " 37 "

" " 41° " 40 "

" " 49° " 50 "

(Berzelius.)

s. H_2O at 15.5° dissolve 33 pts.; at 100° , 100
 re's Dict.)

2 pts. H_2O at 15.5° , and in less than 1 pt.
 H_2O ; sat. solution at 15.5° contains 33.3%, and
 ng solution 50%. (Abl.)

queous solution at 10° contains 15.7%. (Eller.)

queous solution at (?) contains 6.1%. (Mus-

tion in the cold contains 37.5%. (Fourcroy.)

not dissolve as such in H_2O ; $(\text{NH}_4)_2\text{CO}_3$ dis-
 it first, and NH_4HCO_3 later. (Scanlan.)

of carbonate of ammonia + Aq at 12° .

w.	Sp. gr. at 12° .	% Carb. ammon.	Change of sp. gr. for 1°C .
	1.005	1.66	0.0002
	1.010	3.18	0.0002
	1.015	4.66	0.0003
	1.020	6.04	0.0003
	1.025	7.49	0.0003
	1.030	8.93	0.0004
	1.035	10.35	0.0004
	1.040	11.86	0.0004
	1.045	13.36	0.0005
	1.050	14.83	0.0005
	1.055	16.16	0.0005
	1.060	17.70	0.0005
	1.065	19.18	0.0005
	1.070	20.70	0.0005
	1.075	22.25	0.0006
	1.080	23.78	0.0006
	1.085	25.31	0.0006

Sp. gr. of carbonate of ammonia + Aq at
 12° .—Continued

Deg. Tw.	Sp. gr. at 12° .	% Carb. ammon.	Change of sp. gr. for 1°C .
18	1.090	26.82	0.0007
19	1.095	28.33	0.0007
20	1.100	29.93	0.0007
21	1.105	31.77	0.0007
22	1.110	33.45	0.0007
23	1.115	35.08	0.0007
24	1.120	36.88	0.0007
25	1.125	38.71	0.0007
26	1.130	40.34	0.0007
27	1.135	42.20	0.0007
28	1.140	44.29	0.0007
29	1.144	44.90	0.0007

(Lunge, Chem. Ind. 1883. 2.)

Sp. gr. of aqueous solution of salt with com-
 position 31.3% NH_3 , 56.6% CO_2 , 12.1%
 H_2O . 100 pts. of solution contain—

6.58 9.96 14.75 19.83 25.71 pts. salt
 1.0219 1.0337 1.0497 1.0672 1.0863 sp. gr.

29.74 35.85 40.23 44.90 pts. salt.

1.0995 1.1174 1.1297 1.1414 sp. gr.

(J. H. Smith, Chem. Ind. 1883. 3.)

Conc. alcohol dissolves out carbamate and
 leaves carbonate. (Hünefeld, J. pr. 7. 25.)

Insol. in acetone. (Naumann, B. 1904, 37.
 4328.)

Ammonium cerous carbonate, $(\text{NH}_4)_2\text{CO}_3$,
 $\text{Ce}_2(\text{CO}_3)_3 + 6\text{H}_2\text{O}$.

Ppt. Very sl. sol. in conc. $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.
 (Meyer, Z. anorg. 1904, 41. 104.)

Ammonium chromous carbonate, $(\text{NH}_4)_2\text{CO}_3$,
 $\text{CrCO}_3 + \text{H}_2\text{O}$.

Decomp. by moist air; sol. in dil. HCl and
 H_2SO_4 . (Baugé, C. R. 1896, 122. 476.)

Ammonium cobaltous carbonate, $(\text{NH}_4)_2\text{CO}_3$,
 $\text{CoCO}_3 + 4\text{H}_2\text{O}$.

Permanent. Sol. in H_2O . (Deville, A. ch.
 (3) 35. 460.)

$(\text{NH}_4)_2\text{O}$, 2CoO , $4\text{CO}_2 + 9\text{H}_2\text{O}$. Quickly de-
 comp. on air; sol. in H_2O . (Deville.)
 $+12\text{H}_2\text{O}$. Sol. in H_2O .

Ammonium didymium carbonate, $(\text{NH}_4)_2\text{CO}_3$,
 $\text{Di}_2(\text{CO}_3)_3 + 3\text{H}_2\text{O}$.

Insol. in H_2O . (Cleve.)

Ammonium dysprosium carbonate,
 $\text{NH}_4\text{Dy}(\text{CO}_3)_3 + \text{H}_2\text{O}$.

Only sl. sol. in H_2O . (Jantsch, B. 1911, 44.
 1277.)

Ammonium glucinum carbonate, $2(\text{NH}_4)_2\text{CO}_3 \cdot 3\text{GlCO}_3(?)$.

Very sol. in cold, decomp. by hot H_2O . Nearly insol. in alcohol. (Debray.)

Composition is $(\text{NH}_4)_2\text{CO}_3 \cdot 2\text{GlCO}_3 \cdot \text{Gl}(\text{OH})_3 + 2\text{H}_2\text{O}$. (Humpidge, Royal Soc. Proc. 39. 1.)

Ammonium lanthanum carbonate, $\text{La}_2(\text{CO}_3)_3 \cdot (\text{NH}_4)_2\text{CO}_3 + 4\text{H}_2\text{O}$.

Ppt. (Meyer, Z. anorg. 1904, 41. 102.)

Ammonium magnesium carbonate, $(\text{NH}_4)_2\text{Mg}(\text{CO}_3)_2 + 4\text{H}_2\text{O}$.

Sol. in 71 pts. H_2O with decomp.; more sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Divers, Chem. Soc. 51. 196.)

H_2O containing $(\text{NH}_4)_2\text{CO}_3$ dissolves very slightly; more sol. in H_2O containing NH_4Cl . (Favre, A. ch. (3) 10. 473.)

Ammonium magnesium hydrogen carbonate, $(\text{NH}_4)_2\text{Mg}_2\text{H}_2(\text{CO}_3)_4 + 8\text{H}_2\text{O}$, or $12\text{H}_2\text{O}$.

Decomp. on air. (Deville, A. ch. (3) 35. 454.)

Ammonium neodymium carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot \text{Nd}_2(\text{CO}_3)_3 + 4\text{H}_2\text{O}$.

Ppt. Sl. sol. in conc. $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Meyer, Z. anorg. 1904, 41. 106.)

Ammonium nickel carbonate, $\text{NH}_4\text{HCO}_3 \cdot \text{NiCO}_3 + 4\text{H}_2\text{O}$.

Insol. in H_2O . (Deville, A. ch. (3) 35. 452.)

Ammonium praseodymium carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot \text{Pr}_2(\text{CO}_3)_3 + 4\text{H}_2\text{O}$.

Ppt. Insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Meyer, Z. anorg. 1904, 41. 105.)

Ammonium samarium carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot \text{Sm}_2(\text{CO}_3)_3 + 4\text{H}_2\text{O}$.

Ppt.

Ammonium scandium carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot 2\text{Sc}_2(\text{CO}_3)_3 + 6\text{H}_2\text{O}$.

Difficultly sol. in H_2O . Sol. in cold alkali-carbonate + Aq , less sol. in hot. (R. Meyer, Z. anorg. 1910, 67. 410.)

Ammonium tin (stannous) carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot 2\text{SnCO}_3 + 3\text{H}_2\text{O}$.

Decomp. by cold H_2O . (Deville, A. ch. (3) 35. 456.)

Ammonium uranyl carbonate, $2(\text{NH}_4)_2\text{CO}_3 \cdot \text{UO}_2\text{CO}_3$.

Sol. at 15° in 20 pts. H_2O , more abundantly in H_2O containing $(\text{NH}_4)_2\text{CO}_3$. (Ebelmen.)

Insol. in pure H_2O ; sol. in H_2O containing $(\text{NH}_4)_2\text{CO}_3$. Solution is decomp. by boiling. (Berzelius.)

Sol. in $\text{SO}_2 + \text{Aq}$. (Berthier, A. ch. (3) 7. 76.)

$3(\text{NH}_4)_2\text{CO}_3 \cdot 2(\text{UO}_2)_2\text{CO}_3 + 4\text{H}_2\text{O}$. Sol. in H_2O . (Giolitti C. C. 1905, II. 227.)

Ammonium vanadyl carbonate, $3(\text{NH}_4)_2\text{O} \cdot 7\text{VO}_2 \cdot 5\text{CO}_2 + 16\text{H}_2\text{O}$.

Sl. sol. in H_2O .

Sol. in acids and alkalies. (Koppel, Z. anorg. 1905, 45. 350.)

Ammonium yttrium carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot \text{Y}_2(\text{CO}_3)_3 + 2\text{H}_2\text{O}$.

Insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Mosander.)

Ammonium zinc carbonate, basic, $3\text{ZnO} \cdot \text{NH}_4\text{OH} \cdot 2\text{CO}_2 + \text{H}_2\text{O}$.

Insol. in H_2O . (Kassner, Arch. Pharm. (3) 27. 673.)

Ammonium zinc carbonate, $(\text{NH}_4)_2\text{CO}_3 \cdot \text{ZnCO}_3$.

Insol. in H_2O . (Deville.)

Quite sol. in H_2O ; more sol. than $(\text{NH}_4)_2\text{CO}_3$, MgCO_3 . Tolerably permanent in the air. Slowly decomp. by cold, rapidly by hot H_2O .

Very sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Not attacked by alcohol. (Favre, A. ch. (3) 10. 481.)

Barium carbonate, BaCO_3 .

Sol. in 4304 pts. cold, and 2304 pts. boiling H_2O . (Fourcroy.)

Sol. in 47,620 pts. H_2O . (Bineau, A. ch. (3) 51. 290.)

Sol. in 14,137 pts. H_2O at $16-20^\circ$, and 15,421 pts. at 100° . (Fresenius.)

Sol. in 12,027 pts. H_2O at 15° . (Kremer, Pogg. 85. 247.)

Calculated from electrical conductivity of solution, 1 pt. BaCO_3 is sol. in 64,070 pts. H_2O at 8.8° and 45,566 pts. at 24.2° . (Holleman, Z. phys. Ch. 12. 125.)

Solubility in H_2O at t° .

t°	g. sol. in 100 g. H_2O
14	4.32×10^{-4}
18	4.57×10^{-4}
23	4.89×10^{-4}
27	5.22×10^{-4}
32	5.69×10^{-4}
38	6.27×10^{-4}

(Weissenberger, Z. phys. Ch. 1914, 88. 266.)

"Solubility product" = 8.1×10^{-9} mol. l. (McCoy and Smith, J. Am. Chem. Soc. 1911, 33. 473.)

Sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$. (See *barium hydrogen carbonate*.)

Easily sol. in dil. acids. Not acted upon by conc. $\text{HNO}_3 + \text{Aq}$.

Not decomp. by 1 pt. $\text{H}_2\text{SO}_4 + 6$ pts. absolute alcohol. Slowly decomp. by 1 pt.

+6 pts. absolute alcohol. Slowly de-
by 1 pt. H₂C₂O₄+6 pts. absolute al-
t decomp. by absolute alcoholic solu-
of racemic, tartaric, citric, or glacial
acids. (Babington and Phillips, 1816.)
most completely insol. in H₂O containing
OH and (NH₄)₂CO₃, when digested in
a solution and allowed to stand. 1 pt.
)₂ dissolves in 141,000 pts. of such a solu-
(Fresenius.)
t more sol. in NaCl+Aq than in H₂O.
sten.)
in cold NH₄Cl, NH₄NO₃, or NH₄ suc-
e+Aq. (Vogel, J. pr. 7. 453.)
nols. NH₄Cl dissolved in H₂O dissolve 1
BaCO₃ by continued boiling. (Smith,
Mag. J. 9. 540.)
ubility in H₂O increases by addition of
Cl, at first strongly, then less strongly
inally strongly again. (D'Agustino and
grino, Gazz. ch. it. 1908, 38 (1) 532.)
newhat sol. in K₂CO₃+Aq. (Wacken-
, A. 24. 30.)

Solubility of BaCO₃ in KCl+Aq at bpt. of solution

KCl per 100 g. solution	g. BaCO ₃ per 1000 cc. sat. solution
0.15	0.0847
1	0.1781
3	0.2667
10	0.4274
30	0.5550

Cantoni and Goguelia, Bull. Soc. 1905, (3) 33. 13.)

Solubility of BaCO₃ in NaCl+Aq at bpt. of solution

NaCl per 100 g. solution	g. BaCO ₃ per 1000 cc. sat. solution
0.15	0.0587
1	0.0787
3	0.1056
10	0.1575
30	0.2784

(Cantoni and Goguelia, l. c.)

Solubility of BaCO₃ in 10% KCl+Aq at t°.

t°	g. BaCO ₃ per 1000 cc. sat. solution
10	0.2175
20	0.2408
40	0.2972
60	0.3491
80	0.4049

(Cantoni and Goguelia, l. c.)

Solubility of BaCO₃ in 10% NaCl+Aq at t°.

t°	g. BaCO ₃ per 1000 cc. sat. solution
10	0.1085
20	0.1126
40	0.1231
60	0.1303
80	0.1418

(Cantoni and Goguelia, l. c.)

Slowly sol. in conc. Na₂SO₄, MgSO₄, ZnSO₄, Ca(NO₃)₂, or CaCl₂+Aq, but insol. in ZnCl₂+Aq. (Karsten.)
Sl. decomp. by boiling K₂SO₄+Aq.
Sl. decomp. in the cold by 1 pt. K₂SO₄+2 pts. Na₂SO₄+Aq.
Decomp. by salts of Al, Mn, Cr, Fe, U, Bi, Cd, Cu, Hg, Pb, Sn^{II}, Sn^{IV}, Hg₂, Rh, Ir, Au, with pptn. of oxide of metal. (Rose, Tr.)
Pptn. of BaCO₃ is hindered by presence of alkali citrates or metaphosphates.
Sol. in solutions of various salts, as in the case of calcium carbonate (see *Calcium carbonate*). The solvent power of these solutions for barium carbonate is somewhat less than for calcium carbonate.
Insol. in acetone. (Naumann, B. 1904, 37. 4329.)
Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3602.)
Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)
Min. *Witherite*.

Barium hydrogen carbonate, BaH₂(CO₃)₂(?).
100 pts. H₂O containing CO₂ dissolve 0.079 pt. BaCO₃. (Bineau.)
100 pts. H₂O containing CO₂ dissolve 0.17 pt. BaCO₃. (Lassaigne.)
100 pts. H₂O sat. with CO₂ under a pressure of 4-6 atmospheres dissolve 0.725 pt. BaCO₃. Upon evaporating, BaCO₃ is deposited. (Wagner, Z. anal. 6. 167.)
BaCO₃ is sol. in 833 pts. H₂O sat. with CO₂ at 10°. (Lassaigne.)
BaCO₃ is sol. in 830 pts. H₂O sat. with CO₂ at 10°. (Fourcroy.)
BaCO₃ is sol. in 1550 pts. H₂O sat. with CO₂ at 10°. (Bergman.)
100 cc. H₂O sat. with CO₂ dissolve 0.73 g. BaH₂(CO₃)₂. (McCoy and Smith, J. Am. Chem. Soc. 1911, 33. 473.)

Barium calcium carbonate, BaCO₃, CaCO₃.
Min. *Barytocalcite*, *Bromlite*. Sol. in dil. acids.

Barium uranyl carbonate, BaO, 2UO₃, 2CO₂+5H₂O. Decomp. by H₂O. (Blinkoff. Dissert. 1900.)
+8H₂O. Decomp. by H₂O. (Blinkoff.)

Bismuth carbonate, basic, $(\text{BiO})_2\text{CO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in acids. Insol. in $\text{CO}_2 + \text{Aq.}$ (Bergman.)

Completely sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$; sl. sol. in $\text{K}_2\text{CO}_3 + \text{Aq.}$; insol. in $\text{Na}_2\text{CO}_3 + \text{Aq.}$ (Lau-gier.)

Absolutely insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ unless H_3PO_4 or H_3AsO_4 are present. (Berzelius.)

Insol. in $(\text{NH}_4)_2\text{CO}_3$, K_2CO_3 , or $\text{Na}_2\text{CO}_3 + \text{Aq.}$ (Rose.)

Sol. in $\text{NH}_4\text{Cl} + \text{Aq.}$ (Wackenroder.) Insol. in $\text{NH}_4\text{NO}_3 + \text{Aq.}$ (Brett.)

Sol. in $\text{CaCl}_2 + \text{Aq.}$ (Pearson.)

Min. *Bismuthosphærite*.

$3\text{Bi}_2\text{O}_3, \text{CO}_2$. Min. *Bismuthite*. Easily sol. in acids.

$4\text{Bi}_2\text{O}_3, 3\text{CO}_2 + 4\frac{1}{2}\text{H}_2\text{O}$. Min. *Bismuth spar*. Easily sol. in acids.

Bismuth potassium carbonate, $\text{Bi}_2\text{OK}_4(\text{CO}_3)_4 + \text{H}_2\text{O}$.

Decomp. by large quantities of H_2O . (Reynolds, Chem. Soc. 1898, 73. 266.)

Cadmium carbonate, CdCO_3 .

Insol. in H_2O ; easily sol. in acids; insol. in K_2CO_3 , and $\text{Na}_2\text{CO}_3 + \text{Aq.}$; very sl. sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ (Fresenius.)

Easily sol. in NH_4 sulphate, nitrate, and succinate + Aq. (Wittstein.)

Sol. in $\text{KCN} + \text{Aq.}$; sol. in cold $\text{NH}_4\text{Cl} + \text{Aq.}$; less sol. in $\text{NH}_4\text{NO}_3 + \text{Aq.}$ (Brett, 1837.)

Not prevented from pptn. by non-volatile organic substances. (Rose.)

Not pptd. from solutions containing sodium citrate. (Spiller.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+ $\frac{1}{2}\text{H}_2\text{O}$. (Lefort, J. B. 1847. 346.) (Kraut, Z. anorg. 1897, 13. 14.)

Cadmium carbonate hydrazine, $\text{CdCO}_3, 2\text{N}_2\text{H}_4$.

Easily sol. in cold $\text{NH}_4\text{OH} + \text{Aq.}$ (Franzen, Z. anorg. 1908, 60. 281.)

Cæsium carbonate, Cs_2CO_3 .

Very deliquescent, and sol. in H_2O .

100 pts. absolute alcohol dissolve 11.1 pts. Cs_2CO_3 at 19° ; 20.1 pts. Cs_2CO_3 at boiling temp. (Bunsen.)

Cæsium hydrogen carbonate, CsHCO_3 .

Not deliquescent. Sol. in H_2O .

Calcium carbonate basic, $\text{CaO}, \text{CaCO}_3 + \text{H}_2\text{O}$.

Hardened by H_2O , but not dissolved. (Raoult, C. R. 92. 189.)

Calcium carbonate, CaCO_3 .

More sol. in cold than in hot H_2O . (Gmelin.)

When recently pptd., sol. in 8834 pts. boiling, and 10,601 pts. cold H_2O ; much less sol. in H_2O containing NH_4OH and $(\text{NH}_4)_2\text{CO}_3$, 65,246 pts. of which dissolve 1 pt. CaCO_3 . (Fresenius (1846), A. 59. 122.)

Sol. in 16,000 pts. pure H_2O . (Brandes, 1835.)

Sol. in 12,858 pts. pure H_2O at 15° . (Kremers, Pogg. 85. 247.)

Sol. in 16,000–24,000 pts. pure H_2O . (Bucholz.)

1 l. H_2O dissolves 34 mg. CaCO_3 . (Chevallet, Z. anal. 8. 91; Hoffmann, Z. anal. 4. 414.)

1 l. H_2O may contain 0.016 g. CaCO_3 , i. e., 1 pt. is sol. in 62,500 pts. H_2O . (Bineau, A. ch. (3) 51. 290.)

1 l. H_2O dissolves 0.02 g. CaCO_3 , i. e., 1 pt. CaCO_3 is sol. in 50,000 pts. H_2O . (Peligot.) Solubility is much affected by CO_2 of the air.

1 l. H_2O at 16° dissolves 13.1 mg. CaCO_3 . (Schlössing, C. R. 74. 1552.)

Calculated from electrical conductivity of $\text{CaCO}_3 + \text{Aq.}$, 1 pt. CaCO_3 is sol. in 99,500 pts. H_2O at 8.7° , and 80,040 pts. at 23.8° . (Holle-mann, Z. phys. Ch. 12. 125.)

By continued boiling $\text{CaH}_2(\text{CO}_3)_2$, 36 mg. CaCO_3 remain in solution. (Weltzien, A. 186. 165.)

Solubility in H_2O at different pressures.

Pressure in atmos.	Solubility
1	1079
2	1403
4	1820
6	2109

(Engel, C. R. 101. 949.)

100 pts. H_2O dissolve 0.0005 pt. (calculated as CaO) from pptd. CaCO_3 , and 0.0027 pt. from calcspar. (Lubavin, J. russ. Soc. 24. 389.)

1 l. H_2O dissolves 13 mg. CaCO_3 at 18° . (Kohlrausch, Z. phys. Ch. 1893, 12. 241.)

1 l. CO_2 free water dissolves 17.4 mg. CaO or 31.0 mg. CaCO_3 . (Gothe, Ch. Z. 1915, 39. 305.)

CaCO_3 dissolves in 9662 pts. H_2O at 12° . (Pollacci, C. C. 1896, II. 946.)

1 l. H_2O free from CO_2 dissolves 9.6 mg. CaCO_3 . (McCoy and Smith, J. Am. Chem. Soc. 1911, 33. 473.)

Found dissolved in 10,000 pts. sea water. (Davy.)

Pptd. amorphous CaCO_3 dissolves in 1600 pts. sea water. Pptd. crystalline CaCO_3 dissolves in 8000 pts. sea water. (Irvine and Young, Chem. Soc. 56. 344.)

Artificial sea water sat. with CO_2 dissolves CaCO_3 corresponding to 57.27 mg. of combined CO_2 per litre at 15° .

Sea water which contains 52–55 mg. neutral combined CO_2 per litre must be sat. with CaCO_3 . (Cohen, Chem. Soc. 1900, 78 (2) 725.)

For action of $\text{H}_2\text{CO}_3 + \text{Aq.}$ see *Calcium hydrogen carbonate*.

Sol. in H_2SO_4 , even when native. Sol. in acids generally. When treated with acids in closed vessels effervescence ceases on increase of pressure, but is renewed at once on removing it. (Link, 1814.)

Unacted upon by conc. HNO_3 , even when boiling, as $\text{Ca}(\text{NC}_2)_2$ is insol. in conc. HNO_3 .

Not decomp. by mixture of 1 pt. H_2SO_4 and 5 pts. absolute alcohol, but immediately by HNO_3 + absolute alcohol.

Not decomp. by absolute alcoholic solutions of oxalic, racemic, tartaric, citric, or glacial acetic acids. (Babington and Phillips, 1816.)

Unacted upon by glacial $\text{HC}_2\text{H}_3\text{O}_2$, even when boiling.

Freely pptd. CaCO_3 is sol. in cold NH_4Cl + Aq; but the solution becomes cloudy on exposure to air, a portion, however, of CaCO_3 remains dissolved, which cannot be pptd. even by boiling. If ppt. is washed and allowed to stand 24 hours, it is not as sol. in NH_4Cl as at first, but natural CaCO_3 is not wholly insol. in NH_4Cl + Aq; it is, however, much less sol. than MgCO_3 . (Vogel, J. pr. 7. 453.)

Sol. in boiling NH_4Cl + Aq with evolution of NH_3 . (Demarcay, 1834.)

When NH_4OH + Aq, incompletely sat. with CO_2 , is mixed with CaCl_2 + Aq, no ppt. occurs even during several days, if kept in a closed vessel; and only a slight ppt. if the mixture is exposed to the air, but CaCO_3 is pptd. if the solution is boiled.

NH_4OH + Aq wholly sat. with CO_2 produces ppt. when mixed with CaCl_2 + Aq, but pptn. is not complete until heat is applied. Also when an excess of CaCl_2 + Aq is added to a solution of crystallized carbonate of ammonia, only a portion of the CaCO_3 is pptd. until the solution is boiled. (Vogel, 1814.)

When CaCl_2 + Aq mixed with NH_4OH + Aq is exposed to an atmos. of pure CO_2 , no ppt. occurs for several hours, but CaCO_3 is completely pptd. in several days. (Vogel.)

When recently pptd., readily sol. in NH_4Cl , and NH_4NO_3 + Aq. (Brett, 1837; Wackenroder, A. 41. 315.)

When recently pptd., readily sol. in $(\text{NH}_4)_2\text{CO}_3$, $(\text{NH}_4)_2\text{SO}_4$, NH_4NO_3 , NH_4Cl , and NH_4 succinate + Aq. (Wittstein.)

Sol. in $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$ + Aq. (Thomson.)

More sol. in NH_4Cl , or NH_4NO_3 + Aq, or in neutral potassium, or sodium salts + Aq than in H_2O . (Fresenius.)

From solutions in NH_4 salts, NH_4OH , and $(\text{NH}_4)_2\text{CO}_3$ + Aq precipitate CaCO_3 more completely than BaCO_3 . (Fresenius.)

When boiled with NH_4Cl + Aq, CaCO_3 is dissolved, and $(\text{NH}_4)_2\text{CO}_3$ given off. (D. Smith.)

CaCl_2 + Aq prevents pptn. of CaCO_3 in the cold, as do also NH_4Cl , KCl , or NaCl + Aq, but it is pptd. when boiled, if the latter solutions are not too conc. K_2SO_4 , KNO_3 , $(\text{NH}_4)_2\text{SO}_4$, or Na_2SO_4 + Aq have a similar effect. A large excess of $(\text{NH}_4)_2\text{CO}_3$ + Aq when quickly added to CaCl_2 + Aq produces

no ppt. in the cold. Na_2CO_3 , or K_2CO_3 + Aq act likewise. (Storer, Am. J. Sci. (2) 25. 41.)

1 g. CaCO_3 requires 13.98 g. NH_4Cl , 8.380 g. $(\text{NH}_4)_2\text{SO}_4$, or 14.438 g. NH_4NO_3 to effect solution. (Bertrand, Monit. Sci. (3) 10. 477.)

Less sol. in Na than in NH_4 salts, but more than in K salts. (Berthelot.)

When NH_4OH + Aq, partially neutralized by CO_2 , is mixed with CaO_2H_2 + Aq, no cloudiness appears until the mixture is boiled; when more CO_2 has been added to NH_4OH + Aq, a ppt. appears at first, which disappears and only reappears on addition of much CaO_2H_2 + Aq; but NH_4OH + Aq does not dissolve pptd. CaCO_3 . (Vogel.)

Solubility in NH_4 salts + Aq at 25°.

NH_4 salt	Millimols NH_4 salt per l.	Millimols CaO dissolved per l.
NH_4Cl	1000	6.770
	500	5.008
	250	3.724
	125	2.743
NH_4NO_3	500	5.267
	250	3.830
	125	2.779
	62.5	2.004
Triammonium citrate	500	66.87
	250	39.80
	125	22.64
	62.5	14.92

(Rindell, Z. phys. Ch. 1909, 70. 454.)

Solubility of CaCO_3 in NH_4Cl + Aq at 12–18°. Time, 98 days.

g. per l. of sat. solution	
NH_4Cl	CaCO_3
53.5	0.423
100	0.609
200	0.645

(Cantoni and Goguelia, Bull. Soc. 1905, (3) 33. 27.)

Solubility of CaCO_3 in NH_4NO_3 + Aq at 18°.

g. per l. of sat. solution	
NH_4NO_3	CaCO_3
0	0.131
5	0.211
10	0.258
20	0.340
40	0.462
80	0.584

(Berju and Kosminiko, Landw. Vers. Sta. 1904, 60. 422.)

CaO₂H₂+Aq dissolves a little CaCO₃. (Welter and Berthollet, 1789.)

CaO₂H₂+Aq retains a little CaCO₃ in solution at ordinary temperature, which is pptd. on boiling. (Eliot and Storer, Proc. Am. Acad. 1860, 5. 63.)

CaO₂H₂+Aq, mixed with dil. NaOH, KOH, or NH₄OH+Aq, gives no immediate ppt. when CO₂ is passed through it, unless boiled.

Sol. in boiling MgCl₂+Aq even when dilute. Cousté.)

Not decomp. when boiled with K₂SO₄, Na₂SO₄, CaSO₄, MgSO₄, and Na₂B₄O₇+Aq; but partially decomp. by boiling with (NH₄)₂SO₄, K₂SO₃, Na₂SO₃, (NH₄)₂SO₃, Na₂HPO₄, (NH₄)₂HPO₄, K₂HPO₄, Na₂HPO₄, (NH₄)₂HPO₄, K₂HAsO₄, Na₂AsO₄, K₂C₂O₄, (NH₄)₂C₂O₄, NaF, and K₂CrO₄+Aq. With the NH₄ salts the decomposition is complete. (Dulong, A. ch. 82. 286.)

Not decomp. by alkali sulphates+Aq. (Malaguti.)

Precipitation of CaCO₃ is much hindered by alkali citrates or metaphosphates.

Solubility in KCl+Aq at 25°.

Sp. gr. 25°/25°	% KCl	% CaCO ₃
1.000	0.00	0.0013
1.024	3.90	0.0078
1.046	7.23	0.0078
1.072	11.10	0.0076
1.092	13.82	0.0072
1.101	15.49	0.0076
1.122	18.21	0.0070
1.133	19.84	0.0072
1.179	26.00	0.0060

(Cameron and Robinson, J. phys. Chem. 1907, 11. 578.)

Solubility in NaCl+Aq in contact with CO₂ free air at 25°.

Sp. gr. 25°/25°	g. per 100 g. H ₂ O	
	NaCl	CaCO ₃
1.0079	1.60	0.0079
1.0314	5.18	0.0086
1.0466	9.25	0.0094
1.0734	11.48	0.0104
1.0949	16.66	0.0106
1.1346	22.04	0.0115
1.1794	30.50	0.0119

(Cameron, Bell and Robinson, J. phys. Ch. 1907, 11. 396.)

Solubility of CaCO₃ in NaOH+Aq.

Solvent	1 litre dissolves	
	at 18°	at 95°-100°
H ₂ O	12.8 mg. CaCO ₃	20.7 mg. CaCO ₃
ca. 0.0001 n. NaOH	8.7 " "	9.6 " "
ca. 0.001 n. NaOH	4.2 " "	6.9 " "
ca. 0.01 n. NaOH	4.3 " "	5.7 " "

(Le Blanc, Z. anorg. 1906, 51. 185.)

Solubility in K₂SO₄+Aq at 25°.

Sp. gr. 25°/25°	% K ₂ SO ₄	% CaCO ₃
1.010	1.60	0.0104
1.021	3.15	0.0116
1.033	4.73	0.0132
1.048	6.06	0.0148
1.061	7.85	0.0168
1.069	8.88	0.0192
1.083	10.18	0.0192
1.084	10.48	0.0188

(Cameron and Robinson, J. phys. Chem. 1907, 11. 578.)

The solubility of CaCO₃ in Na₂SO₄+Aq in equilibrium with air steadily increases with increasing amounts of CaSO₄ in the solution up to saturation point of the CaSO₄. In the presence of solid CaSO₄ the solubility of CaCO₃ is much decreased. (Cameron and Seidell, J. phys. Chem. 1902, 6. 56.)

See under CaH₂(CO₃).

Solubility in Na₂SO₄+Aq in contact with CO₂ free air at 25°.

g. per 100 g. H ₂ O	
Na ₂ SO ₄	CaCO ₃
0.97	0.0151
1.65	0.0180
4.90	0.0262
12.69	0.0313
14.55	0.0322
19.38	0.0346
23.90	0.0360

(Cameron, Bell and Robinson, J. phys. Ch. 1907, 11. 396.)

Solubility in salts+Aq.

g. salt added per litre	mg. CaO dissolved per litre
0.000	17.4
0.585 g. NaCl	20.05
1.17 g. " "	24.9
2.93 g. " "	31.1
0.85 g. NaNO ₃	24.35
1.70 " "	27.7
4.25 " "	34.5
0.805 g. Na ₂ SO ₄ , 10H ₂ O	25.95
1.61 g. " "	31.15
4.03 g. " "	40.7
0.53 g. Na ₂ CO ₃	8.4
1.06 g. " "	7.2
2.65 g. " "	4.4
0.55 g. CaCl ₂ , 6H ₂ O	9.0
1.10 g. " "	8.4
2.75 g. " "	8.4

The solubility of CaCO₃ in CO₂-free water

is therefore increased by the addition of NaCl, NaNO₃ or Na₂SO₄, 10H₂O, but decreased by the addition of Na₂CO₃ or CaCl₂, 6H₂O.

(Gothe, Ch. Z. 1915, 39. 306.)

Sol. in ferric chloride or nitrate+Aq with evolution of CO₂ and pptn. of Fe₂O₃.H₂O (Fuchs, 1831); also in chlorides or nitrates of Al, Mn, Cr, or U, but not in FeCl₂+Aq.

Sol. in cold SnCl₄+Aq with pptn. of SnO₂. Insol. in conc. Na₂SO₄, MgSO₄, BaCl₂, MgCl₂, Pb(NO₃)₂, or AgNO₃+Aq. (Karsten.)

Abundantly sol. when freshly precipitated in CaCl₂+Aq, and MgSO₄+Aq. (Hunt.)

Absolutely insol. at 15-19° in BaO₃H₂+Aq; also on boiling.

1 l. H₂O containing 3-4 g. MgSO₄ dissolves 1-2 g. CaCO₃, and also 1 g. MgCO₃. (Hunt, Am. J. Sci (2) 26. 109.)

100 pts. NaCl+Aq (2.525% NaCl) dissolve 0.0037 pt. (calculated as CaO) pptd. CaCO₃, and 0.0053 pt. calcspar. (Lubavin, J. russ. Soc. 24. 389.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

Insol. in liquid CO₂. (Büchner, Z. phys. Ch. 1906, 54. 674.)

Alcohol dissolves traces of CaCO₃. (Grischow.)

Sol. in Na citrate+Aq. (Spiller.)

Sol. in Ca succinate+Aq. (Barreswill.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Amorphous. Solubility in H₂O cannot be determined because of its instability. (Kendall, Phil. Mag. 1912, (6) 23. 972.)

Min. Calcite. In contact with air free from CO₂, 1 l. H₂O dissolves at:

25° 50° 100°
0.01433 0.01504 0.01779 g. calcite.

(Kendall, Phil. Mag. 1912, (6) 23. 964.)

In contact with air containing 3.7 pts. CO₂ per 10,000, the solubility of calcite in H₂O was found to be 0.04608 g. per l. at 25° and 0.02925 g. per l. at 50°. (Kendall, Phil. Mag. 1912, (6) 23. 973.)

Min. Aragonite. In contact with air free from CO₂, 1 l. H₂O dissolves at:

25° 50° 100°
0.01528 0.01617 0.01902 g. aragonite.

(Kendall, Phil. Mag. 1912, (6) 23. 964.)

+5H₂O. Efflorescent.
+6H₂O. (Pelouze.)

Calcium hydrogen carbonate, CaH₂(CO₃)₂.
Known only in aqueous solution.
CaCO₃ dissolves in CO₂+Aq.

CaCO₃ is sol. in 1428 pts. H₂O sat. with CO₂ at 0°, and 1136 pts. at 10°. (Lassaigne, J. ch. med. 4. 312.)
Bineau could dissolve, even in large quantities of H₂O sat. with CO₂, only 1/3 enough CaCO₃ to form CaH₂(CO₃)₂.
Chalk dissolves in 994.5 pts. H₂O sat. with CO₂, while Iceland spar requires 3149 pts. (Bischof.)
CaCO₃ is sol. in 1015 pts. H₂O sat. with CO₂ at 21° and 748.3 mm. (Warrington, Chem. Soc. 6. 296.)

Solubility of CaCO₃ in CO₂+Aq at p pressure in atmospheres. CaO+CO₂=mg. CO₂ and CaC dissolved, corresponding to CaCO₃=mg. CaCO₃.

p	CaO+CO ₂	CaCO ₃
0.000504	60.96	74.6
0.000808	72.11	85.0
0.00333	123	137.2
0.03187	218.4	223.1
0.0282	310.4	296.5
0.05008	408.5	360
0.1422	...	533
0.2538	1072	663.4
0.4167	1500	787.5
0.5533	1846	885.5
0.7297	2270	972
0.9841	2864	1086

(Schlösing, C. R. 74. 1522.)

With high pressure, 1 l. H₂O containing CO₂ dissolves at most 3 g. CaCO₃. This maximum is reached at 5° under 4 atmospheres' pressure; at 10-13° under 5 atmospheres; and at 20° under 7 atmospheres. (Caro, Arch. Pharm. (3) 4. 145.)

CaCO₃ is sol. in about 1000 pts. H₂C'C₃+Aq, and solubility is considerably increased by Na₂SO₄ or MgSO₄.

1000 pts. H₂O sat. with CO₂ dissolve pts. Carrara marble at t°, and B=height of barometer in millimetres.

t°	B	Pts. CaCO ₃	t°	B	Pts. CaCO ₃
7.5	754	1.224	22.0	746	0.920
8.5	752	1.202	26.0	740	0.875
9.5	754	1.115	26.5	743	0.860
20.5	741	0.975	27.0	741	0.885
21.5	744	0.935	28.0	737	0.770

Or, from 7.5-9.5°, 1000 pts. H₂O sat. with CO₂ dissolve 1.181 pts. CaCO₃; from 20.5-22°; 0.9487 pt. CaCO₃; from 26-28°, 0.855 pt. CaCO₃.

Other varieties of CaCO_3 are dissolved as follows in 1000 pts. H_2O sat. with CO_2 .

Variety	t°	B	Pts. CaCO_3
Lüneburg chalk . . .	18	740	0.835
Pptd. CaCO_3 . . .	18	740	0.950
Isoland spar. . . .	18	735	1.970
Calcite	12	754	1.223
Traversella	12	754	.212
Dolomite, semi-transparent	11.5	749	0.654
Dolomite, opaque, in small crystals . . .	11.5	755	0.725
Dolomite, opaque, in large crystals . . .	11	746	.224
Dolomite, transparent, in large crystals . .	11	749	1.073
Oolitic limestone . .	15	747	.252
Dolomitic limestone .	15.5	740	.573

(Cossa, Z. anal. 8. 145.)

Solubility of CaCO_3 in H_2O containing CO_2 at various pressures.

CO_2 pressure in atm.	1	2	4	6
Solubility	1079	1403	1820	2109

(Engel, C. R. 1885, 101. 951.)

1 l. H_2O dissolves 0.3850 g. $\text{CaH}_2(\text{CO}_3)_2$ at 15°. (Treadwell, Z. anorg. 1898, 17. 186.)
1 l. of sat. $\text{CaH}_2(\text{CO}_3)_2 + \text{Aq}$, obtained from pure or impure limestone, contains 1.13-1.17 g. CaCO_3 at 15°. (Treadwell, Z. anorg. 1898, 17. 189.)

Solubility of $\text{CaH}_2(\text{CO}_3)_2$ in H_2O containing CO_2 at 15°.

% carbonic acid in gas at 0° and 760 mm.	Hg-partial pressure mm.	Free carbonic acid mg.	mg. $\text{CaH}_2(\text{CO}_3)_2$ in 100 cc. of the solution	mg. Ca
8.94	67.9	157.4	187.2	46.2
6.04	45.9	86.3	175.5	43.3
5.45	41.4	52.8	159.7	39.4
2.18	16.6	48.5	154.0	38.0
1.89	14.4	34.7	149.2	36.8
1.72	13.1	24.3	133.1	32.9
0.79	6.0	14.5	124.9	30.8
0.41	3.1	4.7	82.1	20.3
0.25	1.9	2.9	59.5	14.7
0.08	0.6	...	40.2	9.9
....	38.5	9.5
....	38.5	9.5
....	38.5	9.5

(Treadwell and Reuter, Z. anorg. 1898, 17. 185.)

1 l. H_2O sat. with carbonic acid dissolves 1.30 g. CaCO_3 at 13.2°; 1.45 g. at 2.8°. (Treadwell, Z. anorg. 1898, 17. 189.)
At 30° C. in equilibrium with the air, not more than 3 per cent of the calcium present is combined as CaCO_3 . At lower temperatures and lesser concentrations the percentage of normal carbonate is even less, and practically all the calcium present is combined as $\text{Ca}(\text{HCO}_3)_2$. (Cameron and Briggs, J. phys. Chem. 1901, 5. 549.)
With pressures less than 4.5 atmospheres of CO_2 no other than normal calcium carbonate or a hydrate of the normal carbonate can exist as the solid phase at 0°. (Cameron, J. phys. Chem. 1908, 12. 566.)

Solubility in H_2O in contact with air, containing CO_2 with varying partial pressures at t°.

P = partial pressure of CO_2 .

t = 15°		
P	g. per l.	
	CaCO_3	CO_2
0.8	0.193	0.117
1.5	0.193	0.152
1.7	0.238	0.135
6.8	0.445	0.327
9.9	0.627	0.456
13.6	0.723	0.560
14.6	0.686	0.623
31.6	1.050	1.117
t = 25°		
P	g. per l.	
	CaCO_3	CO_2
0.7	0.159	0.091
1.6	0.177	0.111
4.6	0.341	0.208
7.8	0.446	0.301
16.5	0.539	0.522
30.1	0.743	0.715
35.5	0.755	0.803
t = 40°		
P	g. per l.	
	CaCO_3	CO_2
0.6	0.136	0.078
1.7	0.143	0.085
2.9	0.175	0.106
3.5	0.232	0.169
7	0.284	0.234
14.9	0.384	0.293
22.2	0.427	0.333
31.7	0.480	0.476

Similar results at 20°, 30°, and 35° are also given.
(Leather and Sen, Mem. Dept. Agric. (India) Chem. Ser. 1909, 1. 117; Seidell, Solubilities, 1919.)

Solubility of calcite in H_2O at 25° , in contact with CO_2 under varying pressures. Approximate pressure of CO_2 in atmos-

g. per l. sat. solution		Solid phase
H_2CO_3	$\text{Ca}(\text{HCO}_3)_2$	
0.22	0.67	CaCO_3
2.3	1.58	"
20.6	3.62	"
27.5	4.04	"
34.1	4.21	$\text{Ca}(\text{HCO}_3)_2$
53.2	4.22	"

by and Smith, J. Am. Chem. Soc. 1911, 33. 468.)

H_2O dissolves 2.3374 g. CaCO_3 at 5° a CO_2 pressure of 2 atmos. (Ehlert, Elektrochem. 1912, 18. 727.)

Solubility data for calcite in H_2O containing CO_2 , with and without the presence of NaCl are given by Seyler and Lloyd (Chem. 1909, 95. 346.)

Critical analysis and recalculation of results of Schloesing and others is given by Treadwell (J. Am. Chem. Soc. 1915, 37. 2001). CaCO_3 is not dissolved by CO_2 and H_2O in presence of MgCO_3 . (Leather and Sen, C. A. 181.)

Solubility of 1/10-normal $\text{NaCl} + \text{Aq}$ dissolves 0.1 g. $\text{CaH}_2(\text{CO}_3)_2$ at 15° . (Treadwell and Reuter, Z. anorg. 1898, 17. 193.)

Solubility of $\text{CaH}_2(\text{CO}_3)_2$ in $\text{NaCl} + \text{Aq}$ saturated with carbonic acid at 15° , containing 5 g. NaCl per l. of $\text{NaCl} + \text{Aq}$.

mm. Hg = partial pressure	mg. free CO_2	mg. $\text{CaH}_2(\text{CO}_3)_2$ in 100 cc. of the solution	mg. Ca
128.8	132.5	218.4	53.9
87.2	110.1	214.3	52.9
46.1	23.5	149.2	36.8
24.0	13.5	118.3	29.2
3.8	2.7	73.9	18.2
3.4	0.3	49.0	12.1
...	...	34.9	8.6
...	...	33.7	8.3
...	...	32.9	8.1
...	...	33.2	8.2

Treadwell and Reuter, Z. anorg. 1898, 17. 193.)

Solubility in $\text{NaCl} + \text{Aq}$ at 25°C . and in equilibrium with air.

$\text{Ca}(\text{HCO}_3)_2$		NaCl	
Grams per liter	Reacting wts. per liter	Grams per liter	Reacting wts. per litre
0.1046	0.00065	0.000	0.000
0.1770	0.00110	9.720	0.168
0.2051	0.00128	21.010	0.362
0.2152	0.00134	30.301	0.522
0.2252	0.00140	50.620	0.872
0.2212	0.00138	69.370	1.195
0.2172	0.00135	98.400	1.695
0.1971	0.00123	147.400	2.540
0.1569	0.00095	234.500	4.040
0.1227	0.00076	262.300	4.520

(Cameron and Seidell, J. phys. Chem. 1902, 6. 51.)

Solubility in various salts + Aq under a CO_2 pressure of 2 atmos. at 5° .

Salt	g. salt per 1000 g. H_2O	g. CaCO_3 sol. in 1 l. of solvent
H_2O	...	2.3374
$\text{MgCl}_2 + 6\text{H}_2\text{O}$	6.08	2.3518
	50.0	3.4045
	86.0	4.0826
	350.0	3.3009
	700.0	2.7357
	1150.0	2.2054
	1725.0	1.7058
	2300 (sat.)	1.4060
NaCl	27.96	3.2796
	50.0	3.7399
	86.0	3.7828
	106.9	3.6900
	175.6	3.3495
	263.4	2.8107
	351.2	2.1625 at 8°
$\text{MgSO}_4 + 7\text{H}_2\text{O}$	105.3 (14°) sat. at 14°	2.1768
		0.91356
$\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$	137.7 (14°) sat. at 14°	1.4060
		1.9199

(Ehlert and Hempel, Z. Elektrochem. 1912, 18. 727.)

Solubility of CaCO_3 in $\text{KCl} + \text{Aq}$ at 25° sat. with CO_2 at atmospheric pressure.

% KCl	% CaCO_3
3.90	0.145
7.23	0.150
11.10	0.166
13.82	0.165
15.49	0.167
18.21	0.154
19.84	0.140
26.00	0.126

(Cameron and Robinson, J. phys. Chem. 1907, 11. 579.)

Solubility in NaCl+Aq in contact with CO₂ at atmospheric pressure at 25°.

g. per 100 g. H ₂ O	
NaCl	CaCO ₃
1.45	0.150
5.69	0.160
11.08	0.174
15.83	0.172
19.62	0.159
29.89	0.123
35.85	0.103

(Cameron, Bell and Robinson, J. phys. Ch. 1907, 11. 396.)

Solubility in K₂SO₄+Aq, sat. with CO₂ at atmospheric pressure and 25° temp.

% SO ₂	% CaO
0.69	0.69
1.37	0.69
1.67	0.47 *
2.18	0.30 *
2.99	0.24 *

* Solid phase, CaSO₄, K₂SO₄.
(Cameron and Robinson.)

Solubility in Na₂SO₄+Aq at 24° in equilibrium with air.

Total Ca calc. as Ca(HCO ₃) ₂ . Grams per liter	Ca actually dissolved as Ca(HCO ₃) ₂ . Grams per liter	Na ₂ SO ₄ . Grams per liter
0.0925	0.0925	0.000
0.1488	0.1488	2.800
0.1729	0.1729 +	5.235
0.2330	0.2210	11.730
0.3240	0.3020	36.860
0.3960	0.3440	74.010
0.4580	0.3660	116.100
0.5630	0.3940	184.200
0.5910	0.4060	213.700
0.6650	0.4300	255.900

(Cameron and Seidell, J. phys. Chem. 1902, 6. 53.)

Data are also given for solubility of CaCO₃ in NaCl+Na₂SO₄+Aq, and CaCO₃+CaSO₄ in NaCl+Na₂SO₄+Aq. (Cameron, Bell and Robinson.)

Calcium copper uranium carbonate, CaCO₃, 3CuCO₃, 4U(CO₃)₂+24H₂O.

Sol. in acids.

Calcium lead carbonate, xCaCO₃, yPbCO₃.

Min. *Plumbocalcite*.

Calcium magnesium carbonate, CaCO₃, MgCO₃.

Min. *Dolomite*. 1 l. H₂O sat. with CO₂ at 18° and 750 mm. dissolves 0.31 g. dolomite. (Cossa, B. 2. 697.)

Not obtained by evaporating solution, but can be crystallized from CO₂+Aq between 100° and 200°. (Hoppe-Seyler.)

Dolomite is dissolved by CO₂ and H₂O, but solution is prevented partially by CaCO₃ and wholly by MgCO₃. (Leather and Sen, C. A. 1915. 181.)

Insol. in cold dil. acids. (Dolomieu, J. Phys. 39. 1.)

Insol. in cold acetic acid. (Forchhammer.)

Calcium potassium carbonate, CaK₂(CO₃)₂.

Decomp. by H₂O. (Reynolds, Chem. Soc. 1898, 73. 265; Bütschli, C. A. 1907. 2223). 2CaCO₃, 3K₂CO₃+6H₂O. (Bütschli.)

Calcium sodium carbonate, CaNa₂(CO₃)₂.

Anhydrous. Decomp. by H₂O.

+2H₂O. (Bütschli, C. A. 1907. 2223.)

+5H₂O. Min. *Gaylussite*. Sparingly sol. in H₂O.

Calcium uranyl carbonate, CaCO₃, UO₂CO₃+20H₂O.

Min. *Liebigite*. Sol. in HCl+Aq.

+xH₂O. Decomp. by H₂O. (Blinkoff, Dissert. 1900.)

2CaO, 4UO₂, 3CO₂+24H₂O. Decomp. by H₂O. (Blinkoff, Dissert. 1900.)

Calcium carbonate chloride, CaCO₃, CaCl₂+6H₂O.

Sol. in H₂O with immediate decomp. (Fritzsche, J. pr. 83. 213.)

Cerous carbonate, Ce₂(CO₃)₃+5, and 9H₂O.

Insol. in H₂O, and solution of CO₂ in H₂O. (Vauquelin.)

Somewhat sol. in (NH₄)₂CO₃+Aq. (Jolin.)

Insol. in neutral salt solutions and neutral alkali carbonates+Aq; easily sol. in SO₂+Aq. (Berthier, A. ch. (3) 7. 77.)

Ceric carbonate, Ce(CO₃)₂+½H₂O.

Precipitate. (Hisinger, A. ch. 94. 108.)

Insol. in H₂O. Sol. in slight traces in Na₂CO₃+Aq; sl. sol. in NaHCO₃+Aq, and in (NH₄)₂CO₃+Aq. (Rose.)

Cerous lanthanum carbonate fluoride.

Min. *Butnaxite*, *Hamartite*, *Hydrofluocerite*. Slowly decomp. by HCl+Aq, easily by H₂SO₄.

Cerous potassium carbonate, Ce₂(CO₃)₃, K₂CO₃+3H₂O.

Ppt. (Jolin.)

Ce₂(CO₃)₃, K₂CO₃+12H₂O. Ppt.

Sol. in 30% K₂CO₃+Aq. (Meyer, Z. anorg. 1904, 41. 103.)

Mercuric potassium carbonate, $\text{Ce}_2\text{O}_3(\text{CO}_3)_3$, $4\text{K}_2\text{CO}_3 + 12\text{H}_2\text{O}$.

Crystalline. Sl. sol. in H_2O containing CO_2 ; sol. in dil. H_2SO_4 with decomp. ob, C. R. 1899, **128**. 1098.)

Mercurous sodium carbonate, $\text{Ce}_2(\text{CO}_3)_3$, $2\text{Na}_2\text{CO}_3 + 2\text{H}_2\text{O}$.

Ppt. (Jolin.)

$2\text{Ce}_2(\text{CO}_3)_3$, $3\text{Na}_2\text{CO}_3 + 24\text{H}_2\text{O}(?)$. Ppt. easily decomp. (Meyer, Z. anorg. 1904, **41**. 13.)

Mercurous carbonate, CrCO_3 .

Sol. in much H_2O ; sl. sol. in $\text{KHCO}_3 + \text{Aq}$. Moberg, J. pr. **44**. 328; Moissan, A. ch. (5) **1**. 199.)

Mercuric carbonate, basic, Cr_2O_3 , 2CO_2 .

Precipitate. (Parkmann, Sill. Am. J. (2) **4**. 321.)

Cr_2O_3 , $\text{CO}_2 + 4\text{H}_2\text{O}$. Insol. in H_2O ; sol. in acids; when freshly pptd. is sol. in K_2CO_3 , or $\text{NH}_4)_2\text{CO}_3 + \text{Aq}$, and still more sol. in $\text{KOH} + \text{Aq}$. (Meissner.)

Insol. in ethyl acetate (Naumann, B. **910**, **43**. 314); methyl acetate. (Naumann, **3**. 1909, **42**. 3790.)

$2\text{Cr}_2\text{O}_3$, $\text{CO}_2 + 6\text{H}_2\text{O}$. Precipitate. (Langlois, A. ch. (3) **48**. 502.)

Chromous potassium carbonate,

CrCO_3 , $\text{K}_2\text{CO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O when freshly prepared; slowly polymerizes; stable in dry air, decomp. in moist air; sol. in acids with decomp. (Baugé, C. R. 1898, **126**. 1568.)

Chromous sodium carbonate, $\text{CrNa}_2(\text{CO}_3)_2 + \text{H}_2\text{O}$.

Decomp. when heated. In Aq. solution, passes into the hydrate containing 10 mols. H_2O . (Baugé, C. R. 1897, **125**. 1179.)

$+10\text{H}_2\text{O}$. Very sol. in cold H_2O ; Aq. solution decomp. below 100° ; effloresces in the air; sol. in $\text{HCl} + \text{Aq}$ and $\text{H}_2\text{SO}_4 + \text{Aq}$. (Baugé, C. R. 1897, **125**. 1178.)

Cobaltous carbonate, basic, 5CoO , $2\text{CO}_2 + 4\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in $(\text{NH}_4)_2\text{SO}_4$, $(\text{NH}_4)_2\text{CO}_3$, NH_4NO_3 , and $\text{NH}_4\text{Cl} + \text{Aq}$.

Sol. in cold NH_4NO_3 , and $\text{NH}_4\text{Cl} + \text{Aq}$. (Brett, 1837.)

Sol. in $\text{CO}_2 + \text{Aq}$, and acid alkali carbonates $+ \text{Aq}$, from which it is pptd. on boiling. Very sl. sol. in conc. Na_2CO_3 , or $\text{K}_2\text{CO}_3 + \text{Aq}$; largely sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$, and partly sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Berzelius.)

Not pptd. from solutions containing Na citrate. (Spiller.)

4CoO , $\text{CO}_2 + 4\text{H}_2\text{O}$. Ppt. (Beetz.)

$+3\text{H}_2\text{O}$. (Meigen, C. C. 1905, I. 1363.)

Cobaltous carbonate, basic, 3CoO , $\text{CO}_2 + 2\text{H}_2\text{O}$.

(Meigen, C. C. 1905, I. 1363.)

$3\text{H}_2\text{O}$. (Rose, Pogg. **84**. 551.)

3CoO , $2\text{CO}_2 + 4\text{H}_2\text{O}$. (Bratin, Z. anal. **6**. 76.)

2CoO , $\text{CO}_2 + 3\frac{1}{2}\text{H}_2\text{O}$. Converted into 5CoO , $2\text{CO}_2 + 4\text{H}_2\text{O}$ by H_2O . (Beetz.)

Cobaltous carbonate, CoCO_3 .

Anhydrous. Not attacked by cold conc. HCl , or $\text{HNO}_3 + \text{Aq}$. (Senarmont, A. ch. (3) **30**. 129.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 827.)

Min. *Sphaerocobaltite*. Sl. attacked by cold HNO_3 , or $\text{HCl} + \text{Aq}$.

$+2\frac{1}{3}\text{H}_2\text{O}$. Sol. in acids. (Deville, A. ch. (3) **33**. 95.)

$+6\text{H}_2\text{O}$. (Deville.)

Decomp. by H_2O with formation of a basic carbonate. (Berzelius.)

Cobaltous potassium carbonate, CoCO_3 , $\text{K}_2\text{CO}_3 + 4\text{H}_2\text{O}$.

Decomp. by H_2O . (Deville, A. ch. (3) **33**. 90.)

Ppt. Decomp. by H_2O . (Reynolds, Chem. Soc. 1898, **73**. 264.)

CoCO_3 , $\text{KHCO}_3 + 4\text{H}_2\text{O}$. Decomp. by H_2O . (Deville.)

Cobaltous sodium carbonate, CoCO_3 , $\text{Na}_2\text{CO}_3 + 4\text{H}_2\text{O}$, and $10\text{H}_2\text{O}$.

Decomp. by H_2O . (Deville, A. ch. (3) **33**. 75.)

Cupric carbonate, basic.

The compounds produced by pptn. of copper solutions by carbonates are unstable and possess varying solubilities in solutions of CO_2 . On treatment with solutions of CO_2 , these substances pass over into an apparently stable compound possessing a definite solubility in solutions of CO_2 of definite concentration, which solubility increases with the concentration of CO_2 . Solubility of this compound in various salts $+ \text{Aq}$ is recorded. (Free, J. Am. Chem. Soc. 1908, **30**. 1374.)

8CuO , $\text{CO}_2 + 5\text{H}_2\text{O}$. (Deville, A. ch. (3) **33**. 75.)

6CuO , CO_2 . (Field, Chem. Soc. **14**. 70.)

3CuO , $\text{CO}_2 + 2\text{H}_2\text{O}$. (Favre, A. ch. (3) **10**. 119.)

5CuO , $2\text{CO}_2 + 6\text{H}_2\text{O}$. (Struve.)

2CuO , $\text{CO}_2 + \text{H}_2\text{O}$. Insol. in H_2O ; easily sol. in acids, even $\text{H}_2\text{SO}_4 + \text{Aq}$; sl. sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$, 30,720 pts. of the solution containing 1 pt. CuO . (Jahn.) Sol. in 4690 pts. $\text{H}_2\text{CO}_3 + \text{Aq}$ sat. at 4-6 atmos. pressure. (Wagner.) Sol. in 3833 pts. sat. $\text{H}_2\text{CO}_3 + \text{Aq}$. (Lassaigne, J. ch. méd. **4**. 312.)

Sol. in NH_4 salts $+ \text{Aq}$. Partially sol. in Na_2CO_3 , or $\text{K}_2\text{CO}_3 + \text{Aq}$, and more sol. in

NaHCO₃, or **KHCO₃** + Aq; sol. in (NH₄)₂CO₃ + Aq. (Favre, A. ch. (3) 10. 18.)

Less sol. in (NH₄)₂CO₃ + Aq than CuO in NH₄OH + Aq. (Thomson, 1831.) Sol. in KCN + Aq. (Berzelius.) Sol. in NH₄Cl, or NH₄NO₃ + Aq. (Brett.)

Sol. in ferric salts with pptn. of Fe₂O₃·H₂O.

Insol. in liq. NH₃. (Franklin and Kraus, Am. Ch. J. 1898, 20. 827.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Sol. in ethyl amine carbonate + Aq. (Wurtz.)

Sol. in cane sugar + Aq. (Peschier, Repert. 1820, 6. 85.)

Not pptd. from solutions containing sodium citrate. (Spiller.)

Insol. in pyridine. (Schroeder, Dissert. 1901.)

Min. *Malachite*. Sol. in acids, and NH₄OH + Aq.

+ 2H₂O. (Favre.)

8CuO, 5CO₂ + 7H₂O. (Gröger, Z. anorg. 1900, 24. 137.)

3CuO, 2CO₂ + H₂O. Insol. in H₂O. Sol. in NH₄OH + Aq, also in hot conc. NaHCO₃ + Aq.

Min. *Azurite*.

Copper potassium carbonate, basic, 8CuO, 2K₂CO₃, 7CO₂ + 17H₂O.

Ppt.; decomp. by H₂O. (Gröger, B. 1901, 34. 430.)

Mixture. (Wood and Jones, C. A. 1907. 2667.)

5CuO, 4CO₂, K₂CO₃ + 10H₂O. Decomp. by H₂O. (Deville, A. ch. (3) 33. 102.)

Cupric potassium carbonate, CuCO₃, K₂CO₃.

Decomp. by H₂O. (Wood and Jones, C. A. 1907. 2667.)

+ H₂O. (Wood and Jones.)

+ 4H₂O. Decomp. by H₂O. (Reynolds, Chem. Soc. 1898, 73. 263.)

Could not be obtained. (Wood and Jones.)

2CuCO₃, K₂CO₃ + 4H₂O. Decomp. by H₂O. (Wood and Jones.)

Cupric sodium carbonate, CuCO₃, Na₂CO₃.

Not decomp. by cold H₂O. (Debray, C. R. 49. 218.)

+ 3H₂O.

Cupric zinc carbonate, 2CuO, 3ZnO, 2CO₂ + 3H₂O, or 3CuO, 9ZnO, 4CO₂ + 8H₂O.

Min. *Aurichalcite*. Easily sol. in HCl + Aq.

Cupric carbonate ammonia (cuprammonium carbonate), CuCO₃, 2NH₃.

Decomp. by H₂O. Insol. in alcohol and ether. Sol. in (NH₄)₂CO₃ + Aq. (Favre, A. ch. (3) 10. 116.)

Didymium carbonate, Di₂(CO₃)₂ + H₂O, 6H₂O.

Insol. in H₂O. Only traces dissolve in CO₂ + Aq. Insol. in solutions of alkali carbonate or bicarbonates + Aq. (Marignac, A. ch. (3) 38. 166.) Very sl. sol. in conc. NH₄Cl + Aq. (Rose.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+ 8H₂O. (Cleve, Bull. Soc. (2) 43. 363.)

Didymium potassium carbonate, Di₂(CO₃)₂, K₂CO₃ + 4H₂O.

Insol. in H₂O. (Cleve, Bull. Soc. (2) 43. 363.)

+ 12H₂O. (Cleve.)

Didymium sodium carbonate, 2Di₂(CO₃)₂, 3Na₂CO₃ + 9H₂O.

Ppt. (Cleve.)

Di₂(CO₃)₂, 2Na₂CO₃ + 8H₂O. Ppt. (Cleve.)

Dysprosium carbonate, Dy₂(CO₃)₂ + 4H₂O.

Insol. in H₂O. (Jantsch, B. 1911, 44. 1277.)

Erbium carbonate, Er₂O₃, 2CO₂ + 2H₂O.

Insol. in H₂O. (Höglund.)

Erbium sodium carbonate, Er₂(CO₃)₂, 5Na₂CO₃ + 36H₂O.

Efflorescent. Decomp. by H₂O.

Gadolinium carbonate, basic, Gd(OH)CO₃ + H₂O.

Ppt. (Benedicks, Z. anorg. 1900, 22. 417.)

Glucinum carbonate, basic, 3GdO, CO₂; 4GdO, CO₂; 5GdO, CO₂ + 5H₂O, etc.

Not perceptibly sol. in H₂O or H₂CO₃ + Aq. Decomp. by boiling H₂O. Easily sol. in acids. Sol. in NH₄ salts, and KOH, or NaOH + Aq. Sol. in alkali carbonates, especially (NH₄)₂CO₃ + Aq. (Vauquelin.) Sl. sol. in K₂CO₃ + Aq. When solution in (NH₄)₂CO₃ is boiled, a more basic carbonate is pptd. (Rose.)

Glucinum carbonate, GdCO₃ + 4H₂O.

Efflorescent. Sol. in 278 pts. H₂O. (Klatso, J. pr. 106. 242.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 828.)

No definite carbonate of glucinum exists. (Cameron, J. phys. Chem. 1908, 12. 572.)

um potassium carbonate, 3GlCO_3 ,
 I_2CO_3 .
ly sol. in H_2O , but decomp. by boiling.
y.) Less easily sol. in alcohol.

carbonate, $\text{In}_2(\text{CO}_3)_3$.
Insol. in K_2CO_3 , or $\text{Na}_2\text{CO}_3 + \text{Aq}$.
 $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Winkler, J. pr.

erric) carbonate, basic.
 O_3 , $\text{CO}_2 + 12\text{H}_2\text{O}$. (Wallace, Chem.
358. 410.)
 O_3 , $\text{CO}_2 + 4\text{H}_2\text{O}$, and $8\text{H}_2\text{O}$. (Barrat,
l. 110.)
 I_2O . (Wallace.)
 O_3 , $\text{CO}_2 + 1\frac{1}{2}\text{H}_2\text{O}$. (Rother, Pharm.
is. (3) 4. 576.)
 I_3 , CO_2 . (Parkmann, Sill. Am. J. (2)
)
e and other similar basic salts are ppts.,
decomp. on standing into $\text{Fe}_2\text{O}_3\cdot\text{H}_2\text{O}$.

errous) carbonate, FeCO_3 .
l. in H_2O .
in acids, even in $\text{H}_2\text{CO}_3 + \text{Aq}$.
Carbonate, ferrous hydrogen.
. *Siderite*, *Spathic ore*. Sl. attacked by
ds. Sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$ under pressure.
l. in NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett.)
 I_2O . Sl. sol. in H_2O ; easily sol. in acids;
 $\text{H}_2\text{CO}_3 + \text{Aq}$.
in $\text{NH}_4\text{Cl} + \text{Aq}$. Sol. in ferric salts + Aq
volution of CO_2 and pptn. of $\text{Fe}_2\text{O}_3\cdot\text{H}_2\text{O}$.
ble in an aqueous solution of cane sugar.

lubility in salts + Aq free from CO_2 .

	g. salt per 1000 g. H_2O	1 l. of solvent dissolves g. FeCO_3
	351.2	0.35042
+	2300.0	4.2049
$\frac{1}{4}$ I_2O	137.7 sat. at $+14^\circ$	0.70085 0.93444
$\frac{1}{4}$ +	105.3 sat. at $+18^\circ$	1.4667 2.9334

lert, Z. Elektrochem. 1912, 18. 728.)

errous) hydrogen carbonate,
 $\text{FeH}_2(\text{CO}_3)_2(?)$.
wn only in aqueous solution.
conducting CO_2 at ordinary pressure
h H_2O , in which Fe is suspended, a solu-
ntaining 9.1 pts. FeCO_3 to 10,000 pts.
obtained. (v. Hauer, J. pr. 81. 391.)

100 pts. $\text{H}_2\text{CO}_3 + \text{Aq}$ dissolve 0.72 pt.
 FeCO_3 . (Wagner.)
 FeCO_3 dissolves in 1381 pts. H_2O saturated
with CO_2 , under a pressure of 4–6 atmos-
pheres. (Wagner, J. B. 1867. 135.)
1 l. H_2O dissolves 6.1907 g. FeCO_3 (pure)
under a CO_2 pressure of 2 atmos. (Ehlert,
Z. Elektrochem. 1912, 18. 728.)

Solubility in various salts + Aq in presence of
 CO_2 under pressure of 2 atmos.

Salt	With CO_2 of 2 atmos. pressure	
	g. salt per 1000 g. H_2O	1 l. of solvent dissolves g. FeCO_3
H_2O	6.1907
NaCl	50
	106.9
	175.6
	263.4
	351.2
$\text{MgCl}_2 +$ $6\text{H}_2\text{O}$	86.9	5.8403
	700.0	4.5553
	1150.0	4.4587
	1437.5	4.6934
	1725.0	5.3975
	2300.0	9.0524
Na_2SO_4 $+10\text{H}_2\text{O}$	137.7	7.9428
	sat. at $+14^\circ$	9.5780
$\text{MgSO}_4 +$ $7\text{H}_2\text{O}$	105.3	6.2423
	sat. at $+18^\circ$	7.3922

(Ehlert, Z. Elektrochem. 1912, 18. 728.)

A bicarbonate of ferrous iron is not formed
under pressures of CO_2 up to 5 atmospheres
at 0° . (Cameron, J. phys. Chem. 1908, 12.
571.)

Iron (ferrous) magnesium carbonate, FeCO_3 ,
 MgCO_3 .
Min. *Pistomesite*.
 $\text{FeCO}_3, 2\text{MgCO}_3$. Min. *Mesitite*.

Iron (ferrous) potassium carbonate,
 $\text{FeK}_2(\text{CO}_3)_2 + 4\text{H}_2\text{O}$.
Ppt. Decomp. by H_2O . (Reynolds, Chem.
Soc. 1898, 73. 265.)

Lanthanum carbonate, $\text{La}_2(\text{CO}_3)_3 + \text{H}_2\text{O}$,
 $3\text{H}_2\text{O}$, and $8\text{H}_2\text{O}$.
Insol. in H_2O . $\text{CO}_2 + \text{Aq}$ dissolves traces.
Insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.
Insol. in acetone. (Naumann, B. 1904, 37.
4329.)
Min. *Lanthanite*.

Lanthanum potassium carbonate, $\text{La}_2(\text{CO}_3)_3$, $\text{K}_2\text{CO}_3 + 12\text{H}_2\text{O}$.

Sol. in 30% $\text{K}_2\text{CO}_3 + \text{Aq}$. (Meyer, Z. anorg. 1904, 41. 101.)

Lanthanum sodium carbonate, $2\text{La}_2(\text{CO}_3)_3$, $3\text{Na}_2\text{CO}_3 + 20\text{H}_2\text{O} (?)$.

Ppt. Easily decomp. (Meyer, Z. anorg. 1904, 41. 102.)

Lead carbonate, basic, 2PbCO_3 , PbO_2H_2 ; 5PbCO_3 , $3\text{PbO}_2\text{H}_2$; 3PbCO_3 , PbO_2H_2 ; 5PbCO_3 , PbO_2H_2 .

White Lead. Insol. in H_2O . Nearly insol. in $\text{H}_2\text{CO}_3 + \text{Aq}$, even under pressure. Sol. in dil., insol. in conc. $\text{KOH} + \text{Aq}$. Insol. in normal, or acid alkali carbonates + Aq . (Böttger.)

Sol. in cold dil. $\text{NH}_4\text{Cl} + \text{Aq}$. (Brett.)

PbCO_3 , PbO_2H_2 . Very sl. sol. in H_2O . (Yorke.)

2PbCO_3 , PbO_2H_2 .

Solubility is less than 0.0002 millimol Pb in 1 liter H_2O at 18° . (Pleissner, C. C. 1907, II. 1056.)

When not exposed to air, sol. in 32,000 pts. $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ (0.2 g. per l.); 26,000 pts. $\text{KNO}_3 + \text{Aq}$ (0.2 g. per l.); 23,000 pts. $\text{CaCl}_2 + \text{Aq}$ (0.2 g. per l.); 4600 pts. $\text{NH}_4\text{NO}_3 + \text{Aq}$ (0.2 g. per l.); 4300 pts. H_2O sat. with CO_2 .

When exposed to air in beakers, sol. in 43,000 pts. $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ (0.2 g. per l.); 43,000 pts. $\text{KNO}_3 + \text{Aq}$ (0.2 g. per l.); 26,000 pts. $\text{CaCl}_2 + \text{Aq}$ (0.2 g. per l.); 26,000 pts. $\text{NH}_4\text{NO}_3 + \text{Aq}$ (0.2 g. per l.); 4300 pts. H_2O sat. with CO_2 (0.2 g. per l.). (Muir, Chem. Soc. 31. 664.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

3PbO , $4\text{PbCO}_3 + 2\text{H}_2\text{O}$. Ppt. (Strömholm, Z. anorg. 1904, 38. 446.)

Lead carbonate, PbCO_3 .

Sol. in 50,551 pts. H_2O at ordinary temp.

Sol. in 23,450 pts. H_2O with little ammonium acetate, carbonate, and free ammonia; and in somewhat less H_2O , containing much ammonium nitrate with carbonate and free ammonia. (Fresenius, A. 59. 124.)

Calculated from electrical conductivity of $\text{PbCO}_3 + \text{Aq}$, 1 l. H_2O dissolves 3 mg. PbCO_3 at 10° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

Solubility is 0.0002 millimol. Pb in 1 liter H_2O at 18° . (Pleissner, C. C. 1907, II. 1056.)

Sl. sol. in H_2O . 1.5×10^{-3} g. are contained in 1 l. of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 604.)

Easily sol. in acids, even $\text{HC}_2\text{H}_3\text{O}_2$; but not decomp. by conc. $\text{HNO}_3 + \text{Aq}$ on account of insolubility of $\text{Pb}(\text{NO}_3)_2$ in $\text{HNO}_3 + \text{Aq}$. Insol. in a mixture of 1 pt. H_2SO_4 and 6 pts. absolute alcohol, or in an alcoholic solution of racemic or tartaric acids.

Insol. in $\text{H}_2\text{CO}_3 + \text{Aq}$. (Jahn, A. 28. 11) Very sl. sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$, but solution prevented by traces of various salts. (Terman.) Sol. in 7144 pts. sat. $\text{H}_2\text{CO}_3 + \text{Aq}$. (Lassaigne, J. ch. méd. 4. 312.) H_2O sat. with CO_2 under 4–6 atmos. pressure dissolves 0.5 pt. PbCO_3 . (Wagner, Z. anal. 6. 167.)

Solubility of PbCO_3 in $\text{H}_2\text{CO}_3 + \text{Aq}$ at :

mg. per l.	
CO_2	PbCO_3
0	1.75
2.8	6
5.4	7
14.4	8.2
26	9.9
43.5	10.9
106	15.4

(Pleissner, C. C. 1907, II. 1056.)

Sol. in $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$, and $\text{NH}_4\text{Cl} + \text{Aq}$ (Weppen, 1837.) Sol. in $\text{KOH} + \text{Aq}$; not solutely insol. at ord. temp. in an excess K_2CO_3 , or $\text{Na}_2\text{CO}_3 + \text{Aq}$, and still more sol. at 100° ; but absolutely insol. in NaHCO_3 , or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Rose.) In $\text{NH}_4\text{OH} + \text{Aq}$; sol. in KOH or $\text{NaOH} + \text{Aq}$; decomp. by boiling $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$. (Berzelius.)

Sl. decomp. (Persoz), not at all decomp. (Malaguti) by alkali sulphates + Aq .

Partially decomp. by boiling with K_2SO_4 , $(\text{NH}_4)_2\text{SO}_4$, CaSO_4 , MgSO_4 , Na_2HPO_4 , $\text{NaNH}_4\text{HPO}_4$, K_2SO_4 , Na_2SO_4 , $(\text{NH}_4)_2\text{SO}_4$, Na_2HPO_4 , $\text{Na}_2\text{B}_4\text{O}_7$, K_2AsO_4 , $\text{K}_2\text{C}_2\text{O}_4$, $\text{Na}_2\text{C}_2\text{O}_4$, NaF , $\text{K}_2\text{CrO}_4 + \text{Aq}$. With the NH_4 salts, decomp. is complete. (Dulong, A. ch. 290.)

Easily sol. in hot $\text{NH}_4\text{Cl} + \text{Aq}$. (Berzelius.)

When 1 mol. PbCO_3 is boiled with 1 l. $\text{K}_2\text{C}_2\text{O}_4$, 15% of the PbCO_3 is decomp.; with 1 mol. K_2CO_3 , 93.28% is decomp. (Malaguti.)

Not decomp. by $\text{K}_2\text{SO}_4 + \text{Aq}$. (Rose.)

Insol. in liquid NH_3 . (Franklin, Am. J. 1898, 20. 828.)

Sol. in an aqueous solution of acetic acid. (Mercer, 1844.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Not pptd. in presence of Na citrate. (Spiller.)

Min. (cerussite).

Lead sodium carbonate, 4PbCO_3 , Na_2CO_3 .

Insol. in H_2O . (Berzelius, Pogg. 47. 19)

Lead carbonate bromide, PbCO_3 , PbBr_2 .

Insol. in H_2O . (Storer's Dict.)

carbonate chloride, PbCO_3 , PbCl_2 .
in H_2O . (Miller, Chem. Soc. (2) 8.)

Phosgenite. Easily sol. in acids.

carbonate iodide, PbCO_3 , PbI_2 .
in H_2O . (Poggiale.)

carbonate sulphate, PbCO_3 , PbSO_4 .
Lanarkite. Sol. in HNO_3 + Aq with
of PbSO_4 .
 O_3 , PbSO_4 . Min. *Leadhillite*. As

carbonate, Li_2CO_3 .

pts. H_2O dissolve 1 pt. Li_2CO_3 . (Vau-
A. ch. 7. 284.)

pts. H_2O at 13° dissolve 0.769 pt.
at 102° , 0.778 pt. Li_2CO_3 . (Kremers,
3. 48.)

pts. H_2O , cold or hot, dissolve 1.2 pts.
(Troost, A. ch. (3) 51. 103.)

pts. H_2O dissolve 1.4787 pts. at 15° ,
at 100° . (Draper, C. N. 55. 169.)

pts. H_2O dissolve pts. Li_2CO_3 at t° .

Pts. Li_2CO_3	t°	Pts. Li_2CO_3
1.539	75	0.866
1.406	100	0.728
1.329	102	0.796
1.181	..	.

pt. is dissolved at 102° in less than
and 0.955 in 1 hour. (Beketow, J.
c. 1884. 591.)

solution at 15° has sp. gr. 1.014, and
1 g. Li_2CO_3 to 70 g. H_2O , while solu-
at 0° has sp. gr. 1.0168 and contains
 CO_2 in 64.6 g. H_2O . By long spon-
evaporation at 15° a solution can
ined of 1.0278 sp. gr. containing 1 g.
in 45.57 g. H_2O . (Flückiger, Arch.
(3) 25. 549.)

illing for an instant with H_2O a solu-
obtained, which has sp. gr. 1.0074 and
1 g. Li_2CO_3 to 139 g. H_2O . (Flücki-
h. Pharm. (3) 26. 543.)

7 mol. is sol. in 1 l. H_2O at 25° . (Roth-
Z. phys. Ch. 1909, 69. 531.)

Li_2CO_3 + Aq contains at:

15° 75°
723 0.833% by wt. Li_2CO_3 .

chugreff, Z. anorg. 1914, 86. 159.)

solution boils at 102° . (Kremers.)

sol. in CO_2 + Aq than in H_2O . 100

pts. sat. CO_2 + Aq dissolve 5.25 pts. Li_2CO_3 .
(Troost.) See LiHCO_3 .
Sol. in NH_4 salts + Aq.

Solubility in salts + Aq at 25° .

C = concentration of salt solution in g.-
equiv. per l.
S = solubility of Li_2CO_3 in g.-equiv. per l.

Salt	C	S
KNO_3	0 25	0 3647
	0 50	0 3688
	0 75	0 3676
	1 00	0 3656
	1 50	0 3490
	2 00	0 3268
KCl	0 10	0 3553
	0 25	0 3590
	0 50	0 3782
	0 75	0 3832
	1 00	0 3835
	1 50	0 3731
NaCl	2 00	0 3558
	0 10	0 3569
	0 25	0 3601
	0 50	0 3867
	0 75	0 3956
	1 00	0 3946
K_2SO_4 2	1 50	0 3601
	2 00	0 3776
	0 25	0 4028
	0 50	0 4356
	1 00	0 4860
Na_2SO_4 2	0 50	0 4411
	1 00	0 4926
	2 00	0 5534
NH_4Cl	0 10	0 3902
	0 25	0 4677
	0 50	0 5659
	0 75	0 6270
	1 00	0 6810
	1 50	0 7463
$(\text{NH}_4)_2\text{SO}_4$ 2	2 00	0 7739
	4 00	0 7881
	0 25	0 5059
	0 50	0 7863
	1 00	0 9804
	1 50	1 109
KClO_4	2 00	1 174
	0 1	0 3500
	0 2	0 3570
	0 4	0 3616

(Geffcken, Z. anorg. 1905, 43. 198.)

Insol. in liquid NH_3 . (Franklin, Am. Ch.
J. 1898, 20. 828.)

Solubility in organic compds. + Aq. at 25°. Solubility in H ₂ O at 25° = 0.1687 mols. litre.			Solubility in organic compds. + Aq. at 25° -Continued.		
Organic compd.	Normality of the solution	Mol. Li ₂ CO ₃ sol. in 1 l.	Organic compd.	Normality of the solution	Mol. Li ₂ CO ₃ sol. in 1 l.
Methyl alcohol	0.250	0.1604	Thio-urea	0.125	0.1667
	0.5	0.1529		0.250	0.1643
	1.0	0.1394		0.5	0.1600
Ethyl alcohol	0.125	0.1614		1.0	0.1523
	0.250	0.1555	Dimethyl-pyrone	0.125	0.1562
	0.5	0.1417		0.250	0.1460
	1.0	0.1203		0.5	0.1284
Propyl alcohol	0.125	0.1604		1.0	0.0992
	0.250	0.1524	Ammonia	0.125	0.1653
	0.5	0.1380		0.250	0.1630
	1.0	0.1097		0.5	0.1577
Tertiary Amyl alcohol	0.125	0.1564		1.0	0.1466
	0.250	0.1442	Diethylamine	0.125	0.1589
	0.5	0.1224		0.250	0.1481
	1.0	0.0899		0.5	0.1283
Acetone	0.125	0.1600		1.0	0.0937
	0.250	0.1515	Pyridine	0.125	0.1503
	0.5	0.1366		0.250	0.1347
	1.0	0.1104		0.5	0.1091
Ether	0.125	0.1580		1.0	0.1091
	0.250	0.1476	Piperidine	0.125	0.1584
	0.5	0.1300		0.250	0.1320
Formaldehyde	0.125	0.1668		0.5	0.1009
	0.250	0.1653		1.0	0.1009
	0.5	0.1606	Urethane	0.125	0.1604
	1.0	0.1531		0.250	0.1525
Glycol	0.125	0.1660		0.5	0.1377
	0.250	0.1629		1.0	0.1113
	0.5	0.1565	Acetamide	0.250	0.1614
	1.0	0.1472		0.5	0.1520
Glycerine	0.125	0.1670		1.0	0.1358
	0.250	0.1647	Acetonitrile	0.125	0.1618
	0.5	0.1613		0.250	0.1556
	1.0	0.1532		0.5	0.1178
Mannitol	0.125	0.1705		1.0	0.1178
	0.250	0.1737	Mercuric cyanide	0.125	0.1697
	0.5	0.1778		0.250	0.1704
Glucose	0.125	0.1702	(Rothmund, Z. phys. Ch. 1909, 69. 531.)		
	0.250	0.1728	Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3601.)		
	0.5	0.1752	Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)		
	1.0	0.1778	Lithium hydrogen carbonate, LiHCO ₃ .		
Sucrose	0.125	0.1693	100 pts. H ₂ O dissolve 5.501 pts. at 13°.		
	0.250	0.1689	(Bewad, B. 17. 406 R.)		
	0.5	0.1661			
	1.0	0.1557			
Urea	0.125	0.1686			
	0.250	0.1673			
	0.5	0.1643			
	1.0	0.1605			

arbonate, basic, $\text{Mg}_3\text{C}_2\text{O}_7 + 10\text{H}_2\text{O} + 2\text{CO}_2 + 3\text{H}_2\text{O}$ or $2\text{MgCO}_3 + 2\text{H}_2\text{O}$. (Fritzsche, Pogg. 37.)

alba, 3MgCO_3 , $\text{Mg}(\text{OH})_2 + 2\text{O}_2$, $\text{Mg}(\text{OH})_2 + 5\text{H}_2\text{O}$, or $(\text{OH})_2 + 7\text{H}_2\text{O}$.

in H_2O . Sol. in 10,000 pts.

pts. cold, and 9000 pts. hot

containing CO_2 .

sol. in acids.

dil. $\text{HCl} + \text{Aq}$.

NH_4 sulphate, nitrate, or sucrose in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Witt-cold Na_2CO_3 , K_2CO_3 , K_2SO_4 , $+ \text{Aq}$ (Longchamp); also in separating out on heating. 7. 455.) Slowly sol. in conc. or $\text{ZnSO}_4 + \text{Aq}$. (Karsten.)

$+ \text{Aq}$. (Dulong.)

salts $+ \text{Aq}$ with evolution of of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$. (Fuchs.)

ng Co , Ni , Zn , Mn , or Cu rides $+ \text{Aq}$.

omagnesianite, 4MgO , $3\text{CO}_2 +$

sol. in considerable amount in $\text{MgH}_2(\text{CO}_3)_2 + \text{Aq}$. (Engel,)

arbonate, MgCO_3 .

Insol. in H_2O . 1 l. H_2O diss. MgCO_3 . (Chevalet, Z. anal. 5071 pts. H_2O at 15° . (Krem-combines with H_2O to form $+ 5\text{H}_2\text{O}$, which are less than anhydrous salt. (Engel,)

topic. About 20 g. are sol. in H_2O . (C. R. 1899, 129. 598.)

sol. in 1 l. H_2O at 15° .

sol. in 1 l. H_2O at 15° with $+ \text{CO}_2$ equal to zero.

are sol. in 1 l. H_2O at 15° with $+ \text{CO}_2$ equal to 1 atmos.

Reuter, Z. anorg. 1898, 17.

sol. in 1 l. of CO_2 -free water. 1915, 39. 306.)

H_2O in equilibrium with CO_2 .

CO_2 , $\text{Mg}(\text{HCO}_3)_2$, and CO_2 at

Mg as MgCO_3 (grams per liter)	Mg as $\text{Mg}(\text{HCO}_3)_2$ (grams per liter)
0.01205	0.01205
0.06820	0.06314
0.08676	0.07411

Solubility in H_2O in equilibrium with $\text{Mg}(\text{HCO}_3)_2$ and CO_2 —Continued.

System: MgCO_3 , $\text{Mg}(\text{HCO}_3)_2$, and CO_2 at 30°C .

Total Mg (gram-atoms per liter)	Mg as MgCO_3 Per cent	Mg as $\text{Mg}(\text{HCO}_3)_2$ Per cent
0.00100	50.00	50.00
0.00545	51.92	48.08
0.00667	53.93	46.07

Total salts in solution. Grams per liter	MgCO_3		$\text{Mg}(\text{HCO}_3)_2$	
	Grams per liter	Per cent	Grams per liter	Per cent
0.1144	0.0418	36.5	0.0726	63.5
0.6174	0.2368	38.2	0.3806	61.7
0.7479	0.3012	40.3	0.4467	59.7

In a solution near the saturation point and in equilibrium with atmospheric air upwards of 50 per cent of the magnesium is in the form of the normal carbonate. When the solution is brought in contact with the solid phase, the proportion of the base combined as normal carbonate falls to about 34 per cent, or lower. (Cameron and Briggs, J. phys. Chem. 1901, 5. 552-3.)

For solubility in $\text{H}_2\text{CO}_3 + \text{Aq}$, see *Magnesium hydrogen carbonate*.

Scarcely acted upon by $\text{HCl} + \text{Aq}$. (Senarmont.)

The solubility of MgCO_3 in $\text{NaCl} + \text{Aq}$ when in contact with ordinary air increases with increasing concentration of NaCl up to a maximum, and then decreases. (Cameron and Seidell, J. phys. Chem. 1903, 7. 579.)

Solubility of MgCO_3 in salts $+ \text{Aq}$ in equilibrium with an atmosphere free from CO_2 .

$\text{NaCl} + \text{Aq}$; $t = 23^\circ$.

Weight of liter of solution	Grams NaCl per liter	Grams MgCO_3 per liter	Reacting weights NaCl per liter	Reacting weights MgCO_3 per liter
996.92	0.0	0.176	0.000	0.00210
1016.82	28.0	0.418	0.482	0.00500
1041.09	59.5	0.527	1.025	0.00630
1070.50	106.3	0.585	1.831	0.00699
1094.53	147.4	0.544	2.539	0.00650
1142.48	231.1	0.460	3.981	0.00550
1170.14	272.9	0.393	4.701	0.00470
1199.28	331.4	0.293	5.709	0.00350

(Cameron and Seidell, J. phys. Chem. 1903, 7. 585.)

Solubility of MgCO_3 in salts + Aq.—Continued
 $\text{Na}_2\text{SO}_4 + \text{Aq}; t = 24^\circ$.

Weight of liter of solution	Grams Na_2SO_4 per liter	Grams MgCO_3 per liter	Reacting wts. Na_2SO_4 per liter	Reacting wts. MgCO_3 per liter
1000.00	0.00	0.216	0.000	0.00258
1021.24	25.12	0.586	0.178	0.00700
1047.60	54.76	0.828	0.388	0.00990
1100.00	95.68	1.020	0.678	0.01219
1133.85	160.80	1.230	1.140	0.01470
1157.34	191.90	1.280	1.360	0.01530
1206.03	254.60	1.338	1.804	0.01600
1223.91	278.50	1.338	1.973	0.01600
1241.99	305.10	1.388	2.162	0.01660

 $t = 35.5^\circ$.

Weight of liter of solution	Grams Na_2SO_4 per liter	Grams MgCO_3 per liter	Reacting weights Na_2SO_4 per liter	Reacting weights MgCO_3 per liter
995.15	0.32	0.131	...	0.00156
1032.89	41.84	0.577	0.296	0.00689
1067.23	81.84	0.753	0.579	0.00881
1094.77	116.56	0.904	0.826	0.01080
1120.38	148.56	0.962	1.052	0.01149
1151.70	186.70	1.047	1.323	0.01251
1179.82	224.00	1.068	1.587	0.01300
1196.32	247.20	1.100	1.751	0.01314
1236.52	199.20	1.130	2.120	0.01350

(Cameron and Seidell.)

 $\text{Na}_2\text{CO}_3 + \text{Aq}; t = 25^\circ$.

Weight of liter of solution	Grams Na_2CO_3 per liter	Grams MgCO_3 per liter	Reacting weights Na_2CO_3 per liter	Reacting weights MgCO_3 per liter
996.84	0.00	0.223	0.000	0.00266
1019.89	23.12	0.288	0.220	0.00344
1047.72	50.75	0.510	0.482	0.00620
1062.47	86.42	0.879	0.820	0.01027
1118.91	127.30	1.314	1.209	0.01570
1147.66	160.80	1.636	1.526	0.01955
1166.05	181.90	1.972	1.727	0.02357
1189.38	213.20	2.317	2.024	0.02770

(Cameron and Seidell.)

Solubility in salts + Aq.

g. salt added per litre	mg. MgCO_3 dissolved per litre
0.0	94.4
0.585 g. NaCl	128.3
1.17 g. "	134.4
2.93 g. "	120.95
0.85 g. NaNO_3	122.85
1.70 g. "	138.80
4.25 g. "	137.20

Solubility in salts + Aq.—Continued

g. salt added per litre	mg. MgCO_3 dissolved per litre
0.805 g. $\text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$	145.05
1.61 g. "	162.05
4.03 g. "	150.75
0.53 g. Na_2CO_3	98.6
1.06 g. "	53.5
2.65 g. "	15.7
0.51 g. $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$	47.0
1.02 g. "	39.5
2.55 g. "	35.3

The solubility of MgCO_3 in CO_2 -free w is increased by the addition of NaCl , Na or $\text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$ but decreased by the a tion of Na_2CO_3 or $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$.

(Gothe, Ch. Z. 1915, 39. 306.)

Insol. in liquid NH_3 . (Franklin, Am. J. 1898, 20. 828.)

Insol. in acetone. (Naumann, B. 1: 37. 4329.)

Insol. in acetone and in methylal. (mann, C. C. 1899, II. 1014.)

Insol. in methyl acetate (Naumann, 1909, 42. 3790); ethyl acetate. (Naum B. 1904, 37. 3601.)

Min. *Magnesite*. Very sl. attacked warm conc $\text{HCl} + \text{Aq}$. 100 pta. H_2O diss 0.0027 pt., calculated as MgO . (Lubavin

Solution in H_2O contains 0.018 g. Mg 0.065 g. CO_2 per l. at 20° . (Wells, J. Chem. Soc. 1915, 37. 1705.)

Solution in H_2O containing 27.2 g. N per l. contains 0.028 g. Mg and 0.086 g. per l. at 20° . (Wells, J. Am. Chem. 1915, 37. 1705.)

 $+ \text{H}_2\text{O}$.

$+ 2\text{H}_2\text{O}$. Decomp. by suspension in l into basic salt. (Engel, C. R. 100. 911.)

$+ 3\text{H}_2\text{O}$. Small quantities of this salt wholly dissolved by much H_2O . (Bineau

The solution contains in 100 pta. at—

0°	6.5°	8°	16°
0.15	0.153	0.155	0.179

pta. $\text{MgCO}_3 + 3\text{H}$
(Norgaard, 1850.)

Decomp. by boiling H_2O into a basic in salt and CO_2 . 100 pta. H_2O dissolve 0.1 pt. at 19° . (Fritzsche, Pogg. 37. 304.)

Sol in 48 pta. H_2O , and decomp. by h amt. (Fourcroy.)

100 pta. H_2O dissolve 0.1518 pt. at 19° sol in 658 pta. H_2O at 19° . (Beckurts, J. 1881. 212.)

100 pta. H_2O dissolve 0.0612 pt., calcula as MgO . (Lubavin, J. russ. Soc. 24. 389.)

Solution in H_2O contains 0.36 g. Mg 1.01 g. CO_2 per l. at 20° . (Wells, J. Chem. Soc. 1915, 37. 1707.)

Solubility in H_2O sat. with CO_2 has b determined at 20° , 25° , 30° , 34° and 39°

at CO₂ pressures corresponding with 0.5 to 30.3% CO₂ in the gas phase. (Leather and Sen, Chem. Soc. 1915, 108 (2) 13.)
Easily sol. in acids, even when dil
Not decomp. by 1 pt. H₂SO₄+6 pts. alcohol, or by alcoholic solutions of glacial acetic, racemic, or tartaric acids, but is slowly decomp. by alcoholic solution of citric acid, or HNO₃+abs. alcohol. (Butini, 1827.)
100 pts. NaCl+Aq (2.525%) dissolve 1.1250 pt., calculated as MgO. (Lubavin.)
1% Na₂CO₃+Aq, when mixed with 1% MgSO₄+Aq, cause no ppt., but 1.5-2% solutions ppt. this salt. (Brandes, 1825.)
More sol. in NH₄Cl+Aq than CaCO₃. Sol. in NH₄NO₃+Aq, but less easily than in NH₄Cl+Aq.

Solubility in KHCO₃+Aq at t°. Values are given in mol./litre.

t°	K	Mg	Solid phase
15°	0.0	0.0095	MgCO ₃ +3H ₂ O
	0.0992	0.0131	"
	0.1943	0.0167	"
	0.3992	0.0211	" labile
	0.2681	0.0192	MgCO ₃ +3H ₂ O+MgCO ₃ , KHCO ₃ +4H ₂ O
	0.5243	0.0097	MgCO ₃ , KHCO ₃ +4H ₂ O.
	0.6792	0.0074	"
	0.9810	0.0028	"
25°	0.0	0.0087	MgCO ₃ +3H ₂ O
	0.0985	0.0115	"
	0.2210	0.0149	"
	0.3188	0.0175	"
	0.3434	0.0181	"
	0.4216	0.0205	" labile
	0.4985	0.0207	" "
	0.3906	0.0196	MgCO ₃ +3H ₂ O+MgCO ₃ , KHCO ₃ +4H ₂ O
	0.5893	0.0128	MgCO ₃ , KHCO ₃ +4H ₂ O
	0.6406	0.0117	"
35°	0.0	0.0071	MgCO ₃ +3H ₂ O
	0.1092	0.0098	"
	0.2001	0.0132(?)	"
	0.2811	0.0142	"
	0.3704	0.0163	"
	0.4847	0.0177	"
	0.5807	0.0198	"
	0.5088	0.0184	MgCO ₃ +3H ₂ O+MgCO ₃ , KHCO ₃ +4H ₂ O
	0.6231	0.0153	MgCO ₃ , KHCO ₃ +4H ₂ O
	0.8435	0.0119	"

The experiments were performed in such a way as to prevent, as far as possible, loss of CO₂ from the solutions. (Auerbach, Z. Elektrochem. 1904, 10. 164.)

1 l. H₂O, containing 6% MgSO₄+7H₂O and

a little NaCl, dissolves 5 g. MgCO₃. (Hunt, Sill. Am. J. (2) 42. 49.)
More sol. in cold alkali borates+Aq than in hot. (Wittstein.)
Sol. in Na citrate+Aq.
+4H₂O. Efflorescent.
+5H₂O. Two modifications.
a. Plates. Sol. in 600 pts. H₂O at 0-7°; solution gradually separates out MgCO₃+2H₂O. H₂CO₃+Aq sat. at 3-4 atmos. pressure dissolves 9% at 0-4° MgSO₄+Aq dissolves 4% moist salt at 3-4°, and it is easily sol. in Na₂CO₃, or NaHCO₃+Aq. (Norgaard.)
β. Prisms. More efflorescent than α. Sol. in 600 pts. H₂O but not in MgSO₄, or Na₂CO₃+Aq. Both forms are decomp. by boiling H₂O. (Norgaard.)

Magnesium hydrogen carbonate, MgH₂(CO₃)₂(?).

Known only in solution.
1 l. H₂CO₃+Aq sat. at 1 atmos. pressure dissolves 23.5 g. MgCO₃. (Bineau.)
1 l. carbonic acid water dissolves 0.115 g. magnesite at 18° and 0.75 m. pressure. (Cossa, B. 2. 697.)
1 pt. MgOC₃ dissolves in H₂O saturated with CO₂ at 5° and a pressure of—
1 2 3 4 5 6 atmospheres
in 161 144 134 100.7 110 76 pts. H₂O
(Merkel, Techn. J. B. 1867. 213.)

H₂CO₃+Aq sat. at 3-4 atmos. pressure and 0-4° dissolved 9% MgCO₃+5H₂O. (Norgaard.)
MgCO₃+3H₂O is sol. in 72.4 pts. H₂CC₃+Aq sat. at 20° and ord. pressure; 30.5 pts. H₂CO₃+Aq sat. at 2 atmos. pressure; 26.0 pts. H₂CO₃+Aq sat. at 3 atmos. pressure; 21.1 pts. H₂CO₃+Aq sat. at 4 atmos. pressure; 17.09 pts. H₂CO₃+Aq sat. at 5 atmos. pressure. (Beckurts, J. B. 1881. 212.)

1. H₂O sat. with CO₂ at p pressure and t° dissolves g. MgCO₃.

p atmos.	t°	g. MgCO ₃	p mm.	t°	g. MgCO ₃
1.0	19.5	27.79	751	13.4	28.45
2.1	19.5	33.11	760	19.5	25.79
3.2	19.7	37.3	762	29.3	21.95
4.7	19.0	43.5	764	46	15.7
5.6	19.2	46.2	764	62	10.4
6.2	19.2	48.51	765	70	8.1
7.5	19.5	51.2	765	82	4.9
9.0	18.7	56.59	765	91	2.4
...	765	100	0.0

(Engel and Ville, C. R. 93. 34.)

The low figures of other observers are due to their using basic carbonates. By very careful experiments it was found that 1 l. H₂O

sat. with CO_2 at 1 atmos. pressure and t° dissolved the following amts. of MgCO_3 :

t°	g. CO_2	t°	MgCO_3	t°	MgCO_3
3 5	35 6	18	22.1	40	22.1
12	26.5	30	15.8	50	9 5

(Engel, C. R. 100. 444.)

1.9540 g. are sol. 1 l. H_2O at 15° . (Treadwell and Reuter, Z. anorg. 1898, 17. 202.)

$\text{MgH}_2(\text{CO}_3)_2$ is not stable except in the presence of free CO_2 .

At 15° and 760 mm., a solution having the partial pressure of $\text{CO}_2 = 0$, contains 1.9540 g. $\text{MgH}_2(\text{CO}_3)_2$ and 0.7156 g. MgCO_3 per liter. (Treadwell and Reuter, Z. anorg. 1898, 17. 204.)

Solubility of $\text{MgH}_2(\text{CO}_3)_2$ in H_2O containing carbonic acid, at 15° .

carbonic acid in the gas at 0° and 760 mm	partial pressure mm Hg	mg free CO_2	mg. $\text{MgH}_2(\text{CO}_3)_2$ in 100 cc of the solution	mg. MgCO_3 in 100 cc of the solution	mg. Mg
18 86	143 3	119 0	1210 5		201 6
5 47	41 6	86 6	1210 5		201 6
4 45	33 8	223 5	1210 5		201 6
1 54	11 7		1076 6	77 3	201 6
1 35	10 3		762 9	76.5	149 2
1 07	8.2		595 2	80 7	122 4
0 62	4.7		366 3	70.1	86 5
0 60	4 6		341 7	75 8	78 8
0.33	2 5		263 2	74 8	65 5
0 21	1 6		222 9	77 1	59 4
0 14	1 1		216 9	71 0	56 6
0.03	0 3		203 6	71 1	54 5
			203 3	68 5	53 6
			196 0	70 2	52 9
			203 6	62 5	52 0
			195 4	61 6	51 1
			195 4	64 1	51 8

(Treadwell and Reuter, Z. anorg. 1898, 17. 200.)

No bicarbonate of magnesium is formed under pressures of CO_2 up to five atmospheres at 0° . (Cameron, J. phys. Chem. 1908, 12. 570.)

A critical analysis and recalculation of results of Engel and others is given by Johnston (J. Am. Chem. Soc. 1915, 37. 2001.)

Solubility in $\text{NaCl} + \text{Aq}$ at 23°C . in equilibrium with an atmosphere of CO_2 .

g. NaCl per liter	g. $\text{Mg}(\text{HCO}_3)_2$ per liter
7.0	30 64
56.5	30.18
119.7	27.88
163.9	24.96
224.8	20.78
306.0	10.75

(Cameron and Seidell, J. phys. Chem. 1907. 582.)

Solubility in $\text{Na}_2\text{SO}_4 + \text{Aq}$ at 23°C . in equilibrium with an atmosphere of CO_2 .

Strength of $\text{Na}_2\text{SO}_4 + \text{Aq}$	g. $\text{Mg}(\text{HCO}_3)_2$ in 100 cc
0.0	1.463
12%	1.916
saturated	1.612

(Cameron and Seidell.)

Magnesium potassium carbonate,
 $\text{MgK}_2(\text{CO}_3)_2 + 4\text{H}_2\text{O}$.

Quickly decomp. by cold H_2O . (Deville, A. ch. (3) 33. 87.)

Ppt. Decomp. by H_2O . (Reynolds, Ch. Soc. 1898, 73. 264.)

$\text{MgKH}(\text{CO}_3)_2 + 4\text{H}_2\text{O}$. Insol. in H_2O , decomp. thereby into an insol. basic Mg carbonate, and $\text{MgH}_2(\text{CO}_3)_2$ and KHCO_3 , which dissolve. (Berzelius.)

Magnesium rubidium hydrogen carbonate,
 $\text{MgRbH}(\text{CO}_3)_2 + 4\text{H}_2\text{O}$.

Decomp. in the air. (Erdmann, A. 18 294. 75.)

Magnesium sodium carbonate, MgCO_3 , Na_2CO_3 .

Quickly decomp. with H_2O . (Deville, ch. (3) 33. 89.)
 $+ 15\text{H}_2\text{O}$. (Norgaard.)

Magnesium sodium carbonate sodium chloride, MgCO_3 , Na_2CO_3 , NaCl .

Decomp. by H_2O . (de Schulten, C. 1898, 122. 1427.)

Manganous carbonate, MnCO_3 .

Permanent. Practically insol. in H_2O . Insol. in $\text{H}_2\text{CO}_3 + \text{Aq}$ and in acids generally.

1 l. H_2O dissolves 0.065 g. at 25° . (Ag and Valla, Att. Accad. Line. 1911, 20. 11. 7.) Insol. in liquid NH_3 . (Franklin, Am. J. 1898, 20. 828.)

Insol. in ethyl acetate. (Naumann, 1910, 43. 314.)

Min. *Rhodochrosite*.

$+ 1\frac{1}{2}$, or $1\text{H}_2\text{O}$. Insol. in H_2O . Sol. in acids. Sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$. 1 pt. Mn requires 2000 pts. $\text{H}_2\text{CO}_3 + \text{Aq}$ for soln.

(Lassaigne.) Sol. in 7680 pts. H_2O , and 3840 pts. H_2O containing CO_2 . (Jahn.) When freshly precipitated is sol. in NH_4 salts + Aq. (Wittstein.) Not more sol. in H_2O containing Na_2CO_3 or K_2CO_3 than in pure H_2O . (Ebelmen.) Insol. in NH_4Cl , or NH_4NO_3 + Aq. (Brett.)

Sol. in ferric salts + Aq, with evolution of CO_2 and pptn. of $Fe_2O_3 \cdot H_2O$. (Fuchs.)

Not pptd. in presence of Na citrate. (Spiller.)

Manganous potassium carbonate,
 $MnK_2(CO_3)_2 + 4H_2O$.

Ppt. Decomp. by H_2O alone.

Sl. sol. in $Mn(C_2H_3O_2)_2$ + Aq or K_2CO_3 + Aq. (Reynolds, Chem. Soc. 1898, 73. 264.)

Manganous carbonate hydroxylamine,
 $4MnCO_3, 3NH_2OH + 2H_2O$.

Ppt. Sol. in acids. (Goldschmidt and Syngros, Z. anorg. 5. 138.)

Mercurous carbonate, Hg_2CO_3 .

Ppt. Decomp. by hot H_2O . Sol. in hot or warm NH_4Cl + Aq, but less easily than mercuric carbonate; less sol. in NH_4NO_3 + Aq. (Brett, 1837.)

Sl. sol. in K_2CO_3 + Aq; partially sol. with decomp. in NH_4OH + Aq. (Wittstein.)

Mercuric carbonate, basic, $4HgO, CO_2$.

Can be washed with cold H_2O without decomp. (Millon, A. ch. (3) 19. 368.)

$3HgO, CO_2$. Insol. in cold H_2O . Sol. in CO_2 + Aq; sl. sol. in K_2CO_3 + Aq. Easily sol. in NH_4Cl + Aq. (Berzelius.)

Neodymium potassium carbonate, $Nd_2(CO_3)_3, K_2CO_3 + 12H_2O$.

Ppt. Sol. in 30% K_2CO_3 + Aq. (Meyer, Z. anorg. 1904, 41. 105.)

Neodymium sodium carbonate, $2Nd_2(CO_3)_3, 3Na_2CO_3 + 22H_2O(?)$.

Ppt. Easily decomp.

Sl. sol. in conc. Na_2CO_3 + Aq. (Meyer, Z. anorg. 1904, 41. 106.)

Nickel carbonate, basic, $3NiO, CO_2 + 5H_2O$.

Min. *Zaratite*. Easily sol. in HCl + Aq.

Pptd. nickel carbonate is a basic salt of varying composition. Insol. in H_2O or H_2CO_3 + Aq. Sol. in acids. Sol. in $(NH_4)_2CO_3$ + Aq;

very sl. sol. in Na_2CO_3 + Aq; sol. in warm NH_4Cl + Aq, and KCN + Aq. (Rose.)

Not pptd. in presence of Na citrate. (Spiller.)

Nickel carbonate, $NiCO_3$.

1 l. H_2O dissolves 0.0925 g at 25%. (Ageno and Valla, Att. Accad. Linc. 1911, 20, II. 706.)

Not attacked by cold conc. HCl , or HNO_3 + Aq. (Senarmont, A. ch. (3) 30. 138.)

+ $6H_2O$. Sol. in acids. (Deville, A. ch. (3) 35. 446.)

See also Carbonate, nickel, basic.

Nickel potassium carbonate, $NiCO_3, K_2CO_3 + 4H_2O$.

Ppt. (Deville, A. ch. (3) 33. 96.)

$NiCO_3, KHCO_3 + 4H_2O$. Decomp. by H_2O , but may be washed by $KHCO_3$ + Aq without decomp. (Rose, Pogg. 84. 566.)

Nickel sodium carbonate, $NiCO_3, Na_2CO_3 + 10H_2O$.

Ppt. (Deville.)

Nickel carbonate hydroxylamine, $2Ni(OH)_2, 4NiCO_3, 5NH_2OH + 7H_2O$.

Ppt. (Goldschmidt and Syngros, Z. anorg. 1894, 5. 143.)

$2Ni(OH)_2, 4NiCO_3, 6NH_2OH + 6H_2O$.

Ppt. (Goldschmidt and Syngros.)

Palladious carbonate, $PdCO_3, 9PdO + 10H_2O$.

Insol. in H_2O ; partly sol. in NH_4OH + Aq; sl. sol. in Na_2CO_3 + Aq; sol. in acids. (Kane, 1842.)

Potassium carbonate, K_2CO_3 .

Deliquescent. Very sol. in H_2O with evolution of heat.

Sol. in 1.05 pts. H_2O at 3°; 0.962 pt. at 6°; 0.900 pt. at 12.6°; 0.747 pt. at 25°; and 0.490 pt. at 70°. (Osann.)

Sol. in 0.92 pt. H_2O . (M. R. and P.)

Sol. in 0.922 pt. H_2O at 15°. (Gerlach.)

Sol. in 1 pt. H_2O . (Abl.)

100 pts. H_2O at 15.5° dissolve 100 pts. K_2CO_3 . (Ure's Dict.)

Solubility in 100 pts. H_2O at t°.

t°	Pts. K_2CO_3	t°	Pts. K_2CO_3	t°	Pts. K_2CO_3
0	83.12	40	106.20	80	134.25
10	88.72	50	112.90	90	143.18
20	94.06	60	119.24	100	153.66
30	100.09	70	127.10	135	205.11

(Poggiale, A. ch. (3) 8. 468.)

Solubility in 100 pts. H_2O at t°

t°	Pts. K_2CO_3	t°	Pts. K_2CO_3	t°	Pts. K_2CO_3
0	89.4	46	119	91	148
1	94	47	120	92	149
2	97	48	120	93	150
3	100	49	121	94	151
4	102	50	121	95	151
5	104	51	122	96	152
6	105	52	122	97	153
7	106	53	123	98	154
8	107	54	124	99	155
9	108	55	124	100	156
10	109	56	125	101	157
11	109	57	125	102	158
12	109	58	126	103	159
13	110	59	127	104	160
14	110	60	127	105	161
15	110	61	128	106	162
16	111	62	128	107	163
17	111	63	129	108	164
18	111	64	130	109	166
19	111	65	130	110	167
20	112	66	131	111	168
21	112	67	132	112	169
22	112	68	132	113	171
23	112	69	133	114	172
24	112	70	133	115	173
25	113	71	134	116	175
26	113	72	135	117	176
27	113	73	135	118	178
28	113	74	136	119	179
29	114	75	137	120	181
30	114	76	137	121	182
31	114	77	138	122	184
32	114	78	139	123	185
33	115	79	139	124	187
34	115	80	140	125	188
35	115	81	141	126	190
36	115	82	141	127	191
37	116	83	142	128	193
38	116	84	143	129	195
39	116	85	144	130	196
40	117	86	144	131	198
41	117	87	145	132	200
42	117	88	146	133	201
43	118	89	147	134	203
44	118	90	147	135	205
45	119

(Mulder, Scheik. Verhandel. 1864. 97.)

112 g. are sol. in 100 g. H_2O at 20° . (Frankforter, J. Am. Chem. Soc. 1914, 36. 1106.)Sp. gr. of $K_2CO_3 + Aq$ at 15° .

% K_2CO_3	Sp. gr.	% K_2CO_3	Sp. gr.
0 489	1 0018	11 748	1 1282
0 979	1 0038	12 727	1 1400
1 458	1 0108	13 706	1 1520
2 934	1 0209	14 685	1 1642
4 916	1 0401	15 664	1 1766
6 895	1 0505	16 643	1 1892
8 874	1 0611	17 622	1 2020
10 853	1 0719	18 601	1 2150
12 832	1 0829	19 580	1 2282
14 811	1 0940	20 559	1 2417
16 790	1 1052	21 538	1 2554
18 769	1 1166	22 517	1 2694

Sp. gr. of $K_2CO_3 + Aq$ at 15° —Continued.

% K_2CO_3	Sp. gr.	% K_2CO_3	Sp. gr.
23 496	1 2836	33 286	1 3917
24 475	1 2980	34 265	1 4030
25 454	1 3078	35 244	1 4147
26 432	1 3177	36 223	1 4265
27 412	1 3277	37 202	1 4384
28 391	1 3378	38 181	1 4504
29 369	1 3480	39 160	1 4626
30 349	1 3585	40 139	1 4750
31 328	1 3692	40 304	1 4817
32 307	1 3803		

(Thunersian)

Sp. gr. and boiling-point of $K_2CO_3 + Aq$.

% K_2CO_3	Sp. gr.	B.-pt.	% K_2CO_3	Sp. gr.	B.-pt.
4 7	1 06	100 56°	43 3	1 46	109 44°
9 9	1 11	100 54	45 8	1 50	111 11
13 2	1 15	101 11	48 8	1 54	112 78
16 8	1 19	101 11	52 1	1 58	114 44
20 5	1 22	101 68	56 0	1 63	116 11
24 0	1 25	102 22	60 4	1 70	117 78
27 3	1 28	102 78	65 5	1 80	119 44
30 5	1 31	103 33	71 8	1 95	122 22
33 6	1 34	104 44	79 2	2 15	125 56
36 2	1 38	105 56	84 4	2 40	129 44
39 0	1 41	107 22	100 0	2 60	137 78
41 7	1 44	108 33			

(Dalton.)

Sp. gr. of $K_2CO_3 + Aq$ at 17.5° .

% K_2CO_3	Sp. gr.	% K_2CO_3	Sp. gr.	% K_2CO_3	Sp. gr.
1	1.009	19	1.182	36	1.360
2	1.018	20	1.192	37	1.380
3	1.027	21	1.200	38	1.392
4	1.036	22	1.213	39	1.404
5	1.045	23	1.224	40	1.416
6	1.054	24	1.236	41	1.429
7	1.064	25	1.245	42	1.441
8	1.073	26	1.256	43	1.453
9	1.082	27	1.267	44	1.466
10	1.092	28	1.278	45	1.478
11	1.102	29	1.289	46	1.490
12	1.112	30	1.300	47	1.503
13	1.122	31	1.312	48	1.516
14	1.132	32	1.323	49	0.529
15	1.141	33	1.334	50	1.542
16	1.151	34	1.345	51	1.555
17	1.161	35	1.357	52	1.569
18	1.172

(Hager, Comm. 1883.)

The sp. gr. increases or diminishes between 8° and 20° by a decrease or increase of temp. of 1° by the following amounts:—

K_2CO_3	Corr.
40-50	0.0007
30-40	0.0005
20-30	0.0003
10-20	0.0002

(Hager.)

Sp. gr. of K_2CO_3 + Aq at 15°.

% K_2CO_3	Sp. gr.	% K_2CO_3	Sp. gr.
1	1.00914	28	1.27893
2	1.01829	29	1.28999
3	1.02743	30	1.30105
4	1.03658	31	1.31261
5	1.04572	32	1.32417
6	1.05513	33	1.33573
7	1.06354	34	1.34729
8	1.07396	35	1.35885
9	1.08337	36	1.37082
10	1.09278	37	1.38279
11	1.10258	38	1.39476
12	1.11238	39	1.40673
13	1.12219	40	1.41870
14	1.13199	41	1.43104
15	1.14179	42	1.44388
16	1.15200	43	1.44573
17	1.16222	44	1.46807
18	1.17243	45	1.48041
19	1.18265	46	1.49314
20	1.19286	47	1.50588
21	1.20344	48	1.51861
22	1.21402	49	1.53135
23	1.22459	50	1.54408
24	1.23517	51	1.55728
25	1.24575	52	1.57048
26	1.25681	52.024	1.57079
27	1.26787

(Gerlach, Z. anal. 8. 279.)

Sp. gr. of K_2CO_3 + Aq at 15°.

% K_2CO_3	Sp. gr.	% K_2CO_3	Sp. gr.
5	1.0449	30	1.3002
10	1.0919	40	1.4170
20	1.1920	50	1.5428

(Kohlrausch, W. Ann. 1879. 1.)

K_2CO_3 + Aq containing 10% K_2CO_3 boils at 100.8°

K_2CO_3 + Aq containing 20% K_2CO_3 boils at 102.2°

K_2CO_3 + Aq containing 30% K_2CO_3 boils at 104.5°

K_2CO_3 + Aq containing 40% K_2CO_3 boils at 108.6°

K_2CO_3 + Aq containing 50% K_2CO_3 boils at 115.2°

(Gerlach.)

Sat. K_2CO_3 + Aq containing 158 pts. K_2CO_3 to 100 pts. H_2O forms a crust at 126°; highest temp. observed 134.9°. (Gerlach, Z. anal. 26. 427.)

B.-pt. of K_2CO_3 + Aq containing pts. K_2CO_3 to 100 pts. H_2O . G = according to Gerlach (Z. anal. 26. 459); L = according to Legrand (A. ch. (2) 59. 438).

B.-pt	G	L
101°	11.5	13
102	22.5	22.5
103	32	31
104	40	38.8
105	47.5	46.1
106	54.5	53.1
107	61	59.6
108	67	65.9
109	73	71.9
110	78.5	77.6
111	83.5	83.0
112	88.5	88.2
113	93.5	93.2
114	98.5	98.0
115	103.5	102.8
116	108.5	107.5
117	113.5	112.3
118	117.5	117.1
119	122.5	122.0
120	127.5	127.0
121	132.5	132.0
122	137.5	137.0
123	142.5	142.0
124	147.5	147.1
125	152.5	152.2
126	158	157.3
127	163.5	162.5
128	169.5	167.7
129	175.5	172.9
130	181.5	178.1
131	187.5	183.4
132	193.5	188.8
133	199.5	194.2
133.3	202.5	...
134	199.6
135	205.0

When K_2CO_3 + Aq is sat. with NH_3 , two layers form. When K_2CO_3 is added to NH_4OH + Aq, it dissolves with formation of two layers and evolution of NH_3 . The same takes place also when sat. K_2CO_3 + Aq and NH_4OH + Aq are brought together. (Girard, Bull. Soc. (2) 43. 552.)

Solubility of K_2CO_3 + $KHCO_3$ in H_2O at 0°.

g. per 100 cc. solution		Sp. gr.
K_2CO_3	$KHCO_3$	
0.0	21.2	1.133
11.8	15.3	1.182
16.7	12.6	1.200
23.8	10.3	1.241
34.0	7.6	1.298
43.0	5.9	1.350
51.6	4.9	1.398
60.5	3.8	1.448
81.4	0.0	1.542

(Engel, A. ch. 1888, (6) 13. 348.)

Equilibrium between K_2CO_3 and $KHCO_3$ in H_2O and in contact with the air. System: K_2CO_3 , $KHCO_3$, and CO_2 at $25^\circ C$.

Grams atoms K per liter	Amount of solution used for titration cc.	Amount K combined as K_2CO_3 Per cent	Amount K combined as $KHCO_3$ Per cent
0.393	5	77.5	22.5
0.553	5		
	2	83.9	16.1
	5		
1.025	2	86.8	13.2
	2		
	2		
1.865	2	91.4	8.6
	2		
2.820	2	89.0	11.0
	2		
3.710	2	88.6	11.4
	2		
4.310	1	89.6	10.4
	1		
5.695	1	88.7	11.3
	1		

(Cameron and Briggs, J. phys. Chem. 1901, 5. 546.)

Solubility of $K_2CO_3 + Na_2CO_3$ in H_2O at 25° .

g. per 100 g. solution		g. per 100 g. H_2O		Solid phase
K_2CO_3	Na_2CO_3	K_2CO_3	Na_2CO_3	
32.82	0	112	0	$K_2CO_3 \cdot 2H_2O$
32.0	1.0	110.7	2.2	"
50.7	2.6	108.7	5.7	"
49.0	4.6	105.5	10.0	$K_2CO_3 \cdot 2H_2O + Na_2CO_3 \cdot 12H_2O$
46.5	4.3	94.6	8.8	$Na_2CO_3 \cdot K_2CO_3 \cdot 12H_2O$
46.2	5.2	94.8	10.6	"
41.0	6.3	77.6	11.8	"
37.7	7.0	68.3	12.6	"
31.0	10.5	53.0	17.9	"
29.8	11.3	50.5	19.1	"
25.2	14.1	41.4	23.2	"
22.4	16.6	36.8	27.3	"
19.8	18.7	32.1	30.3	"
19.1	19.7	31.2	32.1	"
15.1	23.2	24.5	37.6	$Na_2CO_3 \cdot K_2CO_3 \cdot 12H_2O + Na_2CO_3 \cdot 10H_2O$
14.5	22.8	23.1	36.4	$Na_2CO_3 \cdot 10H_2O$
10.8	22.7	16.2	34.0	"
10.7	22.4	16.0	33.5	"
4.7	21.9	6.4	29.8	"
0	22.71	0	29.37	"

(Osaka, J. Tok. Chem. Soc. 1911, 32. 870.)

Solubility of $K_2CO_3 + Na_2CO_3$ in H_2O at 24.2° .

In 1000 ccm. H_2O		Solid phase
Na_2CO_3	K_2CO_3	
28.35	Na_2CO_3
354.2	150.03	"
369.7	226.6	"
363.0	243.5	$Na_2CO_3, K_2CO_3 \cdot 6H_2O$
330.8	282.7	"
273.8	344.9	"
187.2	483.9	"
130.0	921.5	"
137.9	982.6	"
112.3	1074.0	$Na_2CO_3, K_2CO_3 \cdot 6H_2O + Na_2CO_3$
95.2	1085.1	"
25.5	1108.6	K_2CO_3
.....	1125.7	"

(Kremann and Zite'k, M. 1909, 30. 317.)

Solubility of $K_2CO_3 + Na_2CO_3$ in H_2O at 10° .

In 1000 ccm. H_2O		Solid phase
Na_2CO_3	K_2CO_3	
119.8	Na_2CO_3
176.4	354.1	"
108.	500.1	"
61.19	0052.9	Na_2CO_3, K_2CO_3
.....	1084.0	K_2CO_3

(Kremann and Zitek, M. 1909, 30. 324.)

Solubility of $K_2CO_3 + KNO_3$ in H_2O at 25.2° .

1 l. of the solution contains	
Mol. K_2CO_3	Mol. KNO_3
0.00	3.217
0.59	2.62
1.35	1.97
2.10	1.46
2.70	1.14
3.58	0.79

(Touren, C. R. 1900, 131. 259.)

Solubility of $K_2CO_3 + KNO_3$ in H_2O at 10° .

In 1000 ccm. H_2O		Solid phase
KNO_3	K_2CO_3	
208.9	KNO_3
26.62	1076.0	KNO_3, K_2CO_3
.....	1084.0	K_2CO_3

(Kremann and Zitek, M. 1909, 30. 325.)

Solubility of $K_2CO_3 + KNO_3$ in H_2O at 24.2° .

In 1000 com. H_2O		Solid phase
KNO_3	K_2CO_3	
3.85	..	KNO_3
5.00	130.3	"
1.67	348.4	"
1.80	371.9	"
3.04	888.1	"
5.78	878.3	"
1.11	1112.2	KNO_3, K_2CO_3

Kremann and Zitek, M. 1909, 30, 316.)

Solubility of $K_2CO_3 + KCl$ in H_2O at 30° .

K_2CO_3	KCl	Solid phase
%	%	
27	0	$K_2CO_3 \cdot 1\frac{1}{2}H_2O$
22	1.03	$K_2CO_3 \cdot 1\frac{1}{2}H_2O + KCl$
66	1.07	KCl
..
64	26.22	KCl
..	28.01	"

Author gives intermediary data.
(de Waal, Dissert. 1910.)Solubility of $K_2CO_3 + KOH$ in H_2O at 30° .

KOH	% K_2CO_3	Solid phase
%	%	
75	0	$KOH \cdot 2H_2O$
14	2.05	$KOH \cdot 2H_2O + K_2CO_3 \cdot 1\frac{1}{2}H_2O$
77	2.50	$K_2CO_3 \cdot 1\frac{1}{2}H_2O$
..
..	53.27	$K_2CO_3 \cdot 1\frac{1}{2}H_2O$

Author gives intermediary data.
(de Waal, Dissert. 1910.)Sol. in liquid NH_3 . (Franklin, Am. Ch. 1898, 30, 828.)Sol. in 9 pts. alcohol of $17^\circ B$. Insol. in dilute alcohol.Not decomp. by 1 pt. H_2SO_4 + 6 pts. absolute alcohol. Not decomp. by 1 pt. HNO_3 + 4 pts. absolute alcohol. Not decomp. by an alcoholic solution of HCl , oxalic, racemic, tartaric, or glacial acetic acids, but is decomp. by alcoholic solution of citric acid.Solubility in methyl alcohol. Composition of liquids in equilibrium with solid K_2CO_3 at t° .

t°	Upper layer			Lower layer		
	% K_2CO_3	% CH_3OH	% H_2O	% K_2CO_3	% CH_3OH	% H_2O
20	21.7	42.2	36.1
20	13.8	52.1	34.1
20	12.4	44.2	8.2	47.6
0	7.6	66.3	26.1	46.3	6.7	47
0	7.4	46.6	6.6	46.8
17	6.2	69.6	24.2	48.3	5.7	46
15	5.0	72.9	22.1	..	4.3	44.7

Bruyn, Z. phys. Ch. 1900, 32, 63 and ff.)

Solubility in ethyl alcohol. Composition of liquids in equilibrium with solid K_2CO_3 at t° .

t°	Upper layer			Lower layer		
	% K_2CO_3	% C_2H_5OH	% H_2O	% K_2CO_3	% C_2H_5OH	% H_2O
-18	0.03	90.3	9.7	51.2	0.2	48.6
0	0.04	91.9	8.1	51.3	0.2	48.5
+17	0.06	91.5	8.4	52.1	0.2	47.7
35	0.07	90.9	9	53.4	0.2	46.4
50	0.09	91.8	8.1	55.3	0.2	44.5
75	0.12	91.4	8.5	57.9	2.0	40.9

(de Bruyn.)

Solubility in ethyl alcohol + Aq at 25° .When K_2CO_3 is dissolved in ethyl alcohol + Aq two layers are formed, the compositions of which are as follows:

Upper layer			Lower layer		
% alcohol	% H_2O	% K_2CO_3	% alcohol	% H_2O	% K_2CO_3
81.25	18.61	0.14	0.82	55.42	43.76
71.67	27.91	0.42	1.79	61.61	38.60
56.98	41.55	1.47	4.02	65.73	30.25
53.92	44.13	2.05	4.88	66.87	28.25
50.21	47.24	2.55	5.54	67.06	27.41
43.93	52.04	3.92	7.71	67.56	24.74
37.64	56.45	5.90	10.54	67.40	22.06
28.43	61.57	10.00	15.73	66.38	17.90

(Cuno, W. Ann. 1909, (4) 28, 664.)

Solubility of K_2CO_3 in alcohol + Aq at 30° .

K_2CO_3	% Alcohol	% H_2O	Solid phase
%	%	%	
53.27	0	46.73	$K_2CO_3 \cdot 1\frac{1}{2}H_2O$
53.09	0.1	46.81	"
..	"
..	"
0.13	90.49	9.38	$K_2CO_3 \cdot 1\frac{1}{2}H_2O$
..	"
..	"
0.04	99.92	0.04	$K_2CO_3 + K_2CO_3 \cdot 1\frac{1}{2}H_2O$

* Solution separates into two layers.
(de Waal, Dissert. Leiden, 1910.)A full discussion of the solubility of K_2CO_3 in methyl, ethyl, propyl, isopropyl, and allyl alcohols is given by Frankforter and Frary (J. phys. Ch. 1913, 17, 402), and Frankforter and Temple (J. Am. Chem. Soc. 1915, 37, 2697).

K_2CO_3 will "salt out" acetone from aqueous solution. The table shows the composition of the solutions at the points at which inhomogeneous solutions of K_2CO_3 , acetone and H_2O just become homogeneous at 20°. 100 g. of the solution contain:

K_2CO_3	H_2O	acetone	K_2CO_3	H_2O	acetone
18.84	73.22	7.94	2.43	55.36	42.21
13.32	71.38	15.30	22.29	72.81	4.90
11.83	70.34	17.83	17.86	73.12	9.02
10.13	69.03	20.84	15.81	72.53	11.66
8.24	67.31	24.45	14.39	71.89	13.72
7.22	65.99	26.79	10.29	69.46	20.25
6.04	64.39	29.57	1.91	54.05	44.04
28.87	69.08	2.05	1.76	52.86	45.38
23.94	71.98	4.08	1.60	51.60	46.80
21.52	72.75	5.33	1.29	49.57	49.14
19.60	73.10	7.70	1.08	47.86	51.06
6.46	65.34	28.20	0.94	46.73	52.33
5.91	64.65	29.44	0.75	44.72	54.53
5.60	63.93	30.47	0.66	43.31	56.03
5.04	62.80	32.16	0.60	42.49	56.91
4.50	61.48	34.02	0.54	41.73	57.73
3.80	59.79	36.41	0.50	40.69	58.81
3.18	57.95	38.87	0.46	40.48	60.06
2.73	56.50	40.77

(Frankforter and Cohen, J. Am. Chem. Soc. 1914, **36**, 1121.)

Insol. in benzonitrile. (Naumann, B. 1914, **47**, 1370.)

Insol. in methyl acetate (Naumann, B. 1909, **42**, 3790); ethyl acetate. (Naumann, B. 1910, **34**, 314.)

Sol. in phenol.

Sol. in 13.5 pts. glycerine of 1.225 sp. gr. (Vogel, N. Repert. **16**, 557.)

100 g. sat. K_2CO_3 + sugar + Aq. contains 22.44 g. K_2CO_3 and 56 g. sugar at 31.25° (Köhler, Z. Ver. Zuckerind. 1897, **47**, 447.) + H_2O .

+ $1\frac{1}{2}H_2O$. Very deliquescent. (Pohl.)

Deliquescent only in very moist air. (Stadeler.)

Sol. in H_2O with evolution of heat. (Pohl.)

Sol. at 17.6° with absorption of heat, at 32° with evolution of heat, and at 25° with neither absorption nor evolution of heat. (Berthelot, C. R. **78**, 1722.)

Sat. solution of $K_2CO_3 + 1\frac{1}{2}H_2O$ in H_2O contains 53.27 g. anhydrous K_2CO_3 in 100 g. solution at 30°. (de Waal, Dissert. 1910.)

+ $2H_2O$. Salt usually given as containing $1\frac{1}{2}H_2O$ contains $2H_2O$. (Gerlach, Z. anal. **26**, 460.)

Sat. solution of $K_2CO_3 + 2H_2O$ contains 112 g. anhydrous K_2CO_3 in 100 g. H_2O at 25°. (Osaka, J. Tok. Ch. Soc. 1911, **32**, 870.)

+ $4H_2O$. Not deliquescent in closed vessels. (Gerlach, *l. c.*)

Potassium hydrogen carbonate, $KHCO_3$.

Not deliquescent.

Sol. in 3.5 pts. H_2O at 15°. (Redwood.) Sol. in 4 pts. H_2O at moderate temperatures. (Bergmann.) Sol. in 0.8333 pt. boiling H_2O (Pelletier); in 4 pts. cold. and 1.2 pts. boiling H_2O (M. R. and P.'s Pharm.). Sol. in 4 pts. H_2O at 18.75°. (Abl.) 100 pts. H_2O at 15.5° dissolve 30 pts. and at 100°, 83 pts. (U're's Dict.) 100 pts. H_2O at 10-11.2° dissolve 26.1 pts. $KHCO_3$ and the sp. gr. of solution is 1.1536. (Anthon, Dimpl. **161**, 216.)

100 pts. H_2O dissolve at—

0°	10°	20°	30°
19.61	23.23	26.91	30.57
40°	50°	60°	70°
34.15	37.92	41.35	45.24

pts. $KHCO_3$. (Poggiale, A. ch. (3) **8**, 468.)

100 pts. H_2O dissolve pts. $KHCO_3$ at t°.

t°	Pts. $KHCO_3$	t°	Pts. $KHCO_3$
0	22.4	40	45.2
20	33.2	60	46.4

(Dibbits, J. pr. (2) **10**, 417.)

Sp. gr. of $KHCO_3$ + Aq at 15° containing 5% $KHCO_3$ = 1.0328; containing 10% $KHCO_3$ = 1.0674. (Kohlrausch, Z. anal. **28**, 472.)

Sol. in 1200 pts. boiling alcohol. (Berthelot.) Insol. in alcohol. (Dumas.)

100 pts. H_2O dissolve 19.3 pts. $KHCO_3$ and 8.3 pts. $NaHCO_3$ if the sat. solution of latter is sat. with former; and 26.1 pts. $KHCO_3$ and 6.0 pts. $NaHCO_3$ if the sat. solution of the former is sat. with the latter, all at 10°. (Mulder, J. B. **1866**, 67.)

Insol. in sat. K_2CO_3 + Aq. (Engel, C. R. **102**, 365.)

Solubility of $KHCO_3 + KNO_3$ in H_2O in an atmosphere of CO_2 .

1 litre of the solution contains			
at 11.5°		at 25.2°	
Mol. $KHCO_3$	Mol. KNO_3	Mol. $KHCO_3$	Mol. KNO_3
0.60	2.33	0.60	3.28
0.39	2.17	0.89	2.84
0.76	2.03	1.33	2.65
1.16	1.92	1.91	2.45
1.55	1.81

This case is complicated by the fact that KNO_3 is more sol. in H_2O sat. with CO_2 than in pure H_2O .

(Touren, C. R. 1906, **131**, 261.)

Insol. in benzonitrile (Naumann, B. 1914, **47**, 1370.)

Insol. in methyl acetate (Naumann, B. 1909, **42**, 3790); ethyl acetate (Naumann, B. 1910, **34**, 314.)

Potassium praseodymium carbonate, $K_2CO_3 \cdot Pr_2(CO_3)_3 + 12H_2O$.

Ppt. Sol. in 0% K_2CO_3 + Aq. (Meyer, Z. anorg. 1904, **41**, 104.)

Potassium samarium carbonate, K_2CO_3 , $Sm_2(CO_3)_3 + 12H_2O$.
(Cleve.)

Potassium silver carbonate, $KAgCO_3$.
Decomp. by H_2O . (de Schulten, C. R. 105. 811.)

Ppt. Decomp. by H_2O . (Reynolds, Chem. Soc. 1898, 73. 265.)

Potassium sodium carbonate, $KNaCO_3 + 6H_2O$.

Slightly efflorescent. Sol. in 0.75 pt. H_2O at 12.5° ; in 0.54 pt. H_2O at 15° .

Sat. solution at 15° has sp. gr. = 1.366. (Stolba, J. pr. 94. 406.)

Decomp. by recrystallizing from H_2O , but crystallizes undecomposed from sat. $K_2CO_3 + Aq$.

Sol. in H_2O . (Osaka, C. A. 1911. 2601.)
See $K_2CO_3 + Na_2CO_3$ under Na_2CO_3 .

+ $3H_2O$. (Kremann and Zite^k, M. 1909. 30. 317.) Does not exist. (Osaka.)

$K_2CO_3, 2Na_2CO_3 + 18H_2O$. Sl. efflorescent. Very sol. in H_2O . (Marignac.)

Potassium stannous carbonate, $K_2CO_3, 2SnCO_3 + 2H_2O$.

Decomp. by H_2O . (Deville.)

Potassium uranyl carbonate, $2K_2CO_3, (UO_2)_2CO_3$.

Sol. without decomp. in 13.5 pts. H_2O at 15° , and in somewhat less warm H_2O . Sol. in boiling H_2O with decomp.

More sol. in K_2CO_3 , or $KHCO_3 + Aq$ than in H_2O . (Rose.)

Insol. in alcohol. (Ebelmen, A. ch. (3) 5. 189.)

Potassium zinc carbonate, $4K_2O, 6ZnO, 11CO_2 + 8H_2O$.

Can be washed with cold H_2O without decomp. (Deville, A. ch. (3) 33. 99.)

Praseodymium carbonate, $Pr_2(CO_3)_3 + 8H_2O$.

Sol. in H_2O . (von Schule, Z. anorg. 1898, 18. 362.)

Praseodymium sodium carbonate, $2Pr_2(CO_3)_3, 3Na_2CO_3 + 22H_2O(?)$.

Ppt. Easily decomp. (Meyer, Z. anorg. 1904, 41. 105.)

Radium carbonate.

Less sol. in H_2O than corresponding Ba comp. (Curie, Dissert. 1903.)

Rubidium carbonate, Rb_2CO_3 .

Very deliquescent, and sol. in H_2O . 100 pts. absolute alcohol dissolve 0.74 pt. Rb_2CO_3 . (Bunsen.)

Rubidium hydrogen carbonate, $RbHCO_3$.

Not deliquescent. Easily sol. in H_2O . (Bunsen.)

Samarium carbonate, $Sm_2(CO_3)_3 + 3H_2O$.

Insol. in H_2O . (Cleve, Bull. Soc. (2) 43. 168.)

Samarium sodium carbonate, $Sm_2(CO_3)_3, Na_2CO_3 + 16H_2O$.

Ppt. (Cleve.)

Scandium carbonate, $Sc_2(CO_3)_3 + 12H_2O$.

(Crookes, Roy. Soc. Proc. 1908, 80, A. 518.)

Scandium sodium carbonate, $Sc_2(CO_3)_3, 4Na_2CO_3 + 6H_2O$.

Difficultly sol. in H_2O .

Sol. in cold, less sol. in hot alkali carbonates + Aq. (R. Meyer, Z. anorg. 1910, 67. 410.)

Silver carbonate, Ag_2CO_3 .

Somewhat sol. in H_2O . Sol. in 31,978 pts. H_2O at 15° . (Kremers, Pogg. 85. 248.) 1 g. Ag_2CO_3 dissolves in 2 l. boiling H_2O . (Joulin, A. ch (4) 30. 260.)

Solubility in H_2O at $25^\circ = 1.16 \times 10^{-4}$ mol./l. (Spencer and Le Pla, Z. anorg. 1910, 65. 14.)

1 l. H_2O at 25° dissolves 1.2×10^{-4} gram atoms of silver. (Abegg and Cox, Z. phys. Ch. 1903, 46. 11.)

Insol. in $H_2CO_3 + Aq$. (Bergman.) Sol. in 961 pts. $H_2CO_3 + Aq$. (Lassaigne.) 1 l. sat. $H_2CO_3 + Aq$ dissolves 0.846 g. Ag_2CO_3 at 15° . (Johnson, C. N. 54. 75.)

Sol. in $(NH_4)_2CO_3 + Aq$ or $NH_4OH + Aq$; sl. sol. in $K_2CO_3 + Aq$. (Wittstein.) Easily sol. in $Na_2S_2O_3 + Aq$. (Herschel, 1819.) Sol. in hot $NH_4Cl + Aq$, and sl. sol. in $NH_4NO_3 + Aq$. (Brett, 1837.) Not pptd. in presence of Na citrate. (Spiller.) Decomp. by $HCl + Aq$, and chlorides + Aq.

Somewhat sol. in conc. $NaNO_3 + Aq$. (de Coninck, Belg. Acad. Bull. 1909, 333.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in alcohol.

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Silver carbonate ammonia.

Easily sol. in H_2O . Sol. in $NH_4OH + Aq$, from which it is precipitated by absolute alcohol. (Berzelius.)

$Ag_2CO_3, 4NH_3$. Ppt. Insol. in alcohol. (Keen, C. N. 31. 231.)

Sodium carbonate, Na_2CO_3 .

Anhydrous. Sol. in H_2O with evolution of heat.

Sol. in 5.967 pts. H_2O at 15° . (Fresenius.) 100 pts. H_2O at 14.6° dissolve 7.74 pts. Na_2CO_3 , or 20.64 pts. $Na_2CO_3, 10H_2O$ is sol. in rather less than 1 pt. boiling H_2O . (Thomson, 1831.)

Sol. in 2 pts. H_2O . (Bergman.)

Sol. in 2 pts. H_2O at 18.75° . (Abl.)

Solubility in 100 pts. H ₂ O at t°.						Possesses four different degrees of solubility, according to different states of molecular constitution and degrees of hydration. (A. ch. (3) 44. 330.) Little more sol. at 34–38° than at 10°; maximum of solubility is probably at 34–38° (Löwel.)
t°	Pts. Na ₂ CO ₃	Pts. Na ₂ CO ₃ + 10H ₂ O	t°	Pts. Na ₂ CO ₃	Pts. Na ₂ CO ₃ + 10H ₂ O	
0	7.08	21.52	25	35.90	171.33	
10	16.66	61.98	30	35.90	241.57	
20	30.83	123.12	104.6	48.50	420.68	

(Poggiale, A. ch. (3) 8. 468.)

Solubility of Na₂CO₃, Na₂CO₃+10H₂O, Na₂CO₃+7H₂O (a), and Na₂CO₃+7H₂O (b) in 100 pts. H₂O at t°.

t°	Sat. solution of Na ₂ CO ₃ + 10H ₂ O contains—		Sat. solution of Na ₂ CO ₃ + 7H ₂ O (b) contains—			Sat. solution of Na ₂ CO ₃ + 7H ₂ O (a) contains—		
	Pts. Na ₂ CO ₃ in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ + 10H ₂ O in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ + 7H ₂ O (b) in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ + 10H ₂ O in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ + 7H ₂ O (a) in 100 pts. H ₂ O	Pts. Na ₂ CO ₃ + 7H ₂ O (b) in 100 pts. H ₂ O
0	6.97	21.33	20.39	58.93	84.28	31.93	112.94	184.93
10	12.06	40.94	26.33	83.94	128.57	37.85	150.77	284.93
15	16.20	63.20	29.58	100.00	160.51	41.55	179.90	384.93
20	21.71	92.82	38.55	122.25	210.58	45.79	220.20	554.93
25	28.50	149.13	38.07	152.36	290.91
30	37.24	273.64	43.45	196.93	447.93
38	51.67	1142.17
104	45.47	539.63

(Löwel, A. ch. (3) 33. 382.)

100 pts. H₂O at 14° dissolve 60.4 pts. Na₂CO₃+10H₂O; at 36°, 833 pts.; at 104°, 445 pts. Solubility increases to 36°, then diminishes. (Payen, A. ch. (3) 43. 233.)
There are apparently two maxima of solubility; the one occurring at 15°, or even lower, as warm solutions cool; the other at 34–38°, when cold solutions are warmed. (Payen, A. ch. (3) 44. 330.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. Na ₂ CO ₃	t°	Pts. Na ₂ CO ₃	t°	Pts. Na ₂ CO ₃
0	7.1	22	23.8	43	46.2
1	7.5	23	25.1	44	46.2
2	7.8	24	26.5	45	46.2
3	8.4	25	28.0	46	46.2
4	8.9	26	29.7	47	46.2
5	9.5	27	31.6	48	46.2
6	10.0	28	33.6	49	46.2
7	10.6	29	35.8	50	46.2
8	11.2	30	38.1	51	46.2
9	11.9	31	41.4	52	46.2
10	12.6	32	46.2	53	46.2
11	13.3	32.5	59.0	54	46.2
12	14.0	33	46.2	55	46.2
13	14.8	34	46.2	56	46.2
14	15.6	35	46.2	57	46.2
15	16.5	36	46.2	58	46.2
16	17.4	37	46.2	59	46.2
17	18.3	38	46.2	60	46.2
18	19.3	39	46.2	61	46.2
19	20.3	40	46.2	62	46.2
20	21.4	41	46.2	63	46.2
21	22.6	42	46.2	64	46.2

Solubility in 100 pts. H₂O at t°—Cont.

t°	Pts. Na ₂ CO ₃	t°	Pts. Na ₂ CO ₃	t°	Pts. Na ₂ CO ₃
65	46.2	79	46.2	93	46.2
66	46.2	80	46.1	94	46.2
67	46.2	81	46.1	95	46.2
68	46.2	82	46.1	96	46.2
69	46.2	83	46.0	97	46.2
70	46.2	84	46.0	98	46.2
71	46.2	85	45.9	99	46.2
72	46.2	86	45.9	100	46.2
73	46.2	87	45.8	101	46.2
74	46.2	88	45.8	102	46.2
75	46.2	89	45.8	103	46.2
76	46.2	90	45.7	104	46.2
77	46.2	91	45.7	105	46.2
78	46.2	92	45.7

(Mulder, Scheik. Verhandel. 1864. 1)

Liabile to form supersaturated solutions. Supersat. Na₂CO₃+Aq (2 pts. Na₂CO₃: 10H₂O: 1 pt. H₂O) may be kept in a closed vessel with cotton wool. (Schröder.)
When supersat. Na₂CO₃+Aq is exposed to low temperatures, the 10H₂O salt crystallizes out; but under other circumstances two salts are formed, each containing 7H₂O. The one is four times as sol. at 10° as the 10H₂O salt, and the other twice as sol. See above. (A. ch. (3) 33. 337.)
See also Na₂CO₃+H₂O, 7H₂O, and 10H₂O.

Sp. gr. of Na₂CO₃+Aq at 15°.

Na ₂ CO ₃	Sp. gr.	% Na ₂ CO ₃	Sp. gr.
0.372	1.0040	7.812	1.0892
0.744	1.0081	8.184	1.0937
1.116	1.0121	8.556	1.0982
1.488	1.0163	8.928	1.1028
1.850	1.0204	9.300	1.1074
2.232	1.0245	9.672	1.1120
2.504	1.0286	10.044	1.1167
2.976	1.0327	10.416	1.1214
3.348	1.0368	10.788	1.1261
3.720	1.0410	11.160	1.1308
4.090	1.0452	11.532	1.1356
4.464	1.0494	11.904	1.1404
4.836	1.0537	12.276	1.1452
5.208	1.0576	12.648	1.1500
5.580	1.0625	13.020	1.1549
5.972	1.0669	13.392	1.1598
6.324	1.0713	13.764	1.1648
6.396	1.0757	14.136	1.1698
6.768	1.0802	14.508	1.1748
7.440	1.0847	14.880	1.1816

(Tünnerman.)

Sp. gr. of Na₂CO₃+Aq at 15°.

%	Sp. gr. if % is Na ₂ CO ₃	Sp. gr. if % is Na ₂ CO ₃ +10H ₂ O
1	1.0105	1.004
2	1.0210	1.008
3	1.0315	1.012
4	1.0420	1.016
5	1.0525	1.020
6	1.0631	1.023
7	1.0737	1.027
8	1.0843	1.031
9	1.0950	1.035
10	1.1057	1.039
11	1.1165	1.043
12	1.1274	1.047
13	1.1384	1.050
14	1.1495	1.054
15	1.058
16	1.062
17	1.066
18	1.070
19	1.074
20	1.078
21	1.082
22	1.086
23	1.090
24	1.094
25	1.099
26	1.103
27	1.106
28	1.110
29	1.114
30	1.119
31	1.123
32	1.126
33	1.130
34	1.135
35	1.139
36	1.143
37	1.147
38	1.150

(Gerlach, Z. anal. 8. 279.)

Sp. gr. of Na₂CO₃+Aq at 17.5°.

% Na ₂ CO ₃	% Na ₂ CO ₃ +10H ₂ O	Sp. gr.	% Na ₂ CO ₃	% Na ₂ CO ₃ +10H ₂ O	Sp. gr.
1	2.70	1.010	9	24.30	1.095
2	5.40	1.020	10	27.00	1.105
3	8.10	1.031	11	29.70	1.116
4	10.18	1.041	12	32.40	1.127
5	13.50	1.052	13	35.10	1.137
6	16.20	1.063	14	37.80	1.148
7	18.90	1.073	15	40.50	1.157
8	21.60	1.084

(Hager.)

Sp. gr. of Na₂CO₃+Aq increases or diminishes by a change of temperature of 1° by the following amounts—

Corr.	% Na ₂ CO ₃
0.0004	13-15
0.00033	8-12
0.00026	3-7

(Hager, Comm. 1883.)

Sp. gr. of conc. Na₂CO₃+Aq at 30°.

Sp. gr.	% Na ₂ CO ₃	g. Na ₂ CO ₃ in 1 l.	Sp. gr.	% Na ₂ CO ₃	g. Na ₂ CO ₃ in 1 l.
1.310	28.13	368.5	1.220	20.47	249.7
1.300	27.30	354.9	1.210	19.61	237.3
1.290	26.46	341.3	1.200	18.76	225.1
1.280	25.62	327.9	1.190	17.90	214.0
1.270	24.78	314.7	1.180	17.04	201.1
1.260	23.93	301.5	1.170	16.18	189.3
1.250	23.08	288.5	1.160	15.32	177.7
1.240	22.21	275.4	1.150	14.47	166.4
1.230	21.33	262.3	1.140	13.62	155.3

(Lunge, Chem. Ind. 1882. 320.)

Sp. gr. of Na₂CO₃+Aq at 23°.

% Na ₂ CO ₃ +10H ₂ O	% Na ₂ CO ₃	Sp. gr.	% Na ₂ CO ₃ +10H ₂ O	% Na ₂ CO ₃	Sp. gr.
1	0.370	1.0038	16	5.929	1.0628
2	0.741	1.0076	17	6.299	1.0668
3	1.112	1.0114	18	6.670	1.0708
4	1.482	1.0153	19	7.041	1.0748
5	1.853	1.0192	20	7.412	1.0789
6	2.223	1.0231	21	7.782	1.0836
7	2.594	1.0271	22	8.153	1.0871
8	2.965	1.0309	23	8.523	1.0912
9	3.335	1.0348	24	8.894	1.0953
10	3.706	1.0388	25	9.264	1.0994
11	4.076	1.0428	26	9.635	1.1035
12	4.447	1.0468	27	10.005	1.1076
13	4.817	1.0508	28	10.376	1.1117
14	5.188	1.0548	29	10.746	1.1158
15	5.558	1.0588	30	11.118	1.1200

Sp. gr. of $\text{Na}_2\text{CO}_3 + \text{Aq}$ at 23° —Continued.

$\frac{\% \text{Na}_2\text{CO}_3}{+10\text{H}_2\text{O}}$	$\frac{\% \text{Na}_2\text{CO}_3}{+10\text{H}_2\text{O}}$	Sp. gr.	$\frac{\% \text{Na}_2\text{CO}_3}{+10\text{H}_2\text{O}}$	$\frac{\% \text{Na}_2\text{CO}_3}{+10\text{H}_2\text{O}}$	Sp. gr.
31	11.488	1.1242	41	15.195	1.1662
32	11.859	1.1284	42	15.556	1.1704
33	12.230	1.1326	43	15.936	1.1746
34	12.600	1.1368	44	16.307	1.1788
35	12.971	1.1410	45	16.677	1.1830
36	13.341	1.1452	46	17.048	1.1873
37	13.712	1.1494	47	17.418	1.1916
38	14.082	1.1536	48	17.789	1.1959
39	14.530	1.1578	49	18.159	1.2002
40	14.824	1.1620	50	18.530	1.2045

(Schiff, A. 113. 186.)

Sp. gr. of $\text{Na}_2\text{CO}_3 + \text{Aq}$ at 23.3° . a = number of grms. $\times \frac{1}{2}$ mol. wt., dissolved in 1000 grms. H_2O ; b = sp. gr. if $a = \text{Na}_2\text{CO}_3$, $10\text{H}_2\text{O}$ ($\frac{1}{2}$ mol. wt. = 143); c = sp. gr. if $a = \text{Na}_2\text{CO}_3$ ($\frac{1}{2}$ mol. wt. = 53).

a	b	c	a	b	c
1	1.048	1.052	5	1.163	1.226
2	1.086	1.100	6	1.182	...
3	1.117	1.145	7	1.198	...
4	1.142	1.187

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of $\text{Na}_2\text{CO}_3 + \text{Aq}$ at 18° .

$\frac{\% \text{Na}_2\text{CO}_3}{+10\text{H}_2\text{O}}$	Sp. gr.	$\frac{\% \text{Na}_2\text{CO}_3}{+10\text{H}_2\text{O}}$	Sp. gr.
5	1.0511	15	1.1590
10	1.1044

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of $\text{Na}_2\text{CO}_3 + \text{Aq}$.

g.-equivalents Na_2CO_3 per l.	t°	Sp. gr. t°/t°
0.002524	16.004	1.0001418
0.005041	16.026	1.0002844
0.01006	16.049	1.000568
0.02501	16.028	1.001413
0.04954	16.050	1.002789
0.10188	16.030	1.005699
0.24646	16.041	1.013598
0.002628	16.051	1.0001473
0.003948	16.088	1.0002216
0.009182	16.081	1.0005181
0.01830	16.089	1.001033
0.10842	16.042	1.006048
0.21570	16.055	1.011910
0.4297	15.14	1.02346
2.5015	16.05	1.12533

(Kohlrausch, W. Ann. 1894, 53. 26.)

Sp. gr. of $\text{Na}_2\text{CO}_3 + \text{Aq}$ at t° . H_2O at $4^\circ = 1$.

t°	$\% \text{Na}_2\text{CO}_3$	Sp. gr.
60°	28.74	1.2971
	25.20	1.2546
	22.25	1.2191
	18.23	1.1746
	14.06	1.1277
80°	28.59	1.2807
	18.26	1.1607

(Wegscheider, M. 1905, 26. 690.)

Sp. gr. of dil $\text{Na}_2\text{CO}_3 + \text{Aq}$ at 20.004° .
Conc. = g. equiv. Na_2CO_3 per l. at 20.004° and 730 mm.

Sp. gr. compared with H_2O at $20.004^\circ = 1$.

Conc.	Sp. gr.
0.0000	1.000,000,0
0.0001	1.000,005,6
0.0002	1.000,011,2
0.0004	1.000,022,5
0.0005	1.000,028,1
0.0010	1.000,056,3
0.0020	1.000,112,7
0.0040	1.000,225,8
0.0050	1.000,282,4
0.0100	1.000,564,8

(Lamb and Lee, J. Am. Chem. Soc. 1913, 35. 1685.)

$\text{Na}_2\text{CO}_3 + \text{Aq}$ containing 5% Na_2CO_3 boils at 100.5° ; 10% Na_2CO_3 at 101.1° ; 15% Na_2CO_3 at 101.8° . (Gerlach.)

Sat. solution boils at 104.4° (Griffiths, 1825); 106° (Kremers); 104° (Payen.)

Sat. solution forms a crust at 104.1° , and contains 42.2 pts. Na_2CO_3 to 100 pts. H_2O ; highest temperature observed, 105° . (Gerlach, Z. anal. 26. 427.)

B.-pt. of $\text{Na}_2\text{CO}_3 + \text{Aq}$ containing pts. Na_2CO_3 to 100 pts. H_2O . G = according to Gerlach (Z. anal. 26. 458); L = according to Legrand (A. ch. (2) 59. 426).

B.-pt.	G	L	B.-pt.	G	L
100.5°	5.2	7.5	103.5°	36.2	41.0
101.0	10.4	14.4	104.0	41.2	44.7
101.5	15.6	20.8	104.5	46.2	47.9
102.0	20.8	26.7	104.63	48.5
102.5	26.0	32.0	105.0	51.2
103.0	31.1	36.8

Less sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$ than in H_2O . (Fresenius.)

See also under Ammonia.

Solubility of $\text{Na}_2\text{CO}_3 + \text{NH}_4\text{Cl}$. See under Ammonium Chloride.

Solubility of $\text{Na}_2\text{CO}_3 + \text{K}_2\text{CO}_3$. See under Carbonate, potassium.

The reciprocal solubility of sodium carbonate and sodium hydrogen carbonate in H₂O has been determined. (de Paepe, C. A. 1911, 2603, and 1912, 2723.)

Solubility of Na₂CO₃+NaHCO₃ in H₂O at 25°.

g. per 100 g. H ₂ O		Solid phase
NaHCO ₃	Na ₂ CO ₃	
0	28.3	Na ₂ CO ₃ .10H ₂ O
2.1	27.3	"
4.2	26.5	Na ₂ CO ₃ .10H ₂ O+NaHCO ₃
5.7	19.2	NaHCO ₃
7.3	12.4	"
9.0	6.2	"
10.1	1.0	"

(de Paepe, Bull. Soc. Chim. Belg. 1911, 25. 174.)

Solubility of Na₂CO₃+NaHCO₃ in H₂O at 25°.

g. per l.		Solid phase
NaHCO ₃	Na ₂ CO ₃	
98.7	0.0	NaHCO ₃
50.8	216.6	NaHCO ₃ +Na ₂ CO ₃ .NaHCO ₃ .2H ₂ O
27.6	276.3	Na ₂ CO ₃ .NaHCO ₃ .2H ₂ O+Na ₂ CO ₃ .10H ₂ O
0.0	276.4	Na ₂ CO ₃ .10H ₂ O

(McCoy and Test, J. Am. Chem. Soc. 1911, 33. 474.)

Equilibrium between Na₂CO₃, NaHCO₃, and CO₂. See under Carbonate, sodium hydrogen.

Solubility of NaNO₃ in Na₂CO₃+Aq at 10°.

In 1000 ccm. H ₂ O		Solid phase
NaNO ₃	Na ₂ CO ₃	
805.0	...	NaNO ₃
704.8	87.5	NaNO ₃ , Na ₂ CO ₃
...	119.8	Na ₂ CO ₃

(Kremann, M. 1909, 30. 325.)

Solubility of NaNO₃ in Na₂CO₃+Aq at 24.2°.

In 1000 ccm. H ₂ O		Solid phase
NaNO ₃	Na ₂ CO ₃	
913.58	NaNO ₃
844.50	59.61	"
627.75	217.85	NaNO ₃ +Na ₂ CO ₃ .7H ₂ O
544.3	246.30
459.6	263.30	Na ₂ CO ₃ .7H ₂ O
....	28.55	Na ₂ CO ₃ .10H ₂ O
....	28.55	Na ₂ CO ₃ .10H ₂ O

(Kremann.)

Solubility of Na₂CO₃+NaBr in H₂O at 30°.

% Na ₂ CO ₃	% NaBr	Solid phase
27.98	0	Na ₂ CO ₃ .10H ₂ O
27.54	2.41	"
26.72	4.06	"
26.23	6.26	Na ₂ CO ₃ .10H ₂ O+Na ₂ CO ₃ .7H ₂ O
23.40	11.00	Na ₂ CO ₃ .7H ₂ O
22.68	12.22	"
19.86	16.88	"
19.57	16.95	Na ₂ CO ₃ .7H ₂ O+Na ₂ CO ₃ .H ₂ O
18.11	19.32	Na ₂ CO ₃ .H ₂ O
8.45	33.39	"
6.90	36.13	"
3.04	44.75	"
2.99	45.31	NaBr.2H ₂ O+Na ₂ CO ₃ .H ₂ O
2.60	45.68	NaBr.2H ₂ O
0	49.40	"

(Cocheret, Dissert. 1911.)

Solubility in NaCl+Aq. 100 pts. H₂O dissolve pts. NaCl and pts. Na₂CO₃+10H₂O, when that salt is in excess at 15°.

Pts NaCl	Pts. Na ₂ CO ₃ +10H ₂ O	Pts. NaCl	Pts. Na ₂ CO ₃ +10H ₂ O
0.00	61.42	23.70	39.06
4.03	53.86	27.93	39.73
8.02	48.00	31.65	41.44
12.02	43.78	35.46	43.77
16.05	40.96	sat.	
19.82	39.46	37.27	45.32

Solubility of anhydrous Na₂CO₃ in 100 pts. NaCl+Aq containing % NaCl at 15°.

% NaCl	Pts. Na ₂ CO ₃	% NaCl	Pts. Na ₂ CO ₃
0	16.408	12	10.488
1	15.717	13	10.244
2	15.060	14	10.041
3	14.438	15	9.880
4	13.851	16	9.762
5	13.299	17	9.686
6	12.783	18	9.655
7	12.305	19	9.667
8	11.864	20	9.725
9	11.461	21	9.828
10	11.097	22	9.997
11	10.773

(Reich, W. A. B. 99, 2b. 433.)

Solubility of Na₂CO₃+NaCl in H₂O at 30°.

% Na ₂ CO ₃	% NaCl	Solid phase
27.98	0	Na ₂ CO ₃ .10H ₂ O
27.48	0.9	"
27.12	3.33	"
26.82	4.15	Na ₂ CO ₃ .10H ₂ O+Na ₂ CO ₃ .7H ₂ O
25.59	5.17	Na ₂ CO ₃ .7H ₂ O
24.26	5.93	"

Solubility of Na ₂ CO ₃ +NaCl in H ₂ O at 30°—Continued.		
% Na ₂ CO ₃	% NaCl	Solid phase
22.75	10.24	Na ₂ CO ₃ ·7H ₂ O + Na ₂ CO ₃ ·H ₂ O
20.72	11.49	
18.00	14.12	NaCl + Na ₂ CO ₃ ·H ₂ O
14.81	16.26	NaCl
9.71	18.76	"
5.05	21.94	"
0	26.47	"

(Cocheret, Dissert. 1911.)

Solubility of Na ₂ CO ₃ +NaI in H ₂ O at 30°.		
% Na ₂ CO ₃	% NaI	Solid phase
27.4	0	Na ₂ CO ₃ ·10H ₂ O
26.5	2.4	"
25.5	4.7	"
25.2	5.2	"
24.4	8.6	"
24.3	9.5	Na ₂ CO ₃ ·10H ₂ O + Na ₂ CO ₃ ·7H ₂ O
23.0	11.2	Na ₂ CO ₃ ·7H ₂ O
20.8	14.0	"
20.0	15.7	"
18.7	18.4	"
15.3	25.4	Na ₂ CO ₃ ·7H ₂ O + Na ₂ CO ₃ ·H ₂ O
13.1	29.1	Na ₂ CO ₃ ·H ₂ O
10.4	33.3	"
6.4	40.4	"
4.2	46.0	"
3.1	49.5	"
2.7	51.0	"
1.5	54.6	"
0.9	57.6	"
0.6	61.2	"
0.3	65.6	NaI·2H ₂ O + Na ₂ CO ₃ ·H ₂ O
0.0	65.5	NaI·2H ₂ O

(Cocheret, Dissert. 1911.)

Insol. in liquid CO₂. (Büchner, Z. phys. Ch. 1906, 54. 674.)
Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)
Insol. in alcohol. (Fresenius.)
Sl. sol. in absolute alcohol; apparently insol. in an alcoholic solution of soap. (Duffy, Chem. Soc. 5. 305.)

Solubility of Na ₂ CO ₃ in ethyl alcohol at 20°.	
Alcohol, wt. per cent	g. Na ₂ CO ₃ per 100 g. solution
44	1.7
46	1.13
48	0.9
50	0.84
54	0.80

(Linebarger, A. Ch. J. 1892, 14. 380.)

Solubility of Na ₂ CO ₃ in alcohol+H ₂ O at		
% Na ₂ CO ₃	% alcohol	Solid phase
27.4	0	Na ₂ CO ₃ ·10H ₂ O
26.61	2.64	"
{ 26.14*	3.41 }	"
{ 1.38	44.81 }	"
0.62	52.99	"
0.61	53.26	"
0.53	55.70	Na ₂ CO ₃ ·10H ₂ O + Na ₂ CO ₃ ·7H ₂ O
0.51	56.56	Na ₂ CO ₃ ·7H ₂ O
0.47	62.61	"
0.40	63.20	"
0.15	72.80	"
0.11	73.06	Na ₂ CO ₃ ·7H ₂ O + Na ₂ CO ₃ ·H ₂ O
0.07	78.19	Na ₂ CO ₃ ·H ₂ O
0.07	82.26	"
0.06	86.76	"
0.06	90.95	"
0.04	93.09	"
0.03	95.06	Na ₂ CO ₃ ·H ₂ O + Na ₂ CO ₃
...	95.65	Na ₂ CO ₃
...	98.46	"

* Conjugated liquid phases.

(Cocheret, Dissert. 1911.)

See also under Na₂CO₃+H₂O, +7H₂O +10H₂O.

Not decomp. by 1 pt. H₂SO₄+6 pts. solute alcohol.

Not decomp. by alcoholic solutions of citric, tartaric, or glacial acetic acids; sl. decomp. by HNO₃+absolute alcohol.

Solubility of Na₂CO₃+NaBr, NaCl NaI in alcohol. Numerical data given by Cocheret (Dissert. 1911), reported in Transactions annuelles internationales des Constantes, for 1911.

Solubility of Na ₂ CO ₃ in propyl alcohol at	
Alcohol, wt. per cent	g. Na ₂ CO ₃ per 100 g. solution
28	4.4
38	2.7
44	1.7
46	1.5
48	1.3
50	1.2
54	0.9
62	0.4

(Linebarger, A. Ch. J. 1892, 14. 380.)

A full discussion of the solubility of Na₂CO₃ in propyl, and allyl alcohol is given by Forster and Temple (J. Am. Ch. Soc. 1915 2697).

Insol. in CS₂. (Arctowski, Z. anorg. 1 6. 257.)

Insol. in benzonitrile. (Naumann, B. 1 47. 1370.)

Insol. in methyl acetate (Naumann 1909, 42. 3790); ethyl acetate. (Naumann 1904, 37. 3602.)

solubility in mixtures of pyridine and H₂O —65° to +200°. Solubility curves are given. (Limbosch, Chem. Soc. 1909, 96 (2), 1014.)
sol. in acetone and in methylal. (Eidson, C. C. 1899, II. 1014.)
10 g. glycerine (sp. gr.=1.262) dissolve 1 g. Na₂CO₃ at 15-16°. (Ossendowski, m. J. 1907, 79. 575.)
10 g. sat. solution in glycol contain 3.28 g. Na₂CO₃. (de Coninck, Bull. Soc. Belg. Chem., 21. 141.)
10 g. sat. Na₂CO₃+sugar+Aq contain 64.73 g. sugar at 31.25°. (Zuckerind. 1897, 47. 447.)
H₂O. Takes up H₂O from the air. Less soluble in H₂O at 104° than at 38°; at 15-20°, 100 g. H₂O dissolve 52.4 pts. of this salt, calculated as Na₂CO₃. Insol. in alcohol. (See also under Na₂CO₃.)

Solubility in 100 pts. H₂O at t°.

Corrected t°. (Hydrogen scale)	Pts. anhydrous salt
29.86	50.53
29.89	50.75
31.80	50.31
35.17	49.63
35.37	49.67
35.66	49.37
35.86	49.44
36.45	49.36
36.90	49.29
37.91	49.11
38.92	49.09
40.94	48.51
40.93	48.52
43.94	47.98

(Wells and McAdam, J. Am. Chem. Soc. 1907, 29. 726.)

Solubility in alcohol+Aq.
Composition of the alcohol and water layers in contact with the solid phase Na₂CO₃+H₂O.

Alcohol layer			Water layer		
% alcohol	% salt	% water	% alcohol	% salt	% water
55.8	0.9	43.3	2.3	28.8	68.9
61.0	0.4	38.6	1.2	31.5	67.3
61.0	0.4	38.6	1.2	31.9	66.9
62.0	0.3	37.7	1.1	32.1	66.8
62.9	0.3	36.8	1.0	32.4	66.6

(Ketner, Z. phys. Ch. 1902, 39. 651.)

3H₂O. (Schickendantz, A. 155. 359.)
5H₂O. (Persoz, Pogg. 32. 303.)
Not efflorescent. Sol. in H₂O.
6H₂O. (Mitscherlich, Pogg. 8. 441.)
7H₂O. Efflorescent. Two salts, 7H₂O + 8H₂O of Thomson), and 7H₂O.
See also under Na₂CO₃.

Solubility in 100 pts. H₂O at t°.

Corrected t°. (Hydrogen scale)	Pts. anhydrous salt
30.35	43.50
31.82	45.16
32.86	46.28
34.37	48.22
34.76	48.98
35.15	49.23
35.17	49.34
35.62	50.08

(Wells and McAdam, J. Am. Chem. Soc. 1907, 29. 726.)

Composition of the solutions which can be in equilibrium with Na₂CO₃+7H₂Oβ at different temperatures.

t°	% Na ₂ CO ₃
32.1	31.8
32.5	32.1
33.3	32.7
33.9	33.0
34.5	33.9

(Ketner, Z. phys. Ch. 1902, 39. 646.)

Composition of the alcohol and water layers in contact with the solid phase, Na₂CO₃+7H₂Oβ, at different temperatures.

t°	Alcohol layer			Water layer		
	% alcohol	% salt	% water	% alcohol	% salt	% water
33.2	58.1	0.5	42.4	1.4	31.0	67.6
32.3	56.1	0.6	43.3	1.5	30.2	68.3
31.9	54.8	0.7	44.5	1.7	29.8	68.5
31.45	53.5	0.7	45.8	...	29.3
31.2	52.4	0.8	46.8	...	29.3

(Ketner.)

Composition of the two liquid layers which at different temperatures can be in metastable equilibrium with Na₂CO₃+7H₂Oβ.

t°	Alcohol layer			Water layer		
	% alcohol	% salt	% water	% alcohol	% salt	% water
28.9	46.9	1.3	51.8	2.3	26.3	71.4
26.6	39.1	1.3	59.6	3.3	25.4	71.3
23.0	24.5	6.7	68.8	7.0	20.2	72.8

(Ketner.)

+10H₂O. Efflorescent. Sol. in 1.05 pts. H₂O at 23°, and sat. solution has sp. gr. 1.1995. (Schiff, A. 109. 326.)
Melts in crystal H₂O at 34°. (Tilden, Chem. Soc. 45. 409.)
See above under Na₂CO₃ for further data.

Solubility in 100 pts. H ₂ O at t°.	
Corrected t°. (Hydrogen scale)	Pts. anhydrous salt
27.84	34.20
29.33	37.40
29.85	38.89
30.35	40.12
31.45	43.25
31.66	43.95
31.72	44.21
32.06	45.64

(Wells and McAdam, J. Am. Chem. Soc. 1907, 29. 726.)

Sat. solution at 25° contains 29.37 g. anhyd. Na₂CO₃ in 100 g. H₂O. (Osaka, J. Tok. Ch. Soc. 1911, 32. 870.)

Sat. solution at 25° contains 28.3 g. anhyd. Na₂CO₃ in 100 g. H₂O. (de Paepe, Bull. Soc. Chim. Belg. 1911, 25. 174.)

Sat. solution at 30° contains 27.4–27.98 g. anhyd. Na₂CO₃ in 100 g. of the solution. (Cocheret, Dissert. 1911.)

Sat. solution at 25° contains 27.64 g. anhyd. Na₂CO₃ in 100 cc. of the solution. (McCoy and Test, J. Am. Chem. Soc. 1911, 33. 474.)

Solubility in alcohol.

Composition of the alcohol and water layers in contact with the solid phase, Na₂CO₃+10H₂O, at different temperatures.

t°	Alcohol layer			Water layer		
	% alcohol	% salt	% water	% alcohol	% salt	% water
30.6	47.8	1.2	51.0	2.3	27.8	69.9
29.7	40.0	2.1	57.9	2.9	25.5	71.6
29.0	32.7	3.8	63.5	4.3	22.7	73.0
28.2	23.5	7.3	69.2	7.9	18.6	73.5

(Ketner, Z. phys. Ch. 1902, 39. 651.)

Solubility in alcohol + Aq.

Liquids which can be in equilibrium with Na₂CO₃+10H₂O at 21°.

% alcohol	% salt	% water
0	18.5	81.5
6.2	12.7	81.1
15.3	6.9	77.8
26.1	3.2	70.7
39.2	1.2	59.6
58.2	0.2	41.6
67.1	0.1	32.8
73.3	0.06	26.64

(Ketner.)

Composition of the alcohol liquids which be in equilibrium with Na₂CO₃+10H₂O and Na₂CO₃+7H₂O at different temperatures.

t°	% alcohol	% salt	% water
29°	62.3	0.3	37.4
26°	67.8	0.1	32.1
21°	73.3	0.06	26.6

(Ketner.)

See also under Na₂CO₃+15H₂O. (Jacquelin, A. 80. 241.)

Sodium hydrogen carbonate, NaHCO₃.

100 pts. cold H₂O dissolve 7.7 pts. NaHCO₃. (Schw. J. 6. 52.)

100 pts. H₂O at 11.25° dissolve 8.27 pts. NaHCO₃ form solution of 1.0613 sp. gr. (Anthon, Dingl. 216.)

100 pts. H₂O dissolve at—

0°	10°	20°	30°
8.95	10.04	11.15	12.24 pts. NaHCO ₃
40°	50°	60°	70°
13.35	14.45	15.57	16.69 pts. NaHCO ₃

(Poggiale, A. ch. (3) 8. 468.)

100 pts. H₂O dissolve pts. NaHCO₃ at

t°	Pts. NaHCO ₃	t°	Pts. NaHCO ₃	t°	Pts. NaHCO ₃
0	6.90	21	9.75	42	13
1	7.00	22	9.90	43	13
2	7.10	23	10.05	44	13
3	7.20	24	10.20	45	13
4	7.35	25	10.35	46	13
5	7.45	26	10.50	47	13
6	7.60	27	10.65	48	14
7	7.70	28	10.80	49	14
8	7.85	29	10.95	50	14
9	8.00	30	11.10	51	14
10	8.15	31	11.25	52	14
11	8.25	32	11.40	53	15
12	8.40	33	11.55	54	15
13	8.55	34	11.70	55	15
14	8.70	35	11.90	56	15
15	8.85	36	12.05	57	15
16	9.00	37	12.20	58	16
17	9.15	38	12.35	59	16
18	9.30	39	12.50	60	16
19	9.40	40	12.70	..	.
20	9.60	41	12.90	..	.

(Dibbits, J. pr. (2) 10. 417.)

Experiments with solutions of sodium hydrogen carbonate show that they gradually decompose after a time. (Treadwell, Z. anorg. Chem. 1898, 17. 204.)

The source of error of many solubility terminations of this substance is due to loss of CO₂. Solutions exposed to the air lose CO₂. (McCoy, Am. Ch. J. 1903, 29. 438.)

1 l. sat. solution at 25° contains 96 g. NaHCO₃. (McCoy and Test, J. Am. Chem. Soc. 1911, 33. 474.)

$\text{HCO}_3 + \text{Aq}$ sat. at 16° has sp. gr. = 1.4. (Stolba.)
Slightly insol. in sat. NaCl , or $\text{Na}_2\text{SO}_4 + \text{Aq}$.
Sol. in Ain , B. 5. 121.)
Equilibrium between Na_2CO_3 and NaHCO_3 ,
in H_2O and in contact with the air.
System: Na_2CO_3 , NaHCO_3 , and CO_2 .
Temperature, 25°C .

Amount solution used for titration cc.	Amount Na combined as Na_2CO_3 Per cent	Amount Na combined as NaHCO_3 Per cent
50	8.7	91.3
50		
20	20.0	80.0
20		
10	37.3	62.7
10		
10	59.3	40.7
10		
2	64.0	36.0
2		
3		

Temperature, 37°C .

Amount solution used for titration cc.	Amount Na combined as Na_2CO_3 Per cent	Amount Na combined as NaHCO_3 Per cent
50	10.5	89.5
50		
20	21.1	78.9
20		
10	41.3	58.7
10		
10	64.5	35.5
10		
2	81.9	18.1
2		
2	86.5	13.5
2		
2+	83.4	16.6
2		

Temperature, 50°C .

Amount solution used for titration cc.	Amount Na combined as Na_2CO_3 Per cent	Amount Na combined as NaHCO_3 Per cent
50	22.2	77.8
25		
20	32.9	67.1
20		
10	50.7	49.3
20		
10	70.0	30.0
10		
10	81.0	19.0
10		
2	86.8	13.2
2		
2		
2.1	87.1	12.9
2		

Temperature, 75°C .

Gram atoms Na per liter	Amount solution used for titration cc.	Amount Na combined as Na_2CO_3 Per cent	Amount Na combined as NaHCO_3 Per cent
0.003	50	25.7	74.3
	25		
	25		
0.019	20	34.8	65.2
	20		
0.036	10	55.7	44.3
	10		
0.270	5	79.5	20.5
	5		
0.702	1	85.0	15.0
	1		
6.56	1	84.8	15.2
	1		

System: Na_2CO_3 and NaHCO_3 at 25°C .

Total salts dissolved grams	Na_2CO_3		NaHCO_3	
	Weight Grams	Per cent	Weight Grams	Per cent
0.3555	0.0203	5.71	0.3352	94.29
1.1053	0.1505	13.62	0.9548	86.38
4.0443	1.1041	27.30	2.9402	72.70
14.6558	7.0212	47.91	7.6346	52.09
56.3982	29.8223	52.88	26.5759	47.12

(Cameron and Briggs, J. phys. Chem. 1901, 5. 540.)

100 g. acohol of 0.941 sp. gr. dissolve 1.2 g. NaHCO_3 at 15.5° .

100 g. glycerol dissolve 8 g. NaHCO_3 at 15.5° . (Ossendowski, Pharm. J. 1907, 79. 575.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Sodium dihydrogen tricarbonat,
 $\text{Na}_2\text{H}_2(\text{CO}_3)_3 + 3\text{H}_2\text{O}$.

More sol. than NaHCO_3 , less sol. than Na_2CO_3 in H_2O . (Rose, Pogg. 34. 160.)

100 pts. H_2O dissolve, calculated as $2\text{Na}_2\text{O}$, 3CO_2 —

at 0°	12.63 pts.	at 60°	29.68 pts.
" 10°	15.50 "	" 70°	32.55 "
" 20°	18.30 "	" 80°	35.8 "
" 30°	21.15 "	" 90°	38.63 "
" 40°	23.95 "	" 100°	41.59 "
" 50°	26.78 "		

(Poggiale, A. ch. (3) 8. 468.)

Min. Trona, Urao. See $\text{Na}_2\text{H}(\text{CO}_3)_2 + 2\text{H}_2\text{O}$.

Trisodium hydrogen carbonate, $\text{Na}_3\text{H}(\text{CO}_3)_2 + 2\text{H}_2\text{O}$.

Sol. in H_2O .

True formula of "Trona" and "Urao." (Zepharovich, Zeit. Kryst. 13. 135; de Mondesir, C. R. 104. 1505.)

Sodium thorium carbonate, $3\text{Na}_2\text{CO}_3, \text{Th}(\text{CO}_3)_2 + 12\text{H}_2\text{O}$.

Decomp. by H_2O . (Cleve.)

Sodium uranyl carbonate, $2\text{Na}_2\text{CO}_3, (\text{UO}_2)\text{CO}_3$.

Slowly sol. in H_2O . Solution sat. at 15° has sp. gr. = 1.161. (Anthon, Dingl. 156. 207.)

Sodium yttrium carbonate, $\text{Na}_2\text{CO}_3, \text{Y}_2(\text{CO}_3)_3 + 4\text{H}_2\text{O}$.

Ppt. Not decomp. by cold H_2O . (Cleve.)

Sodium zinc carbonate, $3\text{Na}_2\text{O}, 8\text{ZnO}, 11\text{CO}_2 + 8\text{H}_2\text{O} = 3\text{Na}_2\text{CO}_3, 8\text{ZnCO}_3 + 8\text{H}_2\text{O}$.

Sl. decomp. by pure H_2O . (Wöhler.)

Less easily decomp. by H_2O than most double carbonates. (Deville, A. ch. (3) 33. 101.)

$\text{Na}_2\text{O}, 3\text{ZnO}, 4\text{CO}_2 + 3\text{H}_2\text{O}$. (Kraut, Z. anorg. 1897, 13. 13.)

Sodium carbonate sulphite, $\text{Na}_2\text{CO}_3, 2\text{Na}_2\text{SO}_3 + 21\text{H}_2\text{O}$.

Sol. in hot H_2O , sl. sol. in cold H_2O . (Johnson, J. Soc. Chem. Ind. 1895, 14. 271.)

Strontium carbonate, SrCO_3 .

Sol. in 18,045 pts. H_2O at ordinary temp. (Fresenius.)

Sol. in 12,522 pts. H_2O at 15° . (Kremers, Pogg. 85. 247.)

Sol. in 33,000 pts. H_2O . (Bineau, C. R. 41. 511.)

Less sol. in H_2O than SrSO_4 . (Dulong.)

Sol. in 1536 pts. boiling H_2O . (Hepe, Edinb. Trans. 4. 5.)

Calculated from electrical conductivity of $\text{SrCO}_3 + \text{Aq}$, SrCO_3 is sol. in 121,760 pts. H_2O at 8.8° and 91,468 pts. at 24.3° (Holleman, Z. phys. Ch. 12. 130).

1 l. H_2O dissolves 11 mg. SrCO_3 at 18° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

"Solubility product" = 15.67×10^{-10} mol. litre. (McCoy and Smith, J. Am. Chem. Soc. 1911, 33. 473.)

Sol. in 833 pts. $\text{H}_2\text{CO}_3 + \text{Aq}$ at 10° . (Gmelin.)

Sol. in 56,545 pts. H_2O containing NH_4OH and $(\text{NH}_4)_2\text{CO}_3$.

Quite sol. in $\text{NH}_4\text{Cl} + \text{Aq}$ or $\text{NH}_4\text{NO}_3 + \text{Aq}$, but reprecipitated on addition of NH_4OH and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Fresenius.)

Partially decomp. by boiling with aqueous solutions of K_2SO_4 , Na_2SO_4 , CaSO_4 , $(\text{NH}_4)_2\text{SO}_4$, MgSO_4 , Na_2HPO_4 , $(\text{NH}_4)_2\text{HPO}_4$, K_2SO_3 , Na_2SO_3 , $(\text{NH}_4)_2\text{SO}_3$, $\text{Na}_2\text{B}_4\text{O}_7$,

Na_2AsO_3 , K_2AsO_3 , $\text{K}_2\text{C}_2\text{O}_4$, $\text{Na}_2\text{C}_2\text{O}_4$, NaF , and K_2CrO_4 . Decomp. is complete with the NH_4 salts. (Dulong, A. ch. 82. 286.)

Sl. decomp. by Na_2SO_4 , or $\text{K}_2\text{SO}_4 + \text{Aq}$. (Persoz.)

Easily sol. in NH_4 chloride, nitrate, or succinate + Aq , but less so than BaCO_3 . (Fresenius.) Sol. in ferric salts + Aq , with pptn. of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$. Sol. in Na citrate + Aq . (Spiller.) Not decomp. by a mixture of 1 pt. H_2SO_4 and 6 pts. absolute alcohol, or by alcoholic solutions of tartaric, racemic, citric, or glacial acetic acids; immediately decomp. by $\text{HNO}_3 + \text{absolute alcohol}$, or $\text{H}_2\text{C}_2\text{O}_4 + \text{abs. alcohol}$.

Solubility of SrCO_3 in $\text{NH}_4\text{Cl} + \text{Aq}$.

% NH_4Cl	% SrCO_3
5.35	0.179
10	0.259
20	0.358

(Cantoni and Goguelia, Bull. Soc. 1905, (3) 33. 13.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3602.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Insol. in acetone and in methylal. (Edmann, C. C. 1899, II. 1014.)

Min. *Strontianite*.

Strontium hydrogen carbonate.

SrCO_3 is sol. in 850 pts. of a sat. solution of CO_2 in H_2O .

Strontium uranyl carbonate, $\text{SrO}, 2\text{UO}_2, 2\text{CO}_2 + 16\text{H}_2\text{O}$.

As Ba comp. (Blinkoff, Dissert. 1900.)

Terbium carbonate.

Ppt. Insol. in excess $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Potratz, C. N. 1905, 92. 3.)

Thallous carbonate, Tl_2CO_3 .

100 pts. H_2O dissolve pts. Tl_2CO_3 (C=according to Crookes; L=according to Lamy) at—

15.5°	18°	62°	100°	100.8°
4.2	5.23	12.85	27.2	22.4 pts. Tl_2CO_3
C	L	L	C	L

Insol. in absolute alcohol (L), and ether (C).

Insol. in acetone and pyridine. (Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Thallous carbonate, acid, $\text{Tl}_2\text{O}, 2\text{CO}_2$.

Rather easily sol. in cold H_2O . (Cartan-jen.)

TiHCO₃. (Giorgio, *Gazz. ch. it.* 1894, 24, 74-479.)

Thallous carbonate platincyanide, $\text{Ti}_2\text{CO}_3 \cdot \text{Ti}_2\text{Pt}(\text{CN})_6$.

Sol. in hot, insol. in cold H_2O . (Friswell, *Chem. Soc.* (2) 9, 461.)

Thorium carbonate, basic, $2\text{ThO}_2 \cdot \text{CO}_2 + 3\text{H}_2\text{O}$.

Insol. in $\text{CO}_2 + \text{Aq}$, but sol. in excess of alkali carbonates + Aq , if conc.

Tin (stannous) carbonate, $2\text{SnO} \cdot \text{CO}_2$.

Easily decomp. on air; insol. in H_2O or $\text{H}_2\text{CO}_3 + \text{Aq}$. (Deville, *A. ch.* (3) 35, 448.)

Uranyl carbonate, basic, $5(\text{UO}_2)(\text{OH})_2 \cdot 3(\text{UO}_2)\text{CO}_2 + 6\text{H}_2\text{O}$.

Ppt. (Seubert and Elten, *Z. anorg.* 1893, 480.)

Ytterbium carbonate, basic, $\text{Yb}(\text{OH})\text{CO}_2 + \text{H}_2\text{O}$.

Ppt. (Cleve, *Z. anorg.* 1902, 32, 146.)

Ytterbium carbonate, $\text{Yb}_2(\text{CO}_3)_3 + 4\text{H}_2\text{O}$.

Ppt. (Cleve, *Z. anorg.* 1902, 32, 146.)

Yttrium carbonate, $\text{Y}_2(\text{CO}_3)_3 + 3\text{H}_2\text{O}$.

Insol. in H_2O ; very al. sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$. Sol. in $\text{SO}_2 + \text{Aq}$ and all mineral acids. Sol. in NH_4 salts, and alkali carbonates + Aq to some extent. More sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ than in $\text{K}_2\text{CO}_3 + \text{Aq}$. (Berlin.) More sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ than cerium, but 5 or 6 times less sol. than glucinum carbonate. (Vauquelin.) Sol. in large excess of $\text{KHCO}_3 + \text{Aq}$. (Rose.) Slowly sol. in NH_4 salts + Aq . (Berzelius.)

Zinc carbonates, basic, $8\text{ZnO} \cdot \text{CO}_2 + 2\text{H}_2\text{O}$; $5\text{ZnO} \cdot 2\text{CO}_2 + 3$, or $7\text{H}_2\text{O}$; $3\text{ZnO} \cdot \text{CO}_2 + \text{H}_2\text{O}$; $11\text{ZnO} \cdot 4\text{CO}_2 + 14\text{H}_2\text{O}$; $14\text{ZnO} \cdot 5\text{CO}_2 + 9\text{H}_2\text{O}$; $2\text{ZnO} \cdot \text{CO}_2 + \text{H}_2\text{O}$; $8\text{ZnO} \cdot 3\text{CO}_2 + 5\text{H}_2\text{O}$, etc.

All ppts. formed from Zn salts and carbonates + Aq . Sol. in 2000-3000 pts. cold H_2O , separates out on heating and does not redissolve on cooling. (Schindler.) Sol. in 20,895 pts. H_2O at 15° . (Kremers, *Pogg.* 85, 248.) Sol. in 44,600 pts. H_2O at ord. temp. (Fresenius.)

Sol. in 1428 pts. sat. $\text{H}_2\text{CO}_3 + \text{Aq}$. (Lassaigne.) Sol. in 189 pts. $\text{H}_2\text{CO}_3 + \text{Aq}$ sat. at 4-6 atmos. (Wagner, *Z. anal.* 6, 107.) Easily sol. in KOH , NaOH , NH_4OH , $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$, and in acids. Somewhat sol. in alkali bicarbonates and NH_4 salts + Aq . (Fresenius.) Sol. in hot (Fuchs), also cold (Brett, 1857) $\text{NH}_4\text{Cl} + \text{Aq}$; less sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett.)

Sol. in all NH_4 salts + Aq excepting $(\text{NH}_4)_2\text{S} + \text{Aq}$. (Terreil, *Bull. Soc.* (2) 9, 441.)

Insol. in Na_2CO_3 , or $\text{K}_2\text{CO}_3 + \text{Aq}$. Sol. in ferric salts + Aq with pptn. of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$. (Fuchs, 1831.)

The carbonates described by Boussingault, Wackenroder, Rose, and probably all salts between ZnO , CO_2 , and $5\text{ZnO} \cdot 2\text{CO}_2$ are mixtures. (Kraut, *Z. anorg.* 1897, 13, 1-15.)

$3\text{ZnO} \cdot \text{CO}_2 + 2\text{H}_2\text{O}$. Min. *Zinc bloom*, *Hydrozincite*.

$\text{ZnCO}_3 \cdot 3\text{ZnO} \cdot \text{H}_2\text{O}$. Min. *Auricalcite*.

Zinc carbonate, ZnCO_3 .

1 l. H_2O at 15° dissolves 0.01 g.; 1 l. H_2O dissolves 1.64×10^{-4} mols., or 0.206 g. ZnCO_3 at 25° . (Ageno and Valla, *Att. acc. Linc.* 1911, 20, II, 706.)

1 l. 5.85% $\text{NaCl} + \text{Aq}$ dissolves 0.0586 g.; 1 l. 7.45% $\text{KCl} + \text{Aq}$ dissolves 0.0477 g. ZnCO_3 . (Essen, *Gm.-K.* 4, 1, 680.)

Sol. in acids, $\text{KOH} + \text{Aq}$, and NH_4 salts + Aq .

Sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$.

Solubility in various salts + Aq .

Solvent	g. ZnCO_3 sol. in 1 l. of the solvent
10% $\text{NaNO}_3 + \text{Aq}$	0.058981
sat. $\text{NaNO}_3 + \text{Aq}$	0.149000
5% $\text{NaCl} + \text{Aq}$	0.021730
10% $\text{NaCl} + \text{Aq}$	0.046564
sat. $\text{NaCl} + \text{Aq}$	0.130380
10% $\text{Na}_2\text{SO}_4 + \text{Aq}$	0.009313
sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$	0.015521

(Ehlert, *Z. Elektrochem.* 1912, 18, 728.)

Insol. in liquid NH_3 . (Franklin, *Am. Ch. J.* 1898, 20, 830.)

Insol. in acetone. (Eidmann, *C. C.* 1899, II, 1014; Naumann, *B.* 1904, 37, 4329.)

Insol. in methyl acetate (Naumann, *B.* 1909, 42, 3790); ethyl acetate. (Naumann, *B.* 1910, 43, 314.)

Min. *Calamine*, *Smithsonite*.

Calamine is sol. in $\text{NH}_4\text{OH} + \text{Aq}$ only in the presence of NH_4 salts. (Brandhorst, *Zeit. angew. Ch.* 1904, 17, 513.)

+ $\frac{1}{2}\text{H}_2\text{O}$. (Mikusch, *Z. anorg.* 1908, 56, 386.)

+ H_2O . (Belar, *Zeit. Kryst.* 1890, 17, 126.)

Zinc carbonate ammonia, $\text{ZnCO}_3 \cdot \text{NH}_3$.

Slowly decomp. by H_2O , but not on the air, or by boiling with alcohol. (Favre, *A. ch.* (3) 10, 474.)

Zinc carbonate hydroxylamine, $\text{ZnCO}_3 \cdot 2\text{NH}_2\text{OH}$.

Insol. in H_2O . Decomp. by acids. (Goldschmidt and Syngros, *Z. anorg.* 5, 129.)

Zirconium carbonate, $3\text{ZrO}_2 \cdot \text{CO}_2 + 6\text{H}_2\text{O}$.

Decomp. by hot H_2O , all CO_2 being given off. (Hermann.)

Sol. in alkali carbonates + Aq .

Percarbonic acid.

See Percarbonic acid.

Carbonic anhydride, CO₂.

See Carbon dioxide.

Carbonophosphoric acid.**Potassium carbonophosphate, (K₂HPO₄)₂, 2CO₂, 2KHCO₃.**

Known only in solution. (Barillé, C. R. 1903, 137. 566.)

Carbonyl bromide, COBr₂.

Decomp. by H₂O. (Besson, C. R. 1895, 120. 192.)

Carbonyl platinous bromide, CO, PtBr₂.

Sol. in H₂O with almost instant decomp. Sol. in absolute alcohol. (Pullinger, Chem. Soc. 59. 603.)

Quite easily sol. in hot C₆H₆, insol. in ligroine, and can be crystallized from CCl₄. Very easily sol. in HBr + Aq. (Mylius and Förster, B. 24. 2432.)

Carbonyl bromochloride, COClBr.

Decomp. by H₂O. (Besson.)

Carbonyl chloride, COCl₂.

Phosgene. Cold H₂O dissolves 1–2 vols. COCl₂ gas with slow decomposition. Alcohol decomp. immediately. Immediately absorbed by KOH, or NH₄OH + Aq. Very sol. in glacial HC₂H₃O₂, benzene, and most liquid hydrocarbons. (Berthelot, Bull Soc. (2) 13. 14.)

Sol. in SCl₂.

1 vol. AsCl₃ absorbs 10 vols. COCl₂.

Dicarbonyl cuprous chloride, Cu₂Cl₂, 2CO + 4H₂O.

Decomp. by air. (Jones, Am. Ch. J. 1899, 22. 305.)

Carbonyl platinous chloride, 2COCl₂, PtCl₂.

Sl. deliquescent. Easily sol. in H₂O without decomp.; sl. sol. in alcohol. Almost insol. in CCl₄. (Pullinger, Chem. Soc. 59. 600.)

Monocarbonyl platinous chloride, CO, PtCl₂.

Decomp. by H₂O and alcohol; sol. in hot CCl₄. (Schützenberger, A. ch. (4) 15. 100.)

Sol. in conc. HCl + Aq. (Mylius and Förster.)

Dicarbonyl platinous chloride, 2CO, PtCl₂.

Decomp. by H₂O and alcohol. Sol. in CCl₄. (Schützenberger.)

Decomp. by conc. HCl + Aq into CO and CO, PtCl₂. (Mylius and Förster.)

Sesquicarbonyl platinous chloride, 3/2PtCl₂.

Decomp. by H₂O or alcohol. Much sol. in CCl₄ than 2CO, PtCl₂.

Carbonyl platinous iodide, CO, PtI₂.

Not hygroscopic. Insol. in, but slowly comp. by, H₂O. Easily sol. in benzene ether, also in alcohol, which decomp. on heating; sol. in HI + Aq. (Mylius and Förster.)

Carbonyl platinous sulphide, CO, PtS.

Easily decomp. Insol. in ordinary sol. (Mylius and Förster.)

Carbonyl sulphide, COS.

H₂O absorbs 1 vol. COS.

Absorption of COS by H₂O at t°

t°	Coefficient of absorption
0	1.333
10	0.835
20	0.561
30	0.403

(Winkler, Z. phys. Ch. 1906, 55. 35)

1 ccm. H₂O at 13.5° and 756 mm dissolves 0.8 ccm. COS. (Hempel, angew. ch. 1901, 14. 867.)

1 ccm. of a hydrochloric acid solution of Cu₂Cl₂ absorbs about 0.2 ccm. COS. (pel.)

Carbonyl ferrocyanhydric acid.

H₂Fe(CO)(CN)₄.

Very sol. in H₂O; decomp. on heating. (Müller, A. ch. (6) 17. 94.)

Cobalt carbonyl ferrocyanide.

Sl. sol. in H₂O; very sol. in dil. HNO₃. (M.)

Cupric carbonyl ferrocyanide,

Cu₂[Fe(CO)(CN)₄]₂.

Insol. in H₂O, H₂SO₄, or dil. HNO₃. (M.)

Iron (ferric) carbonyl ferrocyanide, FeFeCO(CN)₄.

Insol. in H₂O. Sol. in H₂C₂O₄ + Aq. in acetic, lactic, succinic, tartaric, and other acids + Aq, but easily sol. in the neutral solutions of those acids. Insol. in KCl, or KNO₃ but sensibly sol. in Na₂HPO₄ + Aq. even on warming in very dil. H₂SO₄, or in dil. HCl + Aq. (Müller.)

Potassium carbonyl ferrocyanide,

K₂Fe(CO)(CN)₄ + 3½H₂O.

100 pts. H₂O dissolve 148 pts. (Müller, C. R. 104. 992.)

nyl ferrocyanide,
 $(\text{CO})(\text{CN})_5$.

H_2O ; sl. sol. in dil. H_2SO_4 , HCl ,
 Aq; scarcely attacked by conc.
 Aq. (Müller.)

bonyl ferrocyanide,
 $(\text{CO})(\text{CN})_5 + 6\text{H}_2\text{O}$.
 H_2O . (Müller.)

onyl ferrocyanide,
 $[\text{FeCO}(\text{CN})_5]_2 + 5\text{H}_2\text{O}$.

H_2O , but more easily if H_2O is
 th $\text{HC}_2\text{H}_3\text{O}_2$.

gistic acid.

cericotungstate, $2(\text{NH}_4)_2\text{O}$,
 $16\text{WO}_3 + 2\text{H}_2\text{O}$.

H_2O , but decomp. by boiling
 (Smith, J. Am. Chem. Soc. 1904,

e.

pure H_2O very slowly at ordinary
 attacked by cold conc. H_2SO_4 or
 HNO_3 . Sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$,
 and conc. or dil. $\text{HCl} + \text{Aq}$.
 and Norton, Pogg. 155. 633.)

mide, CeBr_3 .

us. As the chloride. (Robinson,
 Soc. 37. 150.)

acetone. (Eidmann, C. C. 1899,
 Naumann, B. 1904, 37. 4328.)

methyl acetate. (Naumann, B.
 790.)

Very deliquescent. (Jolin.)

d bromide, CeBr_3 , $\text{AuBr}_3 + 8\text{H}_2\text{O}$.

aurate, cerium.

bide, CeC_2 .

by fused alkali nitrates, chlorates,
 and carbonates; and by conc.
 heating. Insol. in conc. HNO_3 ;
 y H_2O and dil. acids. (Moissan,
 122. 359.)

Not attacked by hot conc. acids.
 ine, J. B. 1865. 176.)

oride, CeCl_3 .

us. Deliquescent. Sol. in H_2O
 ig and evolution of heat; sol. in

acetone. (Eidmann, C. C. 1899,
 Naumann, B. 1904, 37. 4328.)

ly scl. in methyl acetate. (Nau-
 1909, 42. 3790.)

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Den-
 rg. 1894, 7. 260.)

H_2O . Deliquescent. (Berzelius.)

by boiling with H_2O . Sol. in 1 pt.
 rd. temp. and 3-4 pts. alcohol.

Ceric chloride.

Known only in solution, which decom-
 poses by slight heat. (Berzelius.)

Cerous mercuric chloride.

Not deliquescent. (v. Bonsdorff.)

$\text{CeCl}_3, 4\text{HgCl}_2 + 10\text{H}_2\text{O}$. Permanent; eas-
 ily sol. in H_2O . (Jolin, Bull. Sec. (2) 21. 533.)

Cerium stannic chloride.

See Chlorostannate, cerium.

Cerous chloride zinc iodide.

Sol. in H_2O and alcohol. (Holzmann, J. pr.
 84. 76.)

Cerous fluoride, CeF_3 .

Insol. ppt.
 $+ \frac{1}{2}\text{H}_2\text{O}$.

Ceric fluoride, CeF_4 .

Insoluble precipitate. (Berzelius.)

$+ \text{H}_2\text{O}$ Insol. in H_2O . (Brauner, B. 14.
 1944.)

Ceric cobaltous fluoride, $2\text{CeF}_4, \text{CoF}_2 + 7\text{H}_2\text{O}$.

Ppt. Easily decomp. by H_2O . (Rimbach,
 A. 1909, 368. 107.)

Ceric cupric fluoride, $2\text{CeF}_4, \text{CuF}_2 + 7\text{H}_2\text{O}$.

Ppt. Decomp. by H_2O . (Rimbach, l. c.)

Ceric nickel fluoride, $2\text{CeF}_4, \text{NiF}_2 + 7\text{H}_2\text{O}$.

Ppt. Decomp. by H_2O . (Rimbach, l. c.)

Ceric potassium fluoride, $2\text{CeF}_4, 3\text{KF} + 2\text{H}_2\text{O}$.

Insol. in H_2O . (Brauner, B. 14. 1944; 15.
 109.)

Could not be obtained pure. (Rimbach,
 l. c.)

Ceric zinc fluoride, $2\text{CeF}_4, \text{ZnF}_2 + 7\text{H}_2\text{O}$.

Ppt. Decomp. by H_2O . (Rimbach, l. c.)

Ceroceric fluoride, $2\text{CeF}_3, \text{CeF}_4$.

Min. *Fluocerite*.

Cerium hydride, CeH_2 .

Decomp. by acids. (Winkler, B. 24. 873.)

CeH_2 . Decomp. in moist air; decomp. by
 hot or cold H_2O ; sol. in acids with evolution
 of H_2 . Decomp. by alkalis. (Muthmann,
 A. 1902, 325. 266.)

Cerous hydroxide, $\text{Ce}_2\text{O}_3, x\text{H}_2\text{O}$.

Easily sol. in acids. Insol. in excess of
 alkali hydroxides + Aq. Sol. in $(\text{NH}_4)_2\text{CO}_3 +$
 Aq.

100 ccm. of a solution in glycerine + Aq
 containing about 60% by vol. of glycerine
 contain 7.9 g. Ce_2O_3 . (Müller, Z. anorg.
 1905, 43. 322.)

Exists in two modifications: one insol. in

cold HCl+Aq; the other sol. in cold HCl+Aq. (Brauner, C. N. 1895, 71. 283.)

Ceric hydroxide, $2\text{CeO}_2, 3\text{H}_2\text{O}$.

Sol. in HNO_3 or H_2SO_4 ; also in HCl+Aq, forming cerous chloride and free chlorine. Insol. in hydrofluoric, acetic, or formic acids+Aq. Somewhat sol. in dil. HNO_3 or HCl+Aq. (Ordway, Am. J. Sci. (2) 26. 205.) Insol. in NH_4OH , KOH, and NaOH+Aq. Sl. sol. in alkali carbonates+Aq. (Dumas.)

Sl. sol. in $(\text{NH}_4)_2\text{CO}_3$ +Aq. (Ordway.)

100 ccm. of a solution in glycerine+Aq containing about 60% by vol. of glycerine contain 0.08 g. CeO_2 . (Müller, Z. anorg. 1905, 43. 232.)

Cerous iodide, CeI_3 .

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4328.)

+9 H_2O . Very deliquescent and sol. in H_2O . (Lange, J. pr. 82. 134.)

Sol. in alcohol.

Cerium nitride, CeN .

Decomp. by H_2O and alkali. Sol. in mineral acids with formation of cerous and ammonium salts. (Muthmann, A. 1902, 325. 272.)

Cerous oxide, Ce_2O_3 .

When ignited, insol. in HCl+Aq; when long digested with H_2SO_4 , is sol. in HCl+Aq with addition of alcohol.

Ceric oxide, CeO_2 .

When ignited, is only dissolved in traces, even on heating, by HCl or HNO_3 +Aq. Sol. in conc. H_2SO_4 when warmed. Sol. in the cold in a solution of KI in HCl+Aq (Bunsen), in a mixture of HCl and FeCl_3 +Aq, or any reducing substance.

Cerium peroxide, Ce_4O_7 .

Insol. in boiling conc. acids. Sol. in H_2SO_4 by long digestion. (Popp, A. 131. 361.)

Probably does not exist. (Rammelsberg, Pogg. 108. 40.)

Ce_2O_3 . (Hermann, J. pr. 30. 184.)

Probably does not exist. (Rammelsberg.)

$\text{CeO}_2 + x\text{H}_2\text{O}$. Sol. in HCl+Aq. (Popp, A. 131. 361); (Lecoq de Boisbaudran, C. R. 100. 605.)

$\text{CeO}_2 + \text{H}_2\text{O}_2$, according to Cleve (Bull. Soc. (2) 43. 57.)

Cerium oxycarbide, $\text{CeC}_2, 2\text{CeO}_2$.

Stable in the air. Slowly attacked by cold H_2O . With hot H_2O and with acids, it gives unsat. hydrocarbons. (Sterba, C. R. 1902, 134. 1058.)

Cerium oxychloride, CeOCl .

Slightly attacked by hot conc. HCl+Aq. Slowly sol. in conc. HNO_3 +Aq. (Wöhler.)

Easily sol. in dil. acids. (Didier, C. 882.)

Cerium oxychloride tungsten trioxide, WO_3 .

(Didier, C. R. 102. 823.)

Cerium selenide.

Insol. in H_2O ; difficultly sol. in (Berzelius.)

Cerium silicide, CeSi_2 .

Insol. in H_2O , by which it is acted only very slowly.

Sol. in HCl and HF+Aq with evolution of H_2 .

Not attacked by alkalis+Aq or N. Aq.

Insol. in organic solvents. (Sterba 1902, 135. 170.)

Cerium silicide, Ce_3Si_2 .

Insol. in acids. (Ullik, W. A. E. 115.)

Cerium disulphide, Ce_2S_4 .

Not decomp. by cold H_2O .

Slowly sol. in cold dil. H_2SO_4 , in acetic acid. Rapidly sol. in warm dil. HCl and acetic acid with decomp. (1908, 41. 3342.)

Cerium sesquisulphide, Ce_2S_3 .

Insol. in, and not decomp. by, 1 easily decomp. by the weakest acid (sander); (Didier, C. R. 100. 1461.)

Monochloramine, NH_2Cl .

Easily sol. in H_2O . (Raschig. Ch. 31. 926.)

Chlorotetramine comps.

See Chlorotetramine comps.

Metachlorantimonic acid, $\text{HSbCl}_4, 4\frac{1}{2}\text{H}_2\text{O}$.

Hydroscopic. Sol. in H_2O with sol. in alcohol, acetone, and acetic acid. (Weinland, Z. anorg. 1905, 44. 43.)

Metachlorantimonic acid ammonia, 2NH_3 .

Sol. in H_2O and in alcohol. (Weinland, Z. anorg. 1905, 44. 59.)

Aluminum metachlorantimonate, $\text{Al} + 15\text{H}_2\text{O}$.

Hydroscopic.

Sol. in dil. HCl+Aq. (Weinland, 36. 254.)

<i>metachlorantimonate</i> , NH_4SbCl_6	Sol. in dil. $\text{HCl} + \text{Aq.}$ (Weinland, B. 1903, 36. 250.)
pic. O. Solution decomp. slowly when y on warming.	Rubidium metachlorantimonate , RbSbCl_6 .
il. HCl . (Weinland, B. 1903, 36.)	Hydroscopic. Sol. in dil. $\text{HCl} + \text{Aq.}$ (Weinland.)
<i>metachlorantimonate ammonia</i> , Cl_3 , 7NH_3 .	Silver metachlorantimonate ammonia , $\text{AgSbCl}_6, 2\text{NH}_3$.
Weinland and Schmid, Z. anorg. 3.)	Decomp. by H_2O . (Weinland and Schmid.)
<i>etachlorantimonate</i> , $\text{Ca}(\text{SbCl}_4)_2 +$	Zinc metachlorantimonate ammonia , $\text{Zn}(\text{SbCl}_4)_2, 4\text{NH}_3$.
pic.	(Weinland and Schmid.)
il. $\text{HCl} + \text{Aq.}$ (Weinland, B. 1903,	Chlorarsenious acid .
	See Arsenyl chloride.
<i>metachlorantimonate</i> , $\text{Cr}(\text{SbCl}_4)_3$, H_2O .	Chlorauric acid , $\text{HAuCl}_4 + 4\text{H}_2\text{O}$.
pic.	Sol. in H_2O , alcohol, and ether.
il. $\text{HCl} + \text{Aq.}$ (Weinland.)	Sol. in POCl_3 . (Walden, Z. anorg. 1900, 25. 212.)
<i>orthochlorantimonate</i> , $\text{CrSbCl}_5 +$	Difficultly sol. in PCl_3 . (Walden.)
pic.	Cryst. with $3\text{H}_2\text{O}$ as stated by Weber and Schottlander and not with $4\text{H}_2\text{O}$ as stated by Thomsen. (Schmidt, C. C. 1906, II. 855.)
il. $\text{HCl} + \text{Aq.}$ (Weinland.)	Chloraurates .
<i>chlorantimonate ammonia</i> , Cl_3 , 5NH_3 .	All chloraurates are easily sol. in H_2O and in alcohol. (v. Bonsdorff, 1829.)
id and Schmid, Z. anorg. 1905, 44.	Ammonium chloraurate , $\text{NH}_4\text{AuCl}_4 + \text{H}_2\text{O}$.
<i>metachlorantimonate</i> , $\text{Gl}(\text{SbCl}_4)_3$, H_2O .	Very easily sol. in H_2O .
rescopic.	+ $2\text{H}_2\text{O}$. Very easily sol. in H_2O .
il. HCl . (Weinland, B. 1903, 36.)	Barium chloraurate , $\text{Ba}(\text{AuCl}_4)_2 + x\text{H}_2\text{O}$.
c) <i>orthochlorantimonate</i> , FeSbCl_5 , O .	Deliquescent in moist air. Sol. in H_2O and alcohol. (v. Bonsdorff, Pogg. 17. 261.)
pic.	Cadmium chloraurate .
ily be cryst. from dil. $\text{HCl} + \text{Aq.}$)	Not deliquescent. Sol. in H_2O and alcohol. (v. Bonsdorff.)
<i>etachlorantimonate</i> , $\text{LiSbCl}_4 +$	Cæsium chloraurate , CsAuCl_4 .
droscopic.	100 pts. aqueous sat. solution contain at:
il. $\text{HCl} + \text{Aq.}$ (Weinland, l. c.)	10° 20° 30° 40° 50°
n <i>pyrochlorantimonate</i> , MgSbCl_5 , O .	0.5 0.8 1.7 3.2 5.4 pts. anhydrous salt,
pic.	60° 70° 80° 90° 100°
il. $\text{HCl} + \text{Aq.}$ (Weinland.)	8.2 12.0 16.3 21.7 27.5 pts. anhydrous salt.
<i>achlorantimonate ammonia</i> , SbCl_4 , 6NH_3 .	(Rosenblatt, B. 19. 2538.)
H_2O . (Weinland and Schmid, Z. 5, 44. 57.)	+ $\frac{1}{2}\text{H}_2\text{O}$. (Wells and Wheeler, Am. J. (3) 44. 157.)
<i>metachlorantimonate</i> , $\text{KSbCl}_4 +$	Calcium chloraurate , $\text{Ca}(\text{AuCl}_4)_2 + 6\text{H}_2\text{O}$.
pic.	Deliquescent. Sol. in H_2O and alcohol. (v. Bonsdorff.)
H_2O . Solution decomp. slowly when ly when warmed.	Cerium chloraurate , $\text{CeCl}_3, \text{AuCl}_3 + 10\text{H}_2\text{O}$.
	Extremely deliquescent. Easily sol. in H_2O and absolute alcohol. (Holzmann, C. C. 1863. 206.)
	+ $13\text{H}_2\text{O}$. (Jolin, Bull. Soc. (2) 21. 534.)

Cobalt chloraurate, $\text{Co}(\text{AuCl}_4)_2 + 8\text{H}_2\text{O}$.

Sol. in H_2O and alcohol. (Topsoë.)

Didymium chloraurate, $\text{DiCl}_2, \text{AuCl}_3 + 10\text{H}_2\text{O}$.

Very deliquescent. (Cleve, Bull. Soc. (2) 43. 361.)

$2\text{DiCl}_2, 3\text{AuCl}_3 + 20\text{H}_2\text{O}$. (Cleve.)

Gadolinium chloraurate, $\text{GdCl}_3, \text{AuCl}_3 + 10\text{H}_2\text{O}$.

Sol. in H_2O . (Benedicks, Z. anorg. 1900, 22. 404.)

Lanthanum chloraurate, $\text{LaCl}_3, \text{AuCl}_3 + 5\text{H}_2\text{O}$.

Deliquescent in moist air. Sol. in H_2O . (Cleve, B. 8. 128.)

Lithium chloraurate, LiAuCl_4 .

100 pts. aqueous solution contain at:

10°	20°	30°	40°
53.1	57.7	62.5	67.3

pts. anhydrous salt,

50°	60°	70°	80°
72.0	76.4	81.0	85.7

pts. anhydrous salt. (Rosenblatt.)

+ $2\text{H}_2\text{O}$. (Antony and Lucchesi, Gazz. ch. it. 1890, 20. 601.)

+ $4\text{H}_2\text{O}$. Not stable.

Sol. in H_2O and alcohol. (Fasbender, C. C. 1894, I. 409.)

Magnesium chloraurate, $\text{Mg}(\text{AuCl}_4)_2 + 8\text{H}_2\text{O}$.

Somewhat deliquescent. Sol. in H_2O and alcohol. (Topsoë.)

+ $12\text{H}_2\text{O}$.

Manganese chloraurate, $\text{Mn}(\text{AuCl}_4)_2 + 8\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O and alcohol. (Topsoë.)

+ $12\text{H}_2\text{O}$.

Nickel chloraurate, $\text{Ni}(\text{AuCl}_4)_2 + 8\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O and alcohol. (Topsoë.)

Potassium chloraurate, KAuCl_4 .

Anhydrous. Very stable. (Lainer, W. A. B. 99, 2b. 247.)

100 pts. solution in H_2O contain at:

10°	20°	30°
27.7	38.2	48.7

pts. anhydrous salt,

40°	50°	60°
59.2	70.0	80.2

pts. anhydrous salt.

(Rosenblatt, B. 19. 2538.)

Sol. in H_2O and alcohol; insol. in ether. (Fasbender, C. C. 1894, I. 409.)

1 pt. is sol. in 4 pts. 98% alcohol. (Fasbender, C. C. 1894, II. 609.)

+ $2\text{H}_2\text{O}$. Efflorescent.

+ $\frac{1}{2}\text{H}_2\text{O}$.

Praseodymium chloraurate, $\text{PrCl}_3, \text{AuCl}_3 + 10\text{H}_2\text{O}$.

Very sol. in H_2O ; sol. in conc. HCl . (Schule, Z. anorg. 1898, 18. 354.)

Rubidium chloraurate, RbAuCl_4 .

100 pts. sat. $\text{RbAuCl}_4 + \text{Aq}$ contain

10°	20°	30°	40°	50°
4.6	9.0	13.4	17.7	22.2

pts. anhydrous

60°	70°	80°	90°	100°
26.6	31.0	35.3	39.7	44.2

pts. anhydrous

(Rosenblatt.)

1 pt. sol. in 54 pts. 98% alcohol. 1 ether. (Fasbender, C. C. 1894, II. 60)

Samarium chloraurate, $\text{SmCl}_3, \text{AuCl}_3 + 10\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O . (Bull. Soc. (2) 43. 165.)

Scandium chloraurate, $3\text{ScCl}_3, 2\text{AuCl}_3 + 21\text{H}_2\text{O}$.

Very deliquescent. (Crookes, Phil. 1910, 210. A, 365.)

Silver chloraurate, AgAuCl_4 .

Decomp. in the air.

Decomp. by H_2O , HCl and NH_3 . (mann, B. 1894, 27. 597.)

Sodium chloraurate, $\text{NaAuCl}_4 + 2\text{H}_2\text{O}$

Easily sol. in H_2O and absolute alc

10°	20°	30°
58.2	60.2	64.0

pts. anhydrous

40°	50°	60°
69.4	77.5	90.0

pts. anhydrous

(Rosenblatt.)

Easily sol. in $\text{NaCl} + \text{Aq}$.

Easily sol. in H_2O , alcohol and ether. (Fasbender, C. C. 1894, I. 409.)

Strontium chloraurate.

Sol. in H_2O . (v. Bonsdorff.)

Thallium chloraurate.

(Carstanjen.)

Ytterbium chloraurate, $\text{YbCl}_3, \text{AuCl}_3$ -

Ppt. (Cleve, Z. anorg. 1902, 22. 13)

Yttrium chloraurate, $\text{YtCl}_3, 2\text{AuCl}_3 +$

Very sol. in H_2O . (Cleve.)

Zinc chloraurate, $\text{Zn}(\text{AuCl}_4)_2 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Topsoë.)

+ $12\text{H}_2\text{O}$. Sol. in H_2O and alcohol. (Bonsdorff.)

Chlorauricyanhydric acid.

Barium chlorauricyanide, Ba[Au(CN)₂Cl₂]₂ + 8H₂O.

Very sol. in H₂O or alcohol. (Lindbom, and Univ. Arsk. 12. No. 6.)

Potassium chlorauricyanide, KAu(CN)₂Cl₂ + H₂O.

Very sol. in H₂O or alcohol.

Strontium chlorauricyanide, Sr[Au(CN)₂Cl₂]₂ + 8H₂O.

Sol. in H₂O.

Zinc chlorauricyanide, Zn[Au(CN)₂Cl₂]₂ + 7H₂O.

Very sol. in H₂O.

Chlorhydric acid, HCl.

Liquid. Miscible with liquid CO₂, and H₂S.

Gas. Absorbed by H₂O with production of much heat.

H₂O absorbs 400–500 vols. at ord. temp. and pressure or a little less than 1 pt. by weight. (Dalton.)

1 vol. H₂O absorbs 480 vols. at 0°; sp. gr. of sat. solution is 1.2109. (Davy.)

1 vol. H₂O absorbs 417 822 vols. at 20°, the vol. increasing to 1.4138 vols.; 1 vol. of HCl + Aq then contains 311 vols. HCl, has sp. gr. 1.1958, and contains 40.39% HCl by weight. (Thomson, 1831.)

1 vol. H₂O absorbs 464 vols. and sat. solution has 1.21 sp. gr. and contains 42.4% HCl by weight. (Wittstein.)

H₂O sat. at 0° contains 480 times its vol. of HCl, and sp. gr. = 1.2109; sat. at ord. temp. contains 38.3% of its weight in HCl, and sp. gr. = 1.192. (Berzelius.)

1 vol. H₂O absorbs V vols. HCl at t° and 760 mm. pressure, and the liquid formed has the given sp. gr., and contains the given per cent HCl.

t°	V	Sp. gr.	% HCl
0	525.2	1.2257	45.148
4	494.7	1.2265	44.361
8	480.3	1.2185	43.828
12	471.3	1.2148	43.277
14	462.4	1.2074	42.829
18	451.2	1.2064	42.344
18 25	450.7	1.2056	42.283
23	435.0	1.2014	41.536

(Deicke, Pogg. 119. 156.)

At 760 mm. pressure 1 g. H₂O absorbs g. HCl at t°.

t°	g. HCl	t°	g. HCl	t°	g. HCl
0	0.825	22	0.710	44	0.618
2	0.814	24	0.700	46	0.611
4	0.804	26	0.691	48	0.603
6	0.793	28	0.682	50	0.596
8	0.783	30	0.673	52	0.589
10	0.772	32	0.665	54	0.582
12	0.762	34	0.657	56	0.575
14	0.752	36	0.649	58	0.568
16	0.742	38	0.641	60	0.561
18	0.731	40	0.633
20	0.721	42	0.626

(Roscoe and Dittmar.)

Conc. HCl + Aq loses HCl, and dil. HCl + Aq loses H₂O on warming, until an acid of constant composition is formed, containing 20.18% HCl, with a sp. gr. of 1.101 at 15°, which can be distilled unchanged at 110°. (Bineau, A. ch. (3) 7. 257.)

The above is true if barometer is at 760 mm., but the composition changes with the pressure as follows—

Mm. Hg	% HCl	Mm. Hg	% HCl	Mm. Hg	% HCl
50	23.2	800	20.2	1700	18.8
100	22.9	900	19.9	1800	18.7
200	22.3	1000	19.7	1900	18.6
300	21.8	1100	19.5	2000	18.5
400	21.4	1200	19.4	2100	18.4
500	21.1	1300	19.3	2200	18.3
600	20.7	1400	19.1	2300	18.2
700	20.4	1500	19.0	2400	18.1
760	20.24	1600	18.9	2500	18.0

(Roscoe and Dittmar.)

Conc. HCl + Aq gradually gives off HCl on the air until it has a sp. gr. 1.128 at 15°, and contains 25.2% HCl. (Bineau, l. c.)

According to Roscoe and Dittmar, this depends on the temperature. If a current of air is passed through HCl + Aq, acid or water is given off according as the acid is strong or weak, until an acid of constant composition for a given temperature is formed, as follows—

Temp.	% HCl	Temp.	% HCl	Temp.	% HCl
0°	25.0	35°	23.9	70°	22.6
5	24.9	40	23.8	75	22.3
10	24.7	45	23.6	80	22.0
15	24.6	50	23.4	85	21.7
20	24.4	55	23.2	90	21.4
25	24.3	60	23.0	95	21.1
30	24.1	65	22.8	100	20.7

From the above it is seen that the acid which distils unchanged at a given pressure, that is, boils at a certain constant temperature, is identical with the acid which undergoes no change in composition by a current of dry air at the same temperature, and under the ordinary pressure, thus—

Mm. Hg	B.-pt.	% HCl	Temp. of air current	% HCl
100	61–62°	22.8	62°	22.9
200	76–77	22.1	77	22.2
300	84–85	21.7	85	21.7
380	91	21.3	91	21.4
490	97	20.9	98	21.1
620	103	20.6

(Roscoe and Dittmar.)

Solubility of HCl in H₂O at 0° under different degrees of pressure. P = partial pressure in mm. Hg, i. e., total pressure minus the tension of aqueous vapour at the given temp.; G = grammes of HCl dissolved in 1 g. H₂O at the pressure P and 0° temp.

P	G	P	G
60	0.613	350	0.751
70	0.628	400	0.763
80	0.640	450	0.772
90	0.649	500	0.782
100	0.657	550	0.791
110	0.664	600	0.800
120	0.670	650	0.808
130	0.676	700	0.817
140	0.681	750	0.824
150	0.686	800	0.831
175	0.697	900	0.844
200	0.707	1000	0.856
225	0.716	1100	0.869
250	0.724	1200	0.882
275	0.732	1300	0.895
300	0.738

(Roscoe and Dittmar, A. 112. 334.)

1 vol. H₂O dissolves 560 vols. HCl at -12°
" " 500 " " 0°
" " 440 " " +20°

(Berthelot, C. R. 76. 779.)

1 vol. H₂O absorbs 480 vols. HCl at 15° to form a solution containing 42.85% HCl with a sp. gr. of 1.215. (Hager.)

Solubility of HCl at low temperatures, and 760 mm. pressure.

t°	Pts. HCl in 1 pt. H ₂ O	t°	Pts. HCl in 1 pt. H ₂ O
0	0.842	-18	0.957
- 5	0.864	-19	0.965
-10	0.898	-20	0.974
-15	0.933	-21	0.983
-17	0.949	-24	1.012

(Roozeboom, R. t. c. 1884, 3. 79.)

Solubility in H₂O at t°.

t°	%HCl
50	61.65
45	61.76
40	62.27
35	62.90
30	63.21
20	64.19
15	64.70
10	65.18
5	65.48
0	65.85
- 5	66.44
-10	66.71
-50	67.29
-20	67.65

(Rupert, J. Am. Chem. Soc. 1909, 31. 860.)

Sp. gr. of HCl + Aq.

Sp. gr.	% HCl	Sp. gr.	% HCl	Sp. gr.	% HCl
1.203	40.66	1.1285	27.21	1.0000	20.44
1.179	37.00	1.1197	25.52	1.0002	19.47
1.162	33.95	1.1127	24.03	1.0000	18.90
1.149	31.35	1.1060	22.70	1.0020	17.79
1.139	29.13	1.1008	21.51	1.0780	17.06

(Thomson, in his System, 2. 189.)

Sp. gr. of HCl + Aq.

Sp. gr.	% HCl	Sp. gr.	% HCl
1.21	42.43	1.10	20.20
1.20	40.80	1.09	18.18
1.19	38.38	1.08	16.16
1.18	36.36	1.07	14.14
1.17	34.34	1.06	12.12
1.16	32.32	1.05	10.10
1.15	30.30	1.04	8.08
1.14	28.28	1.03	6.06
1.13	26.26	1.02	4.04
1.12	24.24	1.01	2.02
1.11	20.30

(Edm. Davy.)

Sp. gr. of HCl + Aq.

Sp. gr.	% HCl	B.-pt.	Sp. gr.	% HCl	B.-pt.
1.199	34.01	49°	1.094	16.08	111°
1.181	31.09	65	1.075	13.16	109
1.166	28.29	76	1.064	11.16	107
1.154	26.57	87	1.047	8.62	105
1.144	24.84	100	1.035	6.92	104
1.136	23.25	103	1.018	3.52	102
1.127	21.06	105	1.009	1.86	101
1.121	20.74	109

(Kirwan and Dalton.)

Sp. gr. of HCl + Aq at 15°.

% HCl	Sp. gr.	% HCl	Sp. gr.
2.22	1.0103	29.72	1.1504
3.80	1.0189	31.50	1.1588
6.26	1.0310	34.24	1.1730
11.02	1.0557	36.63	1.1844
15.20	1.0751	38.67	1.1936
18.67	1.0942	40.51	1.2021
20.91	1.1048	41.72	1.2074
23.72	1.1196	43.09	1.2124
25.96	1.1308

(Kolb, C. R. 74. 337.)

Sp. gr. of HCl + Aq at 15°.

Sp. gr.	% HCl	Sp. gr.	% HCl	Sp. gr.	% HCl
1.2000	40.777	1.1620	32.621	1.1206	24.446
1.1982	40.309	1.1599	32.213	1.1185	24.058
1.1964	39.961	1.1578	31.805	1.1164	23.650
1.1946	39.554	1.1557	31.398	1.1143	23.242
1.1928	39.146	1.1536	30.990	1.1123	22.834
1.1910	38.738	1.1515	30.582	1.1102	22.426
1.1893	38.330	1.1494	30.174	1.1082	22.019
1.1875	37.923	1.1473	29.767	1.1061	21.611
1.1859	37.516	1.1452	29.359	1.1041	21.203
1.1846	37.108	1.1431	28.951	1.1020	20.796
1.1822	36.700	1.1410	28.544	1.1000	20.388
1.1802	36.292	1.1389	28.136	1.0980	19.980
1.1782	35.884	1.1368	27.728	1.0960	19.572
1.1762	35.476	1.1349	27.321	1.0939	19.165
1.1741	35.068	1.1328	26.913	1.0919	18.757
1.1721	34.660	1.1308	26.505	1.0899	18.349
1.1701	34.252	1.1287	26.098	1.0879	17.941
1.1681	33.845	1.1267	25.690	1.0859	17.534
1.1661	33.437	1.1247	25.282	1.0838	17.126
1.1641	33.029	1.1226	24.874	1.0818	16.718

Sp. gr. of HCl+Aq at 15°—Continued.

gr.	% HCl	Sp. gr.	% HCl	Sp. gr.	% HCl
98	16.310	1.0517	10.602	1.0259	5.301
78	15.902	1.0497	10.194	1.0239	4.893
58	15.494	1.0477	9.786	1.0220	4.486
38	15.087	1.0457	9.379	1.0200	4.078
18	14.679	1.0437	8.971	1.0180	3.670
97	14.271	1.0417	8.563	1.0160	3.262
77	13.863	1.0397	8.155	1.0140	2.854
57	13.456	1.0377	7.747	1.0120	2.447
37	13.409	1.0357	7.340	1.0100	2.039
17	12.641	1.0337	6.932	1.0080	1.631
97	12.233	1.0318	6.524	1.0060	1.224
577	11.825	1.0298	6.116	1.0040	0.816
557	11.418	1.0279	5.709	1.0020	0.408
537	11.010

(Ure, Handwörterbuch.)

gr. of HCl+Aq. U=sp. gr. at 15.55° according to Ure; K=sp. gr. at 15° according to Kremers.

HCl	U	K	%HCl	U	K
1	1.005	1.005	22	1 109	1.111
2	1.010	1 010	23	1 114	1 116
3	1.015	1.015	24	1 119	1.121
4	1.020	1.020	25	1 124	1.126
5	1.025	1.025	26	1 128	1.131
6	1.030	1.030	27	1 133	1.136
7	1.034	1.034	28	1 138	1.141
8	1.039	1.039	29	1 143	1.146
9	1.044	1.044	30	1 147	1.151
10	1.048	1.048	31	1 153	1.157
11	1.053	1.053	32	1 157	1.163
12	1.059	1.059	33	1 163	1.169
13	1.064	1 065	34	1 169	1.179
14	1.069	1.070	35	1 174	...
15	1.074	1.075	36	1 179	...
16	1.079	1 080	37	1 183	...
17	1.084	1.085	38	1 188	...
18	1.089	1.090	39	1 193	...
19	1.094	1.095	40	1 197	...
20	1.098	1.100	41	1 203	...
21	1.104	1.105

(Calculated by Gerlach, Z. anal. 8. 292.)

p. gr. of HCl+Aq at 15° (H₂O at 0°=1).

	Sp. gr.	% HCl	Sp. gr.	% HCl	Sp. gr.
	0.9992	15	1.07539	30	1.15079
	1.00503	16	1.08042	31	1.15581
	1.01005	17	1.08545	32	1.16084
	1.01508	18	1.09047	33	1.16587
	1.02010	19	1.09550	34	1.17089
	1.02513	20	1.10052	35	1.17592
	1.03016	21	1.10555	36	1.18095
	1.03518	22	1.11058	37	1.18597
	1.04021	23	1.11560	38	1.191
	1.04524	24	1.12063	39	1.196
	1.05026	25	1.12566	40	1.200
	1.05529	26	1.13068	41	1.204
	1.06031	27	1.13571	42	1.208
	1.06534	28	1.14074	43	1.212
	1.07037	29	1.14516

lb, recalculated by Gerlach, Z. anal. 27. 316.)

Sp. gr. of HCl+Aq at 15°.

% HCl	Sp. gr.	% HCl	Sp. gr.	% HCl	Sp. gr.
5	1.0244	20	1 0982	35	1.1739
10	1.0488	25	1.1234	40	1.1969
15	1.0733	30	1.1488	41	1.2013

(Hager, Adjumenta varia, Leipzig, 1876.)

Sp. gr. of HCl+Aq at 15° (H₂O at 15°=1).

% HCl	Sp. gr.	%HCl	Sp. gr.
44.345	1.21479	34.464	1.17138
43.136	1.21076	25.260	1.12479
41.901	1.20430	19.688	1.09675
41.212	1.20204	14.788	1.07255
39.831	1.19703	6.382	1.03150
37.596	1.18687

(Pickering, B. 26. 277.)

Most accurate table.

Sp. gr. of HCl+Aq at 15° (H₂O at 4°=1).

Sp. gr.	% HCl	Kg. HCl in 1 l.	Sp. gr.	% HCl	Kg.HCl in 1l.
1.000	0.16	0.016	1.105	20.97	0.232
1.005	1.15	0.012	1.110	21.92	0.243
1.010	2.14	0.022	1.115	22.86	0.255
1.015	3.12	0.032	1.120	23.82	0.267
1.020	4.13	0.042	1.125	24.78	0.278
1.025	5.15	0.053	1.130	25.75	0.291
1.030	6.15	0.064	1.135	26.70	0.303
1.035	7.15	0.074	1.140	27.66	0.315
1.040	8.16	0.085	1.145	28.61	0.322
1.045	9.16	0.096	1.150	29.57	0.340
1.050	10.17	0.107	1.155	30.55	0.353
1.055	11.18	0.118	1.160	31.52	0.366
1.060	12.19	0.129	1.165	32.49	0.379
1.065	13.19	0.141	1.170	33.46	0.392
1.070	14.17	0.152	1.175	34.42	0.404
1.075	15.16	0.163	1.180	35.39	0.418
1.080	16.15	0.174	1.185	36.31	0.430
1.085	17.13	0.186	1.190	37.23	0.443
1.090	18.11	0.197	1.195	38.16	0.456
1.095	19.06	0.209	1.200	39.11	0.469
1.100	20.01	0.220

(Lunge and Marchlewski, Z. angew. Ch. 1891. 133.)

Sp. gr. of HCl+Aq at room temp.

% HCl	Sp. gr.
8.14	1.0370
16.125	1.0843
23.045	1.1138

(Wagner, W. Ann. 1883, 18. 264.)

Relation of sp. gr. of HCl+Aq at t° to sp. gr. at 19.5° = 1.0.					
t°	8.9 % HCl sp. gr. = 1.0401	16.6 % HCl sp. gr. = 1.0704	25.5 % HCl sp. gr. = 1.101	35.8 % HCl sp. gr. = 1.133	46.6 % HCl sp. gr. = 1.166
0	0.99557	0.99379	0.99221	0.99079	0.98982
19.5	1.00000	1.00000	1.00000	1.00000	1.00000
40	1.00707	1.00781	1.00877	1.00990	1.01063
60	1.01588	1.01665	1.01794	1.01969	1.02180
80	1.02639	1.02676	1.02791	1.02986	...
100	1.03855	1.03801	1.03867	1.04059	...

(Kremers, Pogg. 108. 115.)

Sp. gr. of HCl+Aq.			Sp. gr. of HCl+Aq at 20°.		
G. equivalents HCl per liter.	t°	Sp. gr. t°/t°	Normality of HCl+Aq.	% HCl.	Sp. gr.
0.005036	17.111	1.0000943	8.42	27.10	1.1336
0.01006	17.125	1.0001892	5.784	19.30	1.0936
0.02008	17.148	1.0003775	3.77	12.94	1.0618
0.04990	17.138	1.000935	2.031	7.17	1.0334
0.09885	17.133	1.001843	1.588	5.65	1.0261
0.19641	17.162	1.003633	1.138	4.05	1.0187
0.29247	17.147	1.005382	0.523	1.90	1.0076
0.48278	17.140	1.008811			
0.4994	17.28	1.00908			
4.994	17.35	1.08390			

(Kohlrausch, W. Ann. 1894, 53. 28.)

Sp. gr. of a normal solution of HCl+Aq at 18°/4° = 1.0165. (Loomis, W. Ann. 1896, 60. 550.)

Sp. gr. of HCl+Aq at 19.5°, when p = per cent strength of solution; d = observed density; w = volume conc. in grams per cc. ($\frac{pd}{100} = w$)

p.	d.	w
36.0	1.1818	0.4255
29.97	1.1511	0.3450
24.35	1.1207	0.2729
18.55	1.0910	0.2024
12.22	1.0587	0.1294
9.148	1.0433	0.0954
6.559	1.0305	0.0676
3.540	1.0159	0.0360
5.345	1.0246	0.0548
1.356	1.0051	0.0136

(Barnes, J. Phys. Chem. 1898, 2. 546.)

Sp. gr. of HCl+Aq at 18° 4°.

g. HCl in 100 g. of solution	Sp. gr.
0.12149	0.99928
0.60757	0.99900
0.040609	0.99887
0.030328	0.99881

(Jahn, Z. phys. Ch. 1900, 33. 567.)

Sp. gr. at 20° of HCl+Aq containing M g. mols. HCl per liter.				
M	0.025	0.05	0.075	0.10
Sp. gr.	1.00034	1.00101	1.00135	1.00180
M	0.25	0.50	0.75	1.0
Sp. gr.	1.00425	1.00849	1.01264	1.01749
M	1.5	2.0		
Sp. gr.	1.02542	1.03414		

(Jones and Pearce, Am. Ch. J. 1907, 33. 730.)

HCl is not absorbed by conc. H₂SO₄+Aq, but in large amounts by anhydrous H₂SO₄ (Aimé.)

Absorption of HCl by H ₂ SO ₄ +Aq.				
Temp. = 17°.				
Sp. gr.	g. per l.		g. per 100 g.	
	HCl	H ₂ SO ₄	HCl	H ₂ SO ₄
1.211	517.8	...	42.7	...
1.220	487.3	22.7	39.9	1.86
1.220	478.8	58.0	39.2	4.75
1.235	455.3	99.3	36.9	8.04
1.260	418.0	161.7	33.2	12.8
1.305	371.4	273.2	28.5	20.9
1.355	306.6	417.7	22.6	30.8
1.430	215.3	638.2	15.0	44.6
1.545	96.7	917.6	6.26	59.4
1.580	51.3	1033.5	3.25	65.4
1.660	10.3	1224.0	0.62	73.7
1.735	1.89	1344.9	0.11	77.5
1.815	1.24	1615.3	0.068	89.0

ption of HCl by H₂SO₄+Aq—Cont.
Temp. = 40°.

g. per l.		g. per 100 g	
HCl	H ₂ SO ₄	HCl	H ₂ SO ₄
421.4	42.2	35.6	3.56
416.4	70.0	34.8	5.86
392.1	107.7	32.4	8.90
346.3	211.2	27.6	16.8
325.4	236.3	25.9	18.8
247.4	383.7	18.5	28.6
161.6	619.4	11.5	44.2
50.9	929.3	3.35	61.1
18.5	1046.0	1.17	66.4
2.9	1207.6	0.17	73.2
1.4	1370.5	0.081	79.4
0.57	1428.4	0.032	81.4
0.52	1478.4	0.029	83.5

Temp. = 70°.

g. per l.		g. per 100 g.	
HCl	H ₂ SO ₄	HCl	H ₂ SO ₄
374.1	18.4	32.7	1.61
357.3	38.9	31.1	3.38
353.8	55.7	30.5	4.80
341.3	93.6	28.9	7.93
277.7	231.9	22.8	18.9
274.3	246.4	22.3	20.0
173.7	476.7	13.2	36.2
96.5	661.8	6.99	48.0
23.6	946.1	1.56	62.7
8.4	1055.0	0.54	67.6
0.86	1371.3	0.05	80.7
0.62	1448.2	0.035	83.0
0.57	1455.2	0.032	83.4

doro, Gazz. ch. it. 1910, 39. II, 626.)

ts. alcohol of 36° B absorb 68 pts. HCl
(Boullay.)

iol of 0.836 sp. gr. dissolves 327 vols.
17.5° and 758 mm. pressure, and the
has sp. gr. = 1.005. (Pierre, A. ch. (3)
)

ty of HCl in methyl alcohol (absolute)
at t°.

% HCl	t°	% HCl
54.6	18	46.9
51.3	31.7	43

(de Bruyn, R. t. c. 11. 112.)

ty of HCl in ethyl alcohol (absolute)
at t°.

% HCl	t°	% HCl
45.4	19.2	41
44.2	23.5	40.2
42.7	32.0	38.1

(de Bruyn, l.c.)

Solubility of HCl in ether at t° and 760 mm.
pressure.

t°	% HCl	t°	% HCl
—9.2	37.51	15	27.62
—5	37.0	20	24.9
0	35.6	25	22.18
+5	33.1	30	19.47
10	30.35

(Schuncke, Z. phys. Ch. 1894, 14. 336.)

Sol. in glacial HC₂H₃O₂, ether, hexane,
benzene, xylene, etc.

Oil of turpentine absorbs 50% HCl.
(Thénard.)

Oil of turpentine absorbs 163 vols. HCl at
22° and 724 mm.; isoterebenthene absorbs
34% at 24° and 724 mm.; metaterebenthene
absorbs 17.7% at 24° and 724 mm. (Berthe-
lot.)

Oil of lavender absorbs 68.7 vols. at 24°.
(Thénard.)

Oil of lavender absorbs 210 vols. without
being saturated; oil of rosemary absorbs 218
vols. at 22°; sol. in 0.4 vol. petroleum.
(Saussure.)

Absorbed by caprylic alcohol. (Bouis.)
Fuming HCl+Aq is sol. in glycerine and
miscible with conc. HC₂H₃O₂.

Solubility of HCl in phenol+Aq at 12°.

Comp. of H ₂ O layer		Ccmp. of phenol layer	
% HCl	% phenol	% HCl	% phenol
0	7.45	0	72
3.1	6.6	0.09	78
6.6	5.3	0.2	80.3
8.0	5.1	0.36	82.6
10.7	4.8	0.52	84.5

Composition of solution in contact with solid phenol.

% H ₂ O	% HCl	% phenol
11.22	0	88.78
14.98	0.52	84.5
84.5	10.7	4.8
80.38	15.64	3.98
72.43	24.37	3.2
60.25	36.25	3.5

(Schreinemakers, Z. phys. Ch. 1912, 79. 553.)

+H₂O. F.-pt.—15.35°.
Very sol. in H₂O but only slightly sol. in
HCl. (Rupert, J. Am. Chem. Soc. 1909, 31.
866.)

+2H₂O. M.-pt.—17.4°.
+3H₂O. M.-pt.—24.8°. (Pickering, B.
1893, 26. 280.)

The composition of the hydrates formed
by HCl at different dilutions is calculated

from determinations of the lowering of the f.-pt. produced by HCl, and of the conductivity and sp. gr. of HCl+Aq. (Jones, Am. Ch. J. 1905, 34. 323.)

Chlorhydric cyanhydric acid, 3HCl, 2HCN.

Decomp. by H_2O or alcohol; sol. in $HC_2H_3O_2$. Insol. in ether, chloroform, or acetic ether. (Claisen, B. 18. 309.)

HCl, HCN. Sol. in H_2O , absolute alcohol, $HC_2H_3O_2$, and $CHCl_3$, with decomp.; decomp. is especially rapid in H_2O . (Gautier, A. ch. (4) 17. 130.)

Chloric acid, $HClO_3$.

Known only in aqueous solution, which can be concentrated in vacuo to a sp. gr. of 1.282 at 14.2° , and then contains 40.10% $HClO_3$, corresponding to $HClO_3 + 7H_2O$; if left longer in vacuo over H_2SO_4 , an acid corresponding to $HClO_3 + 4\frac{1}{2}H_2O$ is obtained. Aqueous solution of HCN, decomp. at 40° . (Kammerer, Pogg. 138. 390.)

Chlorates.

All chlorates except mercurous chlorate are sol. in H_2O ; most of them are deliquescent; many are sol. in alcohol.

Aluminum chlorate, $Al(ClO_3)_3 + 6H_2O$.

Very hygroscopic. (Dobroserdow, G. C. 1904, II. 177.)

+9 H_2O . Very sol. in cold but much less than in hot H_2O . (Dobroserdow.)

Ammonium chlorate, NH_4ClO_3 .

Easily sol. in H_2O ; less sol. in alcohol. Much less sol. in H_2O at 0° than $NaClO_3$. (Storer.)

Very sl. sol. in absolute alcohol. (Wächter, J. pr. 30. 321.)

Barium chlorate, $Ba(ClO_3)_2 + H_2O$.

Sol. in 4 pts. cold, and less hot H_2O . (Chevenix.)

100 pts. H_2O dissolve at:
 0° 20° 40° 60° 80° 100°
 22.8 37.0 52.1 77.5 98.0 126.4 pts. $Ba(ClO_3)_2$.

100 grams sat. $Ba(ClO_3)_2$ +Aq at t° contain grams anhydrous $Ba(ClO_3)_2$.

t°	Grams $Ba(ClO_3)_2$	t°	Grams $Ba(ClO_3)_2$
Eutectic point. -2.749 \pm 0.004	15.28	50	36.69
0	16.90	60	40.05
+10	21.93	70	43.04
20	25.26	80	45.90
25	27.53	90	48.70
30	29.43	99.1	51.17
40	33.16	* 104.6	52.67

* 104.6° is bpt. at 740 mm. pressure = 105.0° at 760 mm. pressure.

(Anschütz, Z. phys. Ch. 1906, 56. 238.)

Solubility of $Ba(ClO_3)_2$ in H_2O .

t°	g. $Ba(ClO_3)_2$ in 100 g. H_2O	Sp. gr.
0	25.5	1.19
10	39.3	1.27
40	74.1	1.35
60	92.1	1.43
80	113.2	1.50
100	120.	1.58
105.6 *		1.70

* Bpt. of sat. solution.

(Carlson, Dissert. 1910.)

Only slight traces dissolve in absolute alcohol. (Wächter, J. pr. 30. 334.)

Sol. in acetone. (Eidmann, C. C. 18. 1014.)

Difficultly sol. in acetone. (Nauman 1904, 37. 4328.)

Insol. in methyl acetate. (Nauman 1909, 42. 3790); ethylacetate (Nauman B. 1910, 43. 314.)

Mercurous chlorate.

Known only in solution, which decomposes on evaporation.

Cadmium chlorate, $Cd(ClO_3)_2 + 2H_2O$.

Very deliquescent; sol. in H_2O and alcohol. Melts in crystal H_2O at 80° . (Wächter, 30. 321.)

Solubility in H_2O .

Sat. solution contains at:

-20°	-15°	0°	+18°
72.18	72.53	74.05	76.36
49°	65°		
80.08	82.95% $Cd(ClO_3)_2$.		

Sp. gr. of solution containing 7% $Cd(ClO_3)_2$ at 18° = 2.284. (Meusser, B. 35. 1422.)

Sol. in acetone. (Naumann, B. 1904 4328.)

Cadmium chlorate ammonia, $Cd(ClO_3)_2 + 6NH_3$.

Ppt. (Ephraim, B. 1915, 48. 49.)

Cesium chlorate, $CsClO_3$.

100 g. H_2O dissolve at:

0°	8°	19.8°	30°	42°
2.46	3.50	6.28	9.53	14
50°	77°	99°		
19.40	41.65	76.5 g. $CsClO_3$		

(Calzolari, Acc. Sc. med. di Ferrara, 85. 150.)

Calcium chlorate, $Ca(ClO_3)_2 + 2H_2O$.

Deliquescent; very sol. in H_2O and alcohol. (Wächter, J. pr. 30. 323.)

its water of crystallization at over
f solution sat. at $18^{\circ} = 1.729$, con-
% $\text{Ca}(\text{ClO}_3)_2$. (Mylius B. 1897,

etone. (Eidmann, C. C. 1899, II.
nann, B. 1904, 37. 4328.)

chlorate.

sol. in H_2O . (Prudhomme, C. C.
8.)

chlorate, $\text{Co}(\text{ClO}_3)_2 + 2\text{H}_2\text{O}$.

, B. 1902, 35. 1418.)

Solubility in H_2O .

tion contains at:

	35°	47°	61°
g	67.09	69.66	76.12%

$\text{Co}(\text{ClO}_3)_2$.

of solution containing 64.19%
at $18^{\circ} = 1.861$. (Meusser, B. 1902,

Very deliquescent. Sol. in H_2O
l. Melts in crystal H_2O at 50° .
J. pr. 30. 321.)

y in H_2O .

tion contains at:

	19°	0°	+10.5°
.61	57.45	61.83%	

$\text{Co}(\text{ClO}_3)_2$.
eusser, B. 1902, 35. 1418.)

rate, basic, $\text{Cu}(\text{ClO}_3)_2, 3\text{Cu}(\text{OH})_2$.
 H_2O . Very sol. in dil. acids. Sol.
nc. $\text{Cu}(\text{ClO}_3)_2 + \text{Aq}$, the solubility
with the conc. and temp. (Bour-
Soc. 1898, (3) 19. 950.)

rate, $\text{Cu}(\text{ClO}_3)_2 + 4\text{H}_2\text{O}$.

y in H_2O .

tion contains at:

	-21°	+0.8°	18°	45°
7.12	58.51	62.17	66.17	

1°
6.9% $\text{Cu}(\text{ClO}_3)_2$.

f the solution containing 62.17%
at $18^{\circ} = 1.695$. (Meusser, B. 1902,

Very deliquescent. Easily sol.
alcohol. Melts in its crystal H_2O
ächter, J. pr. 30. 321.)

$\text{Cu}(\text{ClO}_3)_2 + \text{Aq}$ at 15° .

) ₂	2.106	4.778	6.945
	1.01620	1.03857	1.05714

) ₂	10.016	14.387
	1.0844	1.12531

aube, Gm.-K. 5. 1, 921.)

etone. (Naumann, B. 1904, 37.

Cupric chlorate ammonia, $\text{Cu}(\text{ClO}_3)_2, 4\text{NH}_3$.

Ppt. Not hygroscopic. Insol. in alcohol.

$\text{Cu}(\text{ClO}_3)_2, 6\text{NH}_3$. Not hygroscopic. (Eph-
raim, B. 1915, 48. 46.)

Erbium chlorate, $\text{Er}(\text{ClO}_3)_3 + 8\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O and alcohol.

Glucinum chlorate.

Known only in aqueous solution, which de-
composes on evaporation.

Ferrous chlorate.

Known only in solution.

Ferric chlorate, $\text{Fe}(\text{ClO}_3)_3$.

Sol. in H_2O .

Basic salt. Insol. in H_2O .

Lanthanum chlorate, $\text{La}(\text{ClO}_3)_3$.

Deliquescent. (Cleve.)

Lead chlorate, $\text{Pb}(\text{ClO}_3)_2 + \text{H}_2\text{O}$.

Deliquescent; easily sol. in H_2O and alcohol.
(Wächter, J. pr. 30. 321.)

Sp. gr. of solution sat. at $18^{\circ} = 1.947$ and
contains 60.2% $\text{Pb}(\text{ClO}_3)_2$. (Mylius, B.
1897, 30. 1718.)

100 g. H_2O dissolve 440 g. $\text{Pb}(\text{ClO}_3)_2$ at
 18° ; sp. gr. of sat. solution = 1.63. (Carlson,
Dissert. 1910.)

Lithium chlorate, $\text{LiClO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Very deliquescent and sol. in H_2O . Very
easily sol. in alcohol. Melts at 50° in its
crystal water. (Wächter, J. pr. 30. 321.)

$\text{LiClO}_3 + \text{Aq}$ sat. at 18° contains 75.8%
 LiClO_3 . Sp. gr. = 1.815. (Mylius, B. 1897,
30. 1718.)

483 g. LiClO_3 dissolve in 100 g. H_2O at 15° ;
sp. gr. of solution = 1.82. (Carlson, Dissert.
1910.)

Contains $3\text{H}_2\text{O}$, and is not deliquescent.
(Lagorio, Zeit. f. Kryst. 15. 80.)

Salt is anhydrous. (Retgers, Z. phys. Ch.
5. 449.)

Magnesium chlorate, $\text{Mg}(\text{ClO}_3)_2$.

128.1 g. $\text{Mg}(\text{ClO}_3)_2$ dissolve in 100 g. H_2O
at 19° ; sp. gr. of solution = 1.59. (Carlson,
Dissert. 1910.)

Sp. gr. of solution containing 56.5%
 $\text{Mg}(\text{ClO}_3)_2$ at $18^{\circ} = 1.564$. (Meusser, l.c.)

Sp. gr. of solution sat. at $18^{\circ} = 1.594$, con-
taining 56.3% $\text{Mg}(\text{ClO}_3)_2$. (Mylius, B. 1897,
30. 1718.)

Sol. in acetone. (Naumann, B. 1904, 37.
4328.)

Sol. in acetone. (Eidmann, C. C. 1899,
II. 1014.)

+ $2\text{H}_2\text{O}$. Solubility in H_2O .

Sat. solution contains at:

	39.5°	61°	68°	93°
65.37	69.46	70.69	73.71%	

$\text{Mg}(\text{ClO}_3)_2$.
(Meusser, B. 1902, 35. 1416.)

+4H₂O. Solubility in H₂O.
Sat. solution contains at:
42° 65.5°
63.82 69.12% Mg(ClO₃)₂.
(Meusser, l.c.)

+6H₂O. Very deliquescent and sol. in H₂O. Very easily sol. in alcohol. Melts at 40° in its crystal water. (Wächter, J. pr. 30. 325.)
Solubility in H₂O.
Sat. solution contains at:
-18° 0° +18° 29° 35°
51.64 53.27 56.50 60.23 63.65% Mg(ClO₃)₂.
(Meusser)

Manganous chlorate, Mn(ClO₃)₂.
Known only in solution which decomposes on evaporation. (Wächter.)

Mercurous chlorate, Hg₂(ClO₃)₂.
a. Easily sol. in alcohol and H₂O. (Wächter, J. pr. 30. 321.)
β. Insol. in H₂O; easily sol. in HC₂H₃O₂ + Aq. (Wächter.) Decomp. by boiling H₂O.

Mercuric chlorate, 2HgO, Cl₂O₅ + H₂O.
Deliquescent. Decomp. by H₂O into oxide and an acid salt. (Wächter.)
Sol. in 4 pts. cold H₂O. (Chevenix, 1802.)

Nickel chlorate, Ni(ClO₃)₂ + 4H₂O.
Solubility in H₂O.
Sat. solution contains at:
48.5° 55° 65° 79.5°
67.60 68.78 69.05 75.50% Ni(ClO₃)₂.
(Meusser, B. 1902, 35. 1419.)

+6H₂O. Deliquescent. Easily sol. in H₂O and alcohol. Melts in crystal H₂O at 80°. (Wächter, J. pr. 30. 321.)
Solubility in H₂O.
Sat. solution contains at:
-18° -8° 0° +18° 40°
49.55 51.52 52.66 56.74 64.47% Ni(ClO₃)₂.
Sp. gr. of solution containing 56.74% Ni(ClO₃)₂ at 18° = 1.661.
Goes over into 4H₂O salt at 39°. (Meusser.)
156 g. Ni(ClO₃)₂ dissolve in 100 g. H₂O at 16°; sp. gr. of solution = 1.76. (Carlson, Dissert. 1910.)

Nickel chlorate ammonia, Ni(ClO₃)₂, 6NH₃.
Ppt. (Ephraim, B. 1915, 48. 47.)

Potassium chlorate, KClO₃.
Sol. in H₂O with absorption of heat.
Sol. in about 16 pts. cold, and in much less hot H₂O. (Chevenix, 1802.)
Sol. in 30.03 pts. H₂O at 0°; 17.85 pts. at 13.3°; and in 1.66 pts. at 104.78°. (M. R. and P.)

Sol. in 16 pts. H₂O at 18.75°. (Abl.)
100 pts. H₂O at 15.5° dissolve 6.2 pts.; at 100°, 48 pts. (Ure's Dict.)

100 pts. H₂O dissolve pts. KClO₃ at t° -

t°	Pts. KClO ₃	28	35	40	47	65
Pts. KClO ₃ :	:	9.5	12.3	14.4	18.3	29.1

(Gerardin.)

100 pts. H₂O dissolve pts. KClO₃ at t°.

t°	Pts. KClO ₃	t°	Pts. KClO ₃
0	3.33	35.0	12.05
13.32	5.00	49.08	18.96
15.37	6.03	74.89	35.40
24.43	8.44	104.78	60.24

(Gay-Lussac, A. ch. 11. 314.)

100 pts. H₂O dissolve pts. KClO₃ at t°.

t°	Pts. KClO ₃	t°	Pts. KClO ₃
0	3.3	130	88.5
100	56.5	180	190

(Tilden and Shenstone, Roy. Soc. Proc. 3. 345.)

100 pts. H₂O dissolve pts. KClO₃ at t°.

t°	Pts. KClO ₃	t°	Pts. KClO ₃
120	73.7	160	148
136	98.9	190	183

(Tilden and Shenstone, Phil. Trans. 1884. 22.)

Coefficient of solubility is 3.2 + 0.109t + 0.0043t² between 0° and 35°. (Blarez, C. R. 112. 1213.)

Sat. KClO₃ + Aq contains % KClO₃ at t°.

t°	% KClO ₃	t°	% KClO ₃
-0.5	2.6	92	31.2
-0.3	2.4	106	37.2
+4.5	3.5	130	47.0
4.5	2.9	171	59.8
11	4.7	180	62.1
19	6.1	190	63.1
29	8.9	200	64.2
36	9.9	207	66.0
42	11.4	300	87.0
56	15.1	330	96.7
58	16.6		

(Étard, A. ch. 1894, (7) 2. 528.)

Solubility in H_2O .

% $KClO_3$ in a sat. sol.	Pts. sol. in 100 pts. H_2O	Pts. H_2O to 1 pt. $KClO_3$
3.06	3.14	31.8
3.67	3.82	26.2
4.27	4.45	22.5
5.11	5.35	18.5
6.76	7.22	13.6
7.56	8.17	12.2
8.46	9.26	10.8
10.29	11.47	8.7
11.75	13.31	7.5
13.16	14.97	6.6
15.18	17.95	5.6
16.85	20.27	4.9
18.97	23.42	4.2
20.32	25.50	3.9
22.55	29.16	3.4
24.82	32.99	3.0
26.97	36.93	2.6
29.25	41.35	2.4
31.36	46.11	2.1
33.76	51.39	1.9
35.83	55.54	1.8

Pawlewski, B. 1899, 32. 1041.)

 $KClO_3$ + Aq at 25° contains 675 milli- $KClO_3$. (Calvert, Z. phys. Ch. 1901,) H_2O dissolve at:

0°	20°	40°	60°
3.3	7.4	13.8	24.0 g. $KClO_3$
1.021	1.045	1.073	1.115

80°	100°	104°
37.7	56.5	59.9 g. $KClO_3$
1.165	1.219	1.230

of sat. solution.

(Carlson, Dissert. 1910.)

 H_2O dissolve at:

19.8°	30°	99°
7.15	10.27	57.3 g. $KClO_3$

Isolari, Acc. Sc. med. di Ferrara, 1911, 85. 150.)

 $KClO_3$ + Aq contains at:

68°	81°	$86^\circ (?)$
23.25	23.53	30.46% $KClO_3$

Shugaef, Z. anorg. 1914, 66. 161.)

 $KClO_3$ + Aq, according to Kremer's experiments (Pogg. 96. 62), and Gerlach's relations. (Z. anal. 8. 290.)

Sp. gr.	% $KClO_3$	Sp. gr.
1.007	6	1.039
1.014	7	1.045
1.020	8	1.052
1.026	9	1.059
1.033	10	1.066

Sp. gr. of $KClO_3$ + Aq at 20° containing 1 mol. $KClO_3$ to 100 mols. H_2O = 1.04122. (Nicol, Phil. Mag. (5) 16. 122.)Sp. gr. of $KClO_3$ + Aq at 15° containing 5% $KClO_3$ = 1.0316. (Kohlrausch, W. Ann. 1879. 1.)B.-pt. of $KClO_3$ + Aq containing pts. $KClO_3$ to 100 pts. H_2O .

Pts. $KClO_3$	B.-pt.	Pts. $KClO_3$	B.-pt.
6.5	100.5°	44.6	103.0°
13.2	101.0	53.4	103.5
20.2	101.5	62.2	104.0
27.8	102.0	69.2	104.4
35.8	102.5

(Gerlach, Z. anal. 26. 450.)

Saturated solution boils at 105° . (Kremer.)Saturated solution boils at 104.2° , and contains 61.5 pts. $KClO_3$ to 100 pts. H_2O . (Legrand.)Saturated solution boils at 103.3° , and contains 66.6 pts. $KClO_3$ to 100 pts. H_2O . (Griffiths.)Saturated solution boils at 104.4° . (Gerlach, Z. anal. 26. 427.)Sol. in pure HNO_3 without decomp., but decomp. at once by HNO_3 containing NO_2 . (Millon, A. ch. (3) 6. 92.)Sol. in sat. NH_4Cl + Aq without causing pptn.1 mol. (=129 pts.) $KClO_3$ dissolves in 2493 vols. H_2O ; in 2208 vols. H_2O when 1 mol. (=59 pts.) $NaCl$ is added; in 2060 vols. H_2O with 2 mols. (=118 pts.) $NaCl$; and in 1910 vols. H_2O with 4 mols. (=236 pts.) $NaCl$. (Gladstone, Chem. Soc. 15. 302.) $KClO_3$ is sol. in about—29.50 pts. H_2O .35.50 pts. NH_4OH + Aq conc.39.00 pts. dil. NH_4OH + Aq (1 vol. conc.: 3 vols. H_2O).30.50 pts. HNO_3 + Aq (1 vol. conc. HNO_3 : 5 vols. H_2O).33.0 pts. HCl + Aq (1 vol. conc. HCl : 4 vols. H_2O).48.00 pts. $HC_2H_3O_2$ + Aq (1 vol. commercial $HC_2H_3O_2$: 1 vol. H_2O).31.50 pts. NH_4Cl + Aq (1 pt. NH_4Cl : 10 pts. H_2O).18.00 pts. NH_4NO_3 + Aq (1 pt. NH_4NO_3 : 10 pts. H_2O).34.00 pts. $NH_4C_2H_3O_2$ + Aq (dil. NH_4OH + Aq + dil. $HC_2H_3O_2$ + Aq).32.50 pts. $NaC_2H_3O_2$ + Aq (commercial $HC_2H_3O_2$ + Na_2CO_3 , diluted with 4 vols. H_2O).31.50 pts. $Cu(C_2H_3O_2)_2$ + Aq. (See Stolba, Z. anal. 2. 390.)33.50 pts. cane-sugar (1 pt. cane-sugar: 10 pts. H_2O).36.50 pts. grape-sugar (1 pt. grape-sugar: 10 pts. H_2O). (Pearson, Zeit. Chem. 1869. 662.)Addition of K salts to sat. $KClO_3$ + Aq ppts. $KClO_3$ in such a way, that the sum of the

KClO₃ remaining in solution and the K in the salt added, is a constant, which constant is equal to the solubility of KClO₃, so that the following formula represents the coefficient of solubility of KClO₃ after addition of a K salt, $3.2+0.109t+0.0043t^2 - K$ of salt added. (Blarez, C. R. 112. 1213.)

Solubility of KClO₃+TiClO₃.

100 g. H ₂ O dissolve g. salts		
t°	g. TiClO ₃	g. KClO ₃
0	2.8	3.3
15	10	1.5
50	12.67	16.2
100	57.3	48.2

(Rabe, Z. anorg. 1902, 31. 156.)

Solubility of KClO₃ in KNO₃+Aq.

t°	g. per l.	
	KNO ₃	KClO ₃
19.85	0.00	69.88
	12.65	64.86
	25.29	60.33
	101.19	45.85
	202.38	40.20
23.87	0.00	79.09
	50.59	63.14

(Arrhenius, Z. phys. Ch. 1893, 11. 397.)

Solubility in KCl+Aq at 20° C.

G. KCl in 1 litre	G. KClO ₃ in 1 litre	Sp. gr.
0	71.1	1.050
10	58	1.050
20	49	1.050
30	43	1.050
40	39.5	1.054
50	36.5	1.058
60	34	1.064
70	32	1.070
80	30	1.075
90	28	1.081
100	27	1.086
110	25.5	1.091
120	24.5	1.098
130	23.5	1.103
140	22.5	1.108
150	21.5	1.113
160	21.0	1.119
170	20.5	1.124
180	20.0	1.130
190	20.0	1.135
200	20	1.140
210	20	1.145
220	20	1.150
230	20	1.156
240	20	1.161
250	20	1.168

(Winteler, Z. Elektrochem. 1900, 7. 361.)

Solubility in KOH+Aq at 25°.

KOH+Aq	Millimols KClO ₃ per litre of the solution
1/5-normal	624
1/4-normal	573

(Calvert, Z. phys. Ch. 1901, 33. 541.)

Solubility in H₂O₂ at 25°.

Concentration of H ₂ O ₂ millimols per litre	Millimols KClO ₃ per litre of the solution
1260	730
1310	737

(Calvert, l.c.)

Solubility in 1/4 normal KOH+Aq in presence of H₂O₂ at 25°.

Concentration of H ₂ O ₂ millimols per litre	Millimols KClO ₃ per litre of the solution
15	578
276	584
954	616
1073	673

(Calvert, l.c.)

Moderately sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 828.)

Neither dissolved nor attacked by liquid NO₂. (Frankland, Chem. Soc. 1901, 79. 1361.)

Sol. in 120 pts. alcohol of 83% at 16°. (Wittstein.)

Sol. in 120 pts. alcohol of 77.1%. (Pohl, W. A. B. 6. 595.)

Insol. in absolute alcohol. (Gerardin.)

Solubility of KClO₃ in dil. alcohol. D=sp. gr. of alcohol; S=solubility in 100 pts. alcohol at t°.

D = 0.9904		D = 0.9848		D = 0.9793	
t°	S	t°	S	t°	S
13	4.9	14	4.7	14	3.2
21	6.3	26	7.1	26	5.4
25	7.5	39	9.3	38	7.9
30	9.1	47	12.8	46	10.6
35	10.2	55	16.1	51	12.2
44	13.6	65	22.3	63	17.5
50	16.2	66	22.5	65	19.0

D = 0.9726		D = 0.9573		D = 0.9390	
t°	S	t°	S	t°	S
13	2.2	13	1.9	14.5	1.1
20	3.3	20	2.7	28	2.2
33	5.8	29	3.6	40	3.4
43	7.2	36	4.3	50	4.3
56	11.4	55	7.9	62	6.6
59	12.9	60	9.7	67	7.6
..	63	10.5

Solubility of KClO_3 in dil. alcohol—*Continued.*

D = 0.9111		D = 0.8967		D = 0.8429	
t°	S	t°	S	t°	S
13	0.74	12	6.46	25	0.09
25	1.08	31	1.28	34	0.12
32	1.78	43	1.95	56	0.24
52	3.35	58	3.10	64	0.32

(Gerardin, A. ch. (4) 5. 148.)

Solubility of KClO_3 in alcohol + Aq.

t. % alcohol	g. KClO_3 per 100 g. solution	
	t = 30°	t = 40°
0	9.23	12.23
5	7.72	10.48
10	6.44	8.84
20	4.51	6.40
30	3.21	4.67
40	2.35	3.41
50	1.64	2.41
60	1.01	1.41
70	0.54	0.78
80	0.24	0.34
90	0.06	0.12

(Taylor, J. phys. Ch. 1897, 1. 301.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Very sl. sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Solubility of KClO_3 in acetone + Aq.

wt. % acetone	g. KClO_3 per 100 g. solution	
	t = 30°	t = 40°
0	9.23	12.23
5	8.32	11.10
9.09	7.63	10.28
20	6.09	8.27
30	4.93	6.69
40	3.90	5.36
50	2.90	4.03
60	2.03	2.86
70	1.24	1.68
80	0.57	0.79
90	0.18	0.24

(Taylor, J. phys. Ch. 1897, 1. 301.)

Insol. in methyl acetate. (Naumann, B. 909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Solubility in glycol = 0.9% at ord. temp. de Coninck, Belg. Acad. Bull. 1905. 359.)

100 g. glycerol (sp. gr. 1.256) dissolve 3.54 g. KClO_3 at 15–16°. (Ossendowski, Pharm. J. 907, 79. 575.)Potassium silver chlorate, KClO_3 , AgClO_3 .
(Pfaundler, W. A. B. 46, 2. 266.)Rubidium chlorate, RbClO_3 .100 pts. H_2O dissolve 2.8 pts. at 4.7°; 3.9 pts. at 13°; 4.9 pts. at 18.2°; 5.1 pts. at 19°. (Reissig, A. 127. 33.)100 g. H_2O dissolve 3.1 RbClO_3 at 15°; sp. gr. of solution = 1.07. (Carlson, Dissert. 1910.)100 g. H_2O dissolve at:

0°	8°	19.8°	30°
2.138	3.07	5.36	8.00 g. RbClO_3 ,

42.2°	50°	76°	99°
12.48	15.98	34.12	62.8 g. RbClO_3 .

(Calzolari, Acc. Sc. med. di Ferrara, 1911, 85. 150.)

Scandium chlorate.

(Crookes, Roy. Soc. Proc. 1908, 30. A, 518.)

Silver chlorate, AgClO_3 .Sol. in 10–12 pts. cold H_2O (Vauquelin); in 8–10 pts. cold, and 2 pts. hot H_2O (Chevenix); in 5 pts. cold H_2O (Wächter). Sl. sol. in alcohol (Chevenix); easily sol. in alcohol (Wächter).Silver chlorate ammonia, AgClO_3 , 2NH_3 .Easily sol. in H_2O or alcohol. (Wächter, 1843.)Sodium chlorate, NaClO_3 .

Deliquescent.

Sol. in 3 pts. cold and less hot H_2O . (Wächter; Chevenix.)Sol. in 3 pts. H_2O at 18.75°. (Abl.)100 pts. H_2O dissolve 35.5 pts. NaClO_3 . (Ure's Dict.)100 pts. H_2O dissolve at:

0°	20°	40°	60°
81.9	99	123.5	147.1 pts. NaClO_3 ,

80°	100°	120°
175.6	232.6	333.3 pts. NaClO_3 .

(Kremers, Pogg. 97. 4.)

100 pts. H_2O dissolve 89.3 pts. NaClO_3 at 12°. (Schlösing.)100 g. H_2O dissolve at:

–15°	0°	20°	40°
72	79	101	126 pts. NaClO_3 ,

Sp. gr. 1.380 1.389 1.430 1.472

60°	80°	100°	122° *
155	189	230	286 pts. NaClO_3 .

Sp. gr. 1.514 1.559 1.604 1.654

* Bpt. of sat. solution.

(Carlson, Dissert. 1910.)

100 g. NaClO₃+Aq contain at:
4.78° 19.85° 30.05° 35.10° 44.72°
45.47 48.91 51.22 52.36 54.50 g. NaClO₃.
(Le Blanc and Schmandt, Z. phys. Ch. 1911, 77. 614.)

Sp. gr. of NaClO₃+Aq, containing:
10 15 20 25 30 25% NaClO₃.
1.070 1.108 1.147 1.190 1.235 1.282
(Gerlach, Z. anal. 8. 290.)

Sp. gr. of NaClO₃+Aq at 20° containing 1 mol. NaClO₃ in 100 mols. H₂O = 1.03844. (Nicol, Phil. Mag. (5) 16. 122.)
NaClO₃+Aq containing 7.23% NaClO₃ has sp. gr. 20°/20° = 1.0496. (Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 278.)
Sat. solution boils at 132°, and temp. can be raised to 135° by supersaturation. (Kremers, Pogg. 97. 4.)
Easily sol. in liquid HF. (Franklin, Z. anorg. 1905, 46. 2.)
NaClO₃+NaCl.
100 pts. H₂O dissolve 50.75 pts. NaClO₃+24.4 pts. NaCl at 12°; 100 pts. H₂O dissolve 249.6 pts. NaClO₃+11.5 pts. NaCl at 122°, and when cooled to 12° contain 68.6 pts. NaClO₃+11.5 pts. NaCl. (Schlösing, C. R. 73. 1272.)

Solubility in NaCl+Aq at 20° C.		
G. NaCl in 1 litre	G. NaClO ₃ in 1 litre	Sp. gr.
5	668	1.426
10	661	1.424
15	653	1.423
20	645	1.421
25	638	1.419
30	630	1.418
35	622	1.417
40	615	1.415
45	607	1.414
50	599	1.412
55	590	1.411
60	582	1.409
65	574	1.408
70	566	1.406
75	559	1.405
80	551	1.404
85	544	1.402
90	537	1.401
95	529	1.399
100	522	1.398
105	514	1.396
110	507	1.394
115	499	1.392
120	491	1.391
125	484	1.389
130	476	1.387
135	467	1.385
140	459	1.383
145	451	1.381

Solubility in NaCl+Aq at 20° C.—Continued		
G. NaCl in 1 litre	G. NaClO ₃ in 1 litre	Sp. gr.
150	442	1.379
155	432	1.377
160	423	1.374
165	414	1.372
170	403	1.369
175	393	1.365
180	382	1.362
185	371	1.359
190	360	1.355
195	349	1.350
200	338	1.345
205	326	1.340
210	315	1.335
215	302	1.330
220	287	1.324
225	271	1.319
230	257	1.313
235	243	1.307
240	228	1.301
245	211	1.295
250	197	1.289
255	184	1.283
260	170	1.276
265	150	1.270
270	135	1.263
275	120	1.256
280	105	1.249
285	91	1.241
290	78	1.235
295	67	1.226
300	55	1.217

(Winteler, Z. Elektrochem. 1900, 7. 361.)
Very sol. in liquid NH₃. (Franklin, Am Ch. J. 1898, 20. 829.)
Sol. in 34 pts. alcohol of 83% at 16° and in less hot alcohol. (Wittstein.)
Somewhat more easily sol. in alcohol than NaCl. (Berzelius.)

Solubility of NaClO ₃ in alcohol. (g. NaClO ₃ per l. of solution.)			
t°	Alcohol		
	90 %	75 %	50 %
20	16.1	110.8	311.3
40	22.9	133.5	321.8
60	29.0	155.8	326.8
70	161.3

(Carlson, Dissert. 1910.)
Insol. in methyl acetate. (Naumann, E 1909, 42. 3790); ethyl acetate. (Naumann B. 1910, 48. 314.)
100 g. glycerol dissolve 20 g. NaClO₃ at 15.5°. (Ossendowski, Pharm. J. 1907, 71 575.)

m chlorate, $\text{Sr}(\text{ClO}_3)_2 + 5\text{H}_2\text{O}$.

deliquescent, and sol. in H_2O . (Top-A. B. 66, 2. 29.)

r. of solution sat. at 18° containing $\text{Sr}(\text{ClO}_3)_2 = 1.839$. (Mylius, B. 1897, 30.)

sol. in H_2O , less in alcohol, but more in alcohol than SrCl_2 . (Souchay, A. 102.)

in absolute alcohol. (Wächter.)

chlorate, TlClO_3 .

in H_2O , but decomp. by heating.

in H_2O dissolve at:

	20°	50°	80°	100°
pts. TlClO_3	1.92	12.67	36.65	57.31

(Muir, Chem. Soc. 29. 857.)

$\text{ClO}_3 + \text{Aq}$ sat. at 10° contains 25.637 g. (Roozeboom, Z. phys. Ch. 8. 532.)

H_2O dissolves 0.134 equivalents at 20° ; or 38.51 g. in 1 l. of the solution (mean of 10 experiments). (Noyes and J. Am. Chem. Soc. 1911, 33. 1657.)

Solubility in $\text{Ti}_2\text{SO}_4 + \text{Aq}$ at 20° .

quiv. per l.		Solid phase
	Ti_2SO_4	
3	0.1366	$\text{TlClO}_3 + \text{Ti}_2\text{SO}_4$

(Noyes and Farrell, *l.c.*)

chlorate, $\text{Tl}(\text{ClO}_3)_3 + 4\text{H}_2\text{O}$.

deliquescent; sol. in H_2O . Decomp. in the air. (Gewecke, Z. anorg. 1912, 30.)

m chlorate.

in H_2O . (Popp, A. 131. 179.)

chlorate, $\text{Y}(\text{ClO}_3)_3 + 8\text{H}_2\text{O}$.

deliquescent. Easily sol. in alcohol. Sl. in ether. (Cleve.)

chlorate, $\text{Zn}(\text{ClO}_3)_2 + 4\text{H}_2\text{O}$.

Solubility in H_2O .

Solution contains at:

	30°	40°	55°
% $\text{Zn}(\text{ClO}_3)_2$	67.66	69.06	75.44

Sp. gr. of solution containing 66.52% $\text{Zn}(\text{ClO}_3)_2$ at $18^\circ = 1.916$. (Meusser, B. 1902, 35. 1417.)

sol. in H_2O than chlorates of Mg, Co, Ni; less sol. than chlorate of Cd; more than $\text{Zn}(\text{NO}_3)_2$. (Meusser, *l.c.*)

H_2O . Very deliquescent. Easily sol. in alcohol. Melts in crystal H_2O at 60° . (Lin, A. ch. 95. 113.)

Solubility in H_2O .

Sat. solution contains at:

	-18°	0°	8°	15°
% $\text{Zn}(\text{ClO}_3)_2$	55.62	59.19	60.20	67.32

(Meusser, *l.c.*)

Sp. gr. of solution sat. at 18° containing 65% $\text{Zn}(\text{ClO}_3)_2 = 1.914$. (Mylius, B. 1897, 30. 1718.)

Zinc chlorate ammonia, $\text{Zr}(\text{ClO}_3)_2 \cdot 4\text{NH}_3$.

$\text{Zn}(\text{ClO}_3)_2 \cdot 6\text{NH}_3$. Ppt. (Ephraim, B. 1915, 48. 48.)

Perchloric acid

See Perchloric acid.

Chlorides.

Most chlorides are sol. in H_2O ; a few, however, are insol. or nearly so therein, the chief of which are AgCl , Hg_2Cl_2 , Cu_2Cl_2 , PtCl_2 , and AuCl . Several chlorides are decomp. into insol. basic salts or hydroxides, either by the addition of H_2O , as in the case of BiCl_3 and SbCl_3 , or on evaporating the aqueous solution, as AlCl_3 , ZnCl_2 , MgCl_2 , etc.

Some chlorides are sol. in alcohol or ether. See under each element.

Chlorine, Cl_2 .

The maximum solubility of Cl in H_2O is at 10° (Schönfeld); at $8-10^\circ$ (Gay-Lussac); at $9-10^\circ$ (Pelouze).

Solubility decreases from $9-0^\circ$; at 100° the solubility = 0. (Gay-Lussac.)

$\text{Cl}_2 + \text{Aq}$ sat. at 6° has sp. gr. = 1.003. (Berthelot.)

1 vol. H_2O at t° absorbs vols. Cl reduced to 0° and 760 mm. pressure.

t°	Vols. Cl	t°	Vols. Cl
10	2.5852	26	1.9099
11	2.5413	27	1.8695
12	2.4977	28	1.8295
13	2.4543	29	1.7895
14	2.4111	30	1.7499
15	2.3681	31	1.7104
16	2.3253	32	1.6712
17	2.2828	33	1.6322
18	2.2405	34	1.5934
19	2.1984	35	1.5550
20	2.1565	36	1.5166
21	2.1148	37	1.4785
22	2.0734	38	1.4406
23	2.0322	39	1.4029
24	1.9912	40	1.3655
25	1.9504

(Schönfeld, A. 93. 26.)

1 vol. H₂O absorbs vols. Cl at t° (not corrected).

Vols. Cl	t°	Vols. Cl	t°	Vols. Cl	t°
1.43	0	3.04	8	1.19	50
1.52	3	3.00	10	0.71	70
2.08	6.5	2.37	17	0.15	100
2.17	7	1.61	35

(Gay-Lussac, A. ch. (3) 7. 124.)

1 vol. H₂O at 8° absorbs 3.04 vols. Cl, which is the maximum of solubility. At 50°, 1.09 vols. are absorbed; and at 0°, 1.5 vols. (Pelouze and Fremy.)

1 vol. H₂O at t° dissolves vols. Cl (not corrected).

t°	Vols. Cl	t°	Vols. Cl	t°	Vols. Cl
0	1.75-1.80	12	2.50-2.60	40	1.55-1.60
9	2.70-2.75	14	2.45-2.50	50	1.15-1.20
10	2.70-2.75	30	2.00-2.10	70	0.60-0.65

(Pelouze, A. ch. (3) 7. 188.)

1 vol. H₂O absorbs vols. Cl at t°.

t°	Vols. Cl	t°	Vols. Cl	t°	Vols. Cl
0	1.5-1.6	9	2.65-2.70	14	2.6-2.65
5	2.05-2.1	10	2.9-3.0	16	2.35-2.4
8	2.5-2.6	12	2.65-2.75	30	1.8-1.85

(Riegel and Walz, Berz. J. B. 1846. 72.)

Solubility in H₂O : a = coefficient of solubility.

t°	a	t°	a	t°	a
6.9	2.2931	10.1	2.8741	21.7	2.0422
8.4	2.5469	11.2	2.7267	32.1	1.5766
9.3	2.7135	13.7	2.5079	36.7	1.3802

(Goodwin, B. 15. 3040.)

Goodwin also gives tables for solubility of Cl in HCl and various chlorides, but they do not show evidence of accurate work. (A.M.C.)

Cl₂+Aq contains at 760 mm. pressure:

1.44%	Cl at 0°
1.07%	" " 6°
0.95%	" " 9°
0.87%	" " 12°

(Roozeboom, R. t. c. 1884, 3. 29.)

See also Cl₂+8H₂O.

Solubility of Cl₂ in H₂O.

β¹ = Vol. of Cl (reduced to 0° and 760 mm.) absorbed by 1 vol. H₂O under a total pressure of 760 mm.

q = g. Cl₂ absorbed by 100 g. H₂O under a total pressure of 760 mm.

t°	β ¹	q	t°	β ¹	q
10	3.095	0.980	25	1.985	0.630
11	2.996	948	26	1.937	615
12	2.900	918	27	1.891	600
13	2.808	889	28	1.848	587
14	2.720	861	29	1.808	574
15	2.635	835	30	1.769	562
16	2.553	809	35	1.575	501
17	2.474	784	40	1.414	451
18	2.399	760	45	1.300	415
19	2.328	738	50	1.204	396
20	2.260	716	60	1.006	324
21	2.200	698	70	0.848	274
22	2.143	680	80	0.672	219
23	2.087	662	90	0.380	125
24	2.035	646	100	0.000	000

(Winkler, Landolt and Börnstein, Tab. 4th Ed. 1912, 597.)

1 l. HCl+Aq (38% HCl) dissolves 17.3 g. Cl; 1 l. HCl+Aq (33% HCl) dissolves 11 g. Cl; 1 l. HCl+Aq (3% HCl) dissolves 6.5 g. Cl. (Berthelot, C. R. 91. 191.)

Solubility of Cl₂ in HCl+Aq at 20-21° and 759-761 mm. pressure.

g. HCl per l.	g. Cl ₂ per l.	Coefficient of absorption	Solubility
0	7.23	2.1157	2.2799
3.134	5.30	1.5496	1.6608
6.248	4.94	1.4483	1.5607
9.402	4.76	1.3942	1.5013
12.540	4.85	1.4200	1.5292
15.670	5.10	1.4933	1.6092
31.340	5.81	1.6736	1.8033
62.680	6.38	1.8682	2.0131
94.020	7.19	2.1044	2.2677
125.360	7.76	2.2711	2.4473
156.700	8.58	2.5095	2.7043
188.040	9.23	2.7020	2.9117
219.380	9.93	2.9243	3.1312
250.720	10.68	3.1272	3.3677
282.060	11.87	3.3278	3.5859
313.401	12.03	3.5492	3.8224

(Mellor, Chem. Soc. 1901, 75. 227.)

Solubility of Cl in NaCl+Aq. a = coefficient of solubility.

NaCl = 9.97%.

t°	a	t°	a
7.9	1.8115	18.8	1.2785
11.9	1.5879	22.6	1.0081
15.4	1.3684

Solubility of Cl in NaCl + Aq.—Continued
NaCl = 16.01%.

t°	a	t°	a
6	1.5866	21.4	0.8732
11.6	1.2227	26.9	0.7017
16.4	1.0121

NaCl = 19.66%.

t°	a	t°	a
0	1.6978	15.4	0.9511
9.2	1.2145	20.4	0.7758
9.3	1.2068	21.9	0.7385
14.8	0.9740

(Kumpf, W. Ann. Beibl. 6. 276.)

Solubility of Cl in sat. NaCl + Aq at t° and 760 mm. pressure.

t°	Coefficient of absorption at 0° and 760 mm.	Solubility at 0° and 760 mm.
14.5	0.3607	0.3898
29.0	0.3125	0.3458
60.0	0.1332	0.1625
82.0	0.0586	0.0763

(Kohn and O'Brien, J. Soc. Chem. Ind. 1898, 17. 1100.)

Sat. KCl + Aq absorbs 1/3 less Cl at 15° than pure H₂O. (Dettmer, A. 38. 35.)

1 l. of a solution of CaCl₂ (1 pt. in 15 pts. H₂O) dissolves 2.45 g. Cl at 12°.

1 l. of a solution of MgCl₂ (1 pt. in 15 pts. H₂O) dissolves 2.33 g. Cl at 12°.

1 l. of a solution of MnCl₂ (1 pt. in 15 pts. H₂O) dissolves 2.00 g. Cl at 12°.

Sl. sol. in KOH + Aq. (Fremy.)

Somewhat sol. in liquid NO₂. (Frankland, Chem. Soc. 1901, 79. 1361.)

CCl₄ absorbs 10% of Cl₂ at 13°. (Perkins, Chem. Soc. 1894, 65. 20.)

1 mol. CrOCl₂ dissolves at 0°, 0.70 atom Cl; at -14°, 1.24 atoms; at -21°, 2.31 atoms; and at -24, 3.00 atoms Cl. (Roozeboom, R. t. c. 4. 379.)

Sulphuryl chloride absorbs 71 vols. Cl or 0.136 pt. Cl by weight at 0°. (Schulze, J. pr. (2) 27. 168.)

Insol. in benzene. (Moride.)

Sl. sol. in chloral and iodal. (Dumas.)

Sol. in perchlorethylene. (Faraday.)

Sol. in a very large quantity of ether with decomp.

Coefficient of solubility of Cl₂ in organic liquids at 15°.

Substance	Coefficient of Solubility
Carbon tetrachloride	51.7
Acetic anhydride	39.6
Acetic acid (99.84%)	36.7
" (90 vol. %)	25.3
" (75 vol. %)	16.43
" (65 vol. %)	13.43

(Jones, Chem. Soc. 1911, 99. 392.)

+8H₂O. Critical temp. of decomposition in open vessel = 9.6°; in closed vessel = 28.7°.

Solubility in H₂O.

% Cl₂ = % of Cl₂ in Cl₂ + Aq sat. at t° and 760 mm. in presence of Cl₂ + 8H₂O.

t°	% Cl ₂	t°	% Cl ₂
0	0.505	12.5	1.10
3	0.64	20	1.82
6	0.709	28.5	3.50
9	0.900		

(Roozeboom, R. t. c. 1884, 3. 57.)

Chlorine monoxide, Cl₂O.

Sol. in H₂O. At 0°, H₂O absorbs at least 200 times its volume of Cl₂O gas.

Chlorine trioxide, Cl₂O₃.

Decomp. on air at 57° with explosion.

H₂O absorbs 5-6 vols. Cl₂O₃. (Millon, A. ch. (3) 7. 298.)

H₂C absorbs at 8.5° and 753 mm. press. 8.591 vols. Cl₂O₃. (Brandan.)

100 g. H₂O dissolve at:

8.5°	and	752.9 mm. press.	4.7655 g. Cl ₂ O ₃ .
14°	"	756.3 "	5.0117 "
21°	"	754 "	5.4447 "
93°	"	760 "	5.6508 "

(Brandan, A. 151. 340.)

Does not exist, and above data are for mixture of ClO₂ and Cl. (Garzarolli-Thurnlakh, A. 209. 184.)

Chlorine tetroxide, ClO₂.

H₂O at 4° absorbs about 20 vols. ClO₂ with formation of HClO₂ and HClO₃.

H₂SO₄ at -18° absorbs about 20 vols. ClO₂. (Millon, A. ch. (3) 7. 285.)

Solubility of ClO₂ in H₂O.

t°	g. ClO ₂ per l.
1	> 108.6
10.7	116.7
14.0	> 107.9

ray, Z. phys. Ch. 1906, 54. 569.)

+8H₂O (±1H₂O).

Solubility in H₂O.

t°	g. ClO ₂ per l.	t°	g. ClO ₂ per l.
0.79*	26.98	10	60.06
0	27.59	15.3	60.06
1	29.48	18.2	107.9
5.7	42.10		

* Entertic.
(Bray.)

Chlorine oxide, Cl₂O₁₇.
Very easily decomp. (Millon, A. 46. 281.)
Probably a mixture of ClO₂ and O.

Chlorine heptoxide, Cl₂O₇.
Explosive; decomp. by H₂O; sol. in well cooled benzene with sl. decomp. (Michael, Am. Ch. J. 1909, 23. 447.)

Chlorirididiamine chloride,
Cl₂Ir^{NH₂NH₂Cl}_{NH₂NH₂Cl}
Sl. sol. in cold, easily in hot H₂O. (Skoblikoff, A. 84. 275.)

— nitrate, Cl₂Ir(N₂H₄NO₃)₂.
Sol. in H₂O.

— sulphate, Cl₂Ir(N₂H₄)SO₄.
Sl. sol. in cold, much more easily in hot H₂O.

Chloriridic acid.

Chloriridates.
Most of the chloriridates are very difficultly sol. in H₂O, but a little more sol. than the corresponding chloroplatinates. Insol. or nearly so in alcohol, but not so difficultly sol. as the chloroplatinates. (Rose.)

Ammonium chloriridate, (NH₄)₂IrCl₆.
Sol. in 20 pts. cold H₂O (Vauquelin); sl. sol. in cold, much more in hot H₂O (Claus); sol. in HCl+Aq (Soblewsky); insol. in cold NH₄Cl+Aq (Claus); insol. in alcohol (Berzelius).

100 pts. H₂O dissolve at:
14.4° 26.8° 39.4°
0.699 0.905 1.226 pts. (NH₄)₂IrCl₆,
52.2° 61.2° 69.3°
1.608 2.130 2.824 pts. (NH₄)₂IrCl₆.
(Rimbach and Korten, Z. anorg. 1907, 52. 407.)

Cæsium chloriridate, Cs₂IrCl₆.
Only sl. sol. in H₂O. (Delépine, C. R. 1908, 146. 1268.)

Lithium chloriridate, Li₂IrCl₆.
Somewhat deliquescent; very sol. in H₂O (Antony, Gazz. ch. it. 23, 1. 190.)

Potassium chloriridate, K₂IrCl₆.
Sl. sol. in cold H₂O; sol. in 15 pts. hot H₂O; less sol. in H₂O containing HCl; in alcohol or sat. KCl, and CaCl₂+Aq. Insol. in liquid NH₃. (Gore, Am. Ch. 1898, 20. 829.)

Rubidium chloriridate, Rb₂IrCl₆.
Very sl. sol. in H₂O. (Rimbach, Z. an 1907, 52. 408.)

Sodium chloriridate, Na₂IrCl₆+6H₂O.
Easily sol. in H₂O; sol. in alcohol of 0 sp. gr.

Thallium chloriridate, Tl₂IrCl₆.
Decomp. by hot HCl forming Tl₂Cl₃ (Delépine, C. R. 1909, 149. 1073.)

Chloriridium pentamine comps.
See Iridopentamine chloro comps.

Chloriridosulphurous acid.
Potassium chloriridosulphite, K₄Ir₂Cl₂(SO₃)₄+12H₂O.
Insol. in cold, decomp. by hot H₂O. K₄Ir₂Cl₂(SO₃)₄, 2K₂SO₃. Decomp. by 1 Cl₂Ir₂(SO₃)₂, 8KCl+4H₂O. Sol. in H₂O, insol. in alcohol. (Claus, J. pr. 42. 354.)

Chloriridous acid.
Ammonium chloriridite, (NH₄)₂IrCl₅.
Decomp. by H₂O. (Delépine, C. R. 1 146. 1268.)
+1½H₂O. Sol. in H₂O. (Claus.)
IrCl₅(H₂O)(NH₄)₂. (Delépine.)

Cæsium chloriridite, IrCl₅(H₂O)Cs₂.
(Delépine.)

Lithium chloriridite, Li₂IrCl₅+12H₂O.
Deliquescent; sol. in H₂O and alcoh. (Delépine, C. R. 1914, 158. 1277.)

Lithium sodium chloriridite, Li₂NaIrCl₅ 12H₂O.
Stable in aq. solution in the presence of excess of lithium salt. (Delépine, C. R. 1 158. 1278.)
LiNa₂IrCl₅+12H₂O. Stable in aq. solution in the presence of excess of sodium salt. (Delépine, C. R. 1914, 158. 1278.)

Potassium chloriridite, K₂IrCl₅.
Decomp. by H₂O. (Delépine.)
+3H₂O. Easily sol. in H₂O; insol. in alcohol; insol. in sat. KCl+Aq. (Berzelius) IrCl₅(H₂O)K₂. (Delépine.)

n chloriridite, $\text{IrCl}_3(\text{H}_2\text{O})\text{Rb}_2$.
(ine.)

loriridite, Ag_3IrCl_4 .

in H_2O or acids; sl. sol. in NH_4OH +
(Delépine, Bull. Soc. 1910, (4), 7. 55.)

chloriridite, $\text{Na}_3\text{IrCl}_6 + 12\text{H}_2\text{O}$.

scint; sol. in $\frac{1}{2}$ pt. H_2O . Insol. in
Melts in crystal H_2O at 50° .

chloriridite, Tl_3IrCl_4 .

hot HCl ; pptd. on cooling. (Delé-
R. 1909, 149. 1073.)

tramine chromium comps.

lorotetramine chromium comps.

azoimide, N_3Cl .

. in H_2O . (Raschig, B. 1908, 41.

romo comps:

omochloro comps.

arbonic acid.

rbonyl chloride.

romic acid. CrO_2OH
 Cl .

. only in its salts.

$\frac{1}{2}$. See Chromyl chloride.

am chlorochromate, $\text{NH}_4\text{CrO}_3\text{Cl} =$
 Cl
 ONH_4 .

ol. in H_2O than the K salt. (Peligot,
. 283.)

hlorochromate chloride,
 CrO_3Cl_2 , BaCl_2 .

escent. Very sol. in H_2O . (Prätor-
1. 1.)

. Not deliquescent.

hlorochromate, $\text{Ca}(\text{CrO}_3\text{Cl})_2$.

escent. (Peligot.)

). Very deliquescent. (Prätorius.)

s chlorochromate.

chromyl chloride.

lorochromate, $\text{Co}(\text{CrO}_3\text{Cl})_2 + 9\text{H}_2\text{O}$.
escent; melts at 40° in crystal H_2O .
s.)

hlorochromate, LiCrO_3Cl .

H_2O acidified with HCl without
(Löwenthal, Z. anorg. 1894, 6.

Magnesium chlorochromate, $\text{Mg}(\text{CrO}_3\text{Cl})_2$.

Deliquescent. (Peligot.)

+ $9\text{H}_2\text{O}$. Less deliquescent than the other
chlorochromates. (Prätorius, A. 201. 1.)

Very hygroscopic; sol. in H_2O acidified
with HCl without decomp. (Löwenthal, Z.
anorg. 1894, 6. 359.)

Nickel chlorochromate, $\text{Ni}(\text{CrO}_3\text{Cl})_2 + 9\text{H}_2\text{O}$.

Deliquescent; melts in its crystal H_2O at
 $46-48^\circ$. (Prätorius.)

Potassium chlorochromate, $\text{KCrO}_3\text{Cl} =$
 $\text{CrO}_3(\text{Cl})\text{OK}$.

Sol. in H_2O with decomp. Cryst. from H_2O
containing HCl without decomp. (Peligot.)

Sol. in acetone. (Naumann, B. 1904, 37.
4328.)

Sodium chlorochromate, NaCrO_3Cl .

Deliquescent. (Peligot.)

+ $2\text{H}_2\text{O}$. Deliquescent. (Prätorius.)

Strontium chlorochromate, $\text{Sr}(\text{CrO}_3\text{Cl})_2 +$
 $4\text{H}_2\text{O}$.

Deliquescent; melts in crystal H_2O at 72° .
(Prätorius.)

Thallous chlorochromate, TlCrO_3Cl .

Decomp. by H_2O . (Lachaud and Lepierre,
C. R. 103. 198.)

Zinc chlorochromate, $\text{Zn}(\text{CrO}_3\text{Cl})_2 + 9\text{H}_2\text{O}$.

Deliquescent; melts at 37.5° in crystal H_2O .
(Prätorius.)

Very hygroscopic; very sol. in H_2O and
acids. (Löwenthal, Z. anorg. 1894, 6. 360.)

Dichlorochromium bromide,

$[\text{Cr}(\text{H}_2\text{O})_4\text{Cl}_2]\text{Br}$.

Very deliquescent. Sol. in fuming HBr .
in a mixture of equal volumes ether and fum-
ing HBr , in alcohol and in acetone. (Bjerrum,
B. 1907, 40. 2919.)

Chlorochromotetrammonium comps.

See Chlorotetramine chromium comps.

Chlorocolumbium bromide, $(\text{Cb}_3\text{Cl}_{12})\text{Br}_2 +$
 $7\text{H}_2\text{O}$.

Sol. in a small quantity of cold H_2O .
(Harned, J. Am. Chem. Soc. 1913, 35. 1083.)

Chlorocolumbium chloride, $(\text{Cb}_3\text{Cl}_{12})\text{Cl}_2 +$
 $7\text{H}_2\text{O}$.

Insol. in cold, sol. in boiling H_2O .

Not easily decomp. by boiling with NH_4OH .
Conc. HNO_3 decomp. a boiling solution of
this comp. Completely sol. in conc. alkalis.
(Harned, J. Am. Chem. Soc. 1913, 35. 1080.)

Chlorocolumbium hydroxide, $(\text{Cb}_2\text{Cl}_{12})(\text{OH})_2 + 8\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in acids and alkalies. (Harned, J. Am. Chem. Soc. 1913, **35**. 1082.)

Chloroctamine cobaltic carbonate, $\text{Cl}_4\text{Co}_2(\text{NH}_3)_8\text{CO}_3 + 2\text{H}_2\text{O}$.

Very sol. in H_2O . (Vortmann and Blasberg, B. **22**. 2651.)

$\text{Cl}_2\text{Co}_2(\text{NH}_3)_8(\text{CO}_3)_2 + \text{H}_2\text{O}$. (Vortmann and Blasberg.)

Chloroferrous acid.

Calcium chloroferrite, CaO , CaCl_2 , Fe_2O_3 .

Insol. in H_2O . (le Chatelier, C. R. **99**. 276.)

Dichlorofulminoplatinum,

$\text{Pt}_4\text{N}_4\text{Cl}_2\text{O}_{12}\text{H}_{22}(?)$.

Insol. in H_2O . (v. Meyer, J. pr. (2) **18**. 305.)

Trichlorofulminoplatinum,

$\text{Pt}_4\text{N}_4\text{Cl}_3(\text{OH})\text{O}_{12}\text{H}_{24}(?)$.

Insol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$. (v. Meyer.)

Tetrachlorofulminoplatinum,

$\text{Pt}_4\text{N}_4\text{Cl}_4\text{O}_{12}\text{H}_{24}(?)$

Insol. in H_2O . (v. Meyer.)

Chlorohydroxylonitritoplatin semidiamine nitrite, $(\text{OH})\text{ClNO}_2\text{Pt}(\text{NH}_3)_2\text{NO}_2$.

Easily sol. in hot H_2O . (Cleve.)

Chlorohydroxyloplatin diamine bromide,

$\frac{\text{OH}}{\text{Cl}}\text{Pt}(\text{N}_2\text{H}_5\text{Br})_2$.

Sl. sol. in H_2O .

— **carbonate**, $\frac{\text{OH}}{\text{Cl}}\text{Pt}(\text{N}_2\text{H}_5)_2\text{CO}_3$.

Insol. in H_2O . (Cleve.)

— **chloride**, $\frac{\text{OH}}{\text{Cl}}\text{Pt}(\text{N}_2\text{H}_5\text{Cl})_2$.

Sl. sol. in H_2O . (Cleve.)

— **chromate**, $\frac{\text{OH}}{\text{Cl}}\text{Pt}(\text{N}_2\text{H}_5)_2\text{CrO}_4$.

Nearly insol. in H_2O .

— **dichromate**, $\frac{\text{OH}}{\text{Cl}}\text{Pt}(\text{N}_2\text{H}_5)_2\text{Cr}_2\text{O}_7$.

Ppt. (Cleve.)

— **nitrate** (Raewsky's nitrate),

$\frac{\text{OH}}{\text{Cl}}\text{Pt}(\text{N}_2\text{H}_5\text{NO}_2)_2$.

Sl. sol. in cold, more easily in hot H_2O . (Gerhardt.)

Chlorohyposulphuric acid, $\text{S}_2\text{O}_3\text{Cl}_4$.

See Sulphur oxytetrachloride.

Chloromanganic acid.

See Manganic hydrogen chloride.

Chloromercurosulphrous acid.

Ammonium chloromercurosulphite, $\text{NH}_4\text{SO}_3\text{HgCl}$.

Sol. in H_2O . (Barth, Z. phys. Ch. **1**.)

Barium chloromercurosulphite, $\text{Ba}(\text{SO}_3\text{HgCl})_2$.

Insol. in H_2O . (Barth.)

Potassium chloromercurosulphite, KSO_3HgCl .

Sol. in H_2O . (Barth.)

Sodium chloromercurosulphite, $\text{NaSO}_3\text{HgCl} + \text{H}_2\text{O}$.

Very sol. in H_2O . (Barth.)

Chloromolybdenum bromide, $\text{Cl}_4\text{Mo}_3\text{Br}_2 + 3\text{H}_2\text{O}$.

Insol. in H_2O and dil. acids; sol. in a + $6\text{H}_2\text{O}$. At first easily sol. in H_2O , precipitate soon forms. Can be cryst. from dil. $\text{HBr} + \text{Aq}$. Sol. in alcohol and (Blomstrand.)

Chloromolybdenum potassium bromide, $\text{Cl}_4\text{Mo}_3\text{Br}_2, 2\text{KBr} + 2\text{H}_2\text{O}$.

Decomp. by H_2O . Can be cryst. from + Aq . (Blomstrand.)

Chloromolybdenum chloride, Cl_4Mo_3 molybdenum dichloride, MoCl_5 .

Insol. in H_2O ; easily sol. in $\text{HCl} + \text{H}_2\text{SO}_4 + \text{Aq}$; sl. sol. in HNO_3 ; sol. in $\text{N} + \text{Aq}$, $\text{NaOH} + \text{Aq}$, or $\text{KOH} + \text{Aq}$, with formation of precipitate on boiling; sol. in alcohol and ether. (Blomstrand, J. **1**. 96.)

Very sol. in conc. HCl . (Rosenheim and Kohn, Z. anorg. 1910, **66**. 2.)

+ $3\text{H}_2\text{O}$. Insol. in H_2O .

+ $4\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O . (Liech Kempe, A. **170**. 351.)

+ $6\text{H}_2\text{O}$. Sol. in H_2O , alcohol, or (Blomstrand.)

Chloromolybdenum hydrogen chloride, $\text{Mo}_3\text{Cl}_6, \text{HCl} + 4\text{H}_2\text{O}$.

Sol. in H_2O , but ppt. forms after minutes. (Rosenheim and Kohn, Z. 1910, **66**. 5.)

Chloromolybdenum potassium chloride, $\text{Cl}_4\text{Mo}_3\text{Cl}_2, 2\text{KCl} + 2\text{H}_2\text{O}$.

Decomp. by pure H_2O ; can be recrystallized from $\text{HCl} + \text{Aq}$. (Blomstrand, J. **1**. 108.)

denum hydroxide, $\text{Cl}_4\text{Mo}_2(\text{OH})_2$

H_2O or alcohol. Easily sol. in if fresh, and washed only with H_2O . If washed with warm H_2O , it is insol. If precipitated hot, is insol. in H_2SO_4 or fuming HNO_3 . (Blomstrand, 77. 100.)

denum iodide, $\text{Cl}_4\text{Mo}_2\text{I}_2 + 3\text{H}_2\text{O}$.

Sol. in H_2O and alcohol.

denum potassium iodide, $\text{I}_2, 2\text{KI} + 2\text{H}_2\text{O}$.

by H_2O . Recryst. from $\text{HI} + \text{Aq}$.

denum oxybromide, $\text{Cl}_4\text{Mo}_2\text{OH Br}$

alcohol. (Blomstrand, J. pr. 77.

bdic acid, $(\text{OH}) + 7\text{H}_2\text{O}$.

oscopie. (Weinland, B. 1904, 37.

n tetrachloromolybdate, $(\text{NH}_4)_2 + 2\text{H}_2\text{O}$.

ic. Decomp. by H_2O . Sol. in alkalis and ammonia. (Weinland, 1905, 44. 83.)

rotrimolybdate, acid, $\text{Cl}_{14}(\text{Cs}_2\text{O}) + 22\text{H}_2\text{O}$.

ic. Decomp. by H_2O . Sol. in alkalis, and ammonia. (Wein-

trichloromolybdate, $(\text{OCs}) + \text{H}_2\text{O}$.

ic. Decomp. by H_2O . Sol. in alkalis and ammonia. (Wein-

trachloromolybdate, $(\text{OCs})_2$.

ic. Sol. in H_2O with decomp. e acids, alkalis, and ammonia. anorg. 1905, 44. 83.)

um trichloromolybdate, $(\text{OK}) + \text{H}_2\text{O}$.

ic. Decomp. by H_2O . Sol. in alkalis, and ammonia. (Wein-

tetrachloromolybdate, $(\text{K})_2 + 2\text{H}_2\text{O}$.

ic. Decomp. by H_2O . Sol. in alkalis, and ammonia. (Wein-

Potassium hydrogen chlorotrimolybdate, $\text{Mo}_2\text{O}_{11}\text{Cl}_{14}, \text{K}_2\text{O} + 6\text{H}_2\text{O}$.

Hydroscopic. Decomp. by H_2O . Sol. in dilute acids, alkalis, and ammonia. (Weinland.)

Monorubidium trichloromolybdate, $\text{MoOCl}_3(\text{ORb}) + \text{H}_2\text{O}$.

Hydroscopic. Decomp. by H_2O . Sol. in dilute acids, alkalis, and ammonia. (Weinland.)

Dirubidium tetrachloromolybdate, $\text{MoCl}_4(\text{ORb})_2$.

Hydroscopic. Decomp. by H_2O . Sol. in dilute acid, alkalis, and ammonia. (Weinland.)

Chloronitratoplatinamine nitrite, $\text{Cl Pt}(\text{NH}_2\text{NO}_2)_2$.

Easily sol. in H_2O .

Chloronitratoplatindiamine nitrate, $\text{Cl Pt}(\text{N}_2\text{H}_4\text{NO}_2)_2$.

Decomp. by H_2O with formation of $\text{Cl Pt}[(\text{NH}_2)_2\text{NO}_2]_2$.

— sulphate, $\text{Cl Pt}(\text{N}_2\text{H}_4)_2\text{SO}_4 + \text{H}_2\text{O}$.

Sl. sol. in cold, more easily in hot H_2O .

Chloronitritotetramine cobaltic chloride, $\text{Cl}(\text{NO}_2)\text{Co}(\text{NH}_3)_4\text{Cl}$.

Not very sol. in cold H_2O . (Jørgensen, Z. anorg. 5. 195.)

Chloronitritoplatinsemidiamine chloride, $\text{Cl}_2(\text{NO}_2)\text{Pt}(\text{NH}_3)_2\text{Cl}$.

100 pts. solution in H_2O sat. at 18° contain 1.8 pts. salt; sat. at 100° , 6 pts.

Insol. in abs. alcohol or ether. Not decomp. by conc. HNO_3 , HCl , or $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$, and by H_2SO_4 only at a high heat.

Formula given was $\text{PtN}_6\text{H}_{12}\text{Cl}_3\text{O}_6$. (Peyrone, J. B. 1855. 421.)

— nitrite, $\text{Cl}_2(\text{NO}_2)\text{Pt}(\text{NH}_3)_2\text{NO}_2$.

Sol. in H_2O . (Blomstrand.)

Chlorophosphatoplatindiamine phosphate, $\text{ClPt}(\text{N}_2\text{H}_4)_2\text{PO}_4 + 2\text{H}_2\text{O}$.

Nearly insol. in cold, and only very sl. sol. in hot H_2O . (Raewsky.)

Chloronitrous acid.

Iridium potassium chloronitrite, $\text{Ir}_2\text{Cl}_2(\text{NO}_2)_4, 6\text{KCl}$.

Ppt.; decomp. by boiling H_2O . Sol. in cold H_2O . (Leidié, C. R. 1902, 134. 1583.)

$\text{Ir}_2\text{K}_{12}\text{Cl}_{18}(\text{NO}_2)_8 + 4\text{H}_2\text{O}$. Ppt. (Quennessen, C. R. 1905, 141. 258.)

Chloropalladic acid.

Chloropalladates.

The chloropalladates are generally very sol. in H_2O , and sol. in alcohol. (v. Bonsdorff, Pogg. 17. 264.)

Ammonium chloropalladate, $(\text{NH}_4)_2\text{PdCl}_4$.

Sl. sol. in H_2O . (Berzelius.)

Barium chloropalladate.

Sol. in H_2O and alcohol. (v. Bonsdorff.)

Cadmium chloropalladate.

As above.

Cæsium chloropalladate, Cs_2PdCl_4 .

Nearly insol. in cold H_2O . Decomp. by boiling with H_2O or by hot conc. H_2SO_4 . (Gutbier, B. 1905, 38. 2386.)

Calcium chloropalladate.

Deliquescent; sol. in H_2O and alcohol. (v. Bonsdorff, 1829.)

Glucinum chloropalladate, $\text{GlPdCl}_4 + 8\text{H}_2\text{O}$.

Very hygroscopic, and sol. in H_2O .

Magnesium chloropalladate, $\text{MgPdCl}_4 + 6\text{H}_2\text{O}$.

Deliquescent; sol. in H_2O .

Nickel chloropalladate, $\text{NiPdCl}_4 + 6\text{H}_2\text{O}$.

Extremely deliquescent.

Potassium chloropalladate, K_2PdCl_4 .

Sl. sol. in cold H_2O . Decomp. by long boiling with H_2O . Sl. sol. in dil. $\text{HCl} + \text{Aq}$ without decomp. Insol. in NH_4Cl , KCl , or $\text{NaCl} + \text{Aq}$. Insol. in alcohol. (Berzelius.)

Rubidium chloropalladate, Rb_2PdCl_4 .

Insol. in cold H_2O . Decomp. by boiling with H_2O or by hot conc. H_2SO_4 . (Gutbier, B. 1905, 38. 2387.)

Zinc chloropalladate, $\text{ZnPdCl}_4 + 6\text{H}_2\text{O}$.

Very deliquescent. (v. Bonsdorff.)

Chloropalladous acid

Aluminum chloropalladite, $\text{Al}_2\text{Pd}_2\text{Cl}_{10} + 20\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O , alcohol, or ether. (Welkow, B. 7. 804.)

Ammonium chloropalladite, $(\text{NH}_4)_2\text{PdCl}_4 + \text{H}_2\text{O}$.

Easily sol. in H_2O . Insol. in alcohol. Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Claus.)

Easily sol. in H_2O . (Gutbier, B. 1905, 38. 2386.)

Barium chloropalladite.

Easily sol. in H_2O or alcohol.

Cadmium chloropalladite.

Not deliquescent.

Cæsium chloropalladite, Cs_2PdCl_4 .

Can be cryst. from hot H_2O . (Gutbier, B. 1905, 38. 2386.)

Calcium chloropalladite.

Deliquescent. Sol. in H_2O or alcohol.

Glucinum chloropalladite, $\text{GlPdCl}_4 + 6\text{H}_2\text{O}$.

Very hygroscopic; very sol. in H_2O , or ether. (Welkow.)

Magnesium chloropalladite.

Deliquescent. Easily sol. in H_2O . (v. Bonsdorff.)

Manganese chloropalladite.

Sol. in H_2O and alcohol.

Nickel chloropalladite.

Sol. in H_2O .

Potassium chloropalladite, K_2PdCl_4 .

Much more sol. in hot than cold H_2O . (Joannis, C. R. 95. 295.) Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Berzelius.) Sol. in cold sat. $\text{KCl} + \text{Aq}$. (Gibbs, Sil. Am. J. (2) 31. 70.) Insol. in alcohol. (Wollaston.) Somewhat sol. in alcohol of 0.84 sp. gr., but insol. in absolute alcohol; decomp. on boiling (Berzelius.)

Rubidium chloropalladite, Rb_2PdCl_4 .

Can be cryst. from hot H_2O . (Gutbier, B. 1905, 38. 2387.)

Sodium chloropalladite.

Deliquescent. Sol. in H_2O and alcohol.

Zinc chloropalladite.

Very deliquescent. Sol. in H_2O and alcohol. (v. Bonsdorff.)

Chlorophosphoarsenoiiridic acid,

$3\text{H}_3\text{PO}_3, 3\text{H}_3\text{PO}_4, 5\text{H}_3\text{AsO}_4(?)$.

Very sol. in H_2O . (Geisenheimer.)

Lead chlorophosphoarsenoiiridate,

$3\text{Pb}_2\text{H}_2(\text{PO}_3)_2, 3\text{Pb}_2(\text{PO}_4)_2, 5\text{Pb}_2\text{H}_2(\text{AsO}_4)_2$.

Insol. in H_2O .

Chlorophosphoiridic acid, $2\text{IrCl}_3, 3\text{H}_3\text{PO}_3$.

Very sol. in H_2O . Insol. in alcohol. (Geisenheimer, A. ch. (6) 23. 254.)
 $2\text{IrCl}_3, 3\text{H}_3\text{PO}_4$. Sol. in H_2O and alcohol.

um chlorophosphoiridate, $2\text{IrCl}_3, \text{H}_4\text{P}_2\text{O}_7, 3(\text{NH}_4)_2\text{HPO}_4$.

deliquescent. Very sol. in H_2O . (Cleve.)

orophosphoiridate, $4\text{IrCl}_3, (\text{PO}_4)_3, 3\text{PbH}_2(\text{PO}_3)_2$.

in H_2O or acetic acid; very sol. in dil. Aq. (Geisenheimer.)

lorophosphoiridate, $2\text{IrCl}_3, \text{H}_2\text{PO}_4, 3\text{AgH}_2\text{PO}_3$.

in H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$, and $+ \text{Aq}$. (Geisenheimer.)

hosphoplatinic acid.

loroplatinophosphoric acid.

hosphoric acid.

chlorophosphate, $3\text{ThO}_2, \text{ThCl}_4, \text{H}_2\text{O}$.

in H_2O and acids; decomp. by boil-
 H_2SO_4 and fusing with alkali car-
(Colani, C. R. 1909, 149. 208.)

latinamine chloride, $\text{Cl}_2\text{Pt} \begin{smallmatrix} \text{NH}_3\text{Cl} \\ \text{NH}_3\text{Cl} \end{smallmatrix}$.

about 700 pts. H_2O at 0° , and 33–34
 100° . Not attacked by boiling conc.
 H_2SO_4 . Sol. in boiling $\text{KOH} + \text{Aq}$
mp. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Cleve,
H. 10, 9. 30.)

ite, $\text{Cl}_2\text{Pt}(\text{NH}_2\text{NO}_2)_2$.

in cold, easily in hot H_2O .

ite silver nitrite, $\text{Cl}_2\text{Pt}(\text{NH}_2\text{NO}_2)_2, \text{O}_2$.

sol. in hot, sl. sol. in cold H_2O .

itochloride, $\text{Cl}_2\text{Pt} \begin{smallmatrix} \text{NH}_2\text{NO}_2 \\ \text{NH}_2\text{Cl} \end{smallmatrix}$.

H_2O . (Cleve.)

tin diamine bromide,

$\text{Pt}(\text{N}_2\text{H}_4\text{Br})_2$.

in hot H_2O . (Cleve.)

ride (Gros' chloride),

$\text{Pt}(\text{N}_2\text{H}_4\text{Cl})_2$.

insol. in cold, and only sl. sol. in hot
sl. in hot conc. $\text{KOH} + \text{Aq}$, with de-
Grimm.)

cold $\text{KOH} + \text{Aq}$ without decomp.
sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Buckton.)

(Raewsky.)

roplatinite, $\text{NCl}_2\text{Pt}(\text{N}_2\text{H}_4\text{Cl})_2, \text{PtCl}_4$.

sol. in hot H_2O .

roplatinite, $\text{Cl}_2\text{Pt}(\text{N}_2\text{H}_4\text{Cl})_2, \text{PtCl}_2$.

in H_2O . (Cleve.)

Chloroplatin diamine chromate,

$\text{Cl}_2\text{Pt}(\text{N}_2\text{H}_4)_2\text{CrO}_4$.

Nearly insol. in H_2O . (Cleve.)

— **dichromate**, $\text{Cl}_2\text{Pt}(\text{N}_2\text{H}_4)_2\text{Cr}_2\text{O}_7$.

Sl. sol. in cold, more sol. in hot H_2O .
(Cleve.)

— **nitrate** (Gros' nitrate), $\text{Cl}_2\text{Pt}(\text{N}_2\text{H}_4\text{NO}_3)_2$.

Much more easily sol. in hot than in cold
 H_2O . Sol. in hot $\text{KOH} + \text{Aq}$ with decomp.
Nearly insol. in conc. $\text{HNO}_3 + \text{Aq}$.

— **nitritochloride**, $\text{Cl}_2\text{Pt} \begin{smallmatrix} \text{N}_2\text{H}_4\text{NO}_2 \\ \text{N}_2\text{H}_4\text{Cl} \end{smallmatrix}$.

Ppt. (Jørgensen.)

— **phosphate**.

See **Chlorophosphatoplatin diamine phos-
phate**.

— **sulphate**, $\text{Cl}_2\text{Pt}(\text{N}_2\text{H}_4)_2\text{SO}_4$.

Sl. sol. in both cold or hot H_2O . (Cleve.)
 $+ x\text{H}_2\text{O}$. Sl. sol. in cold, easily in hot
 H_2O . (Grimm.)

— **sulphocyanide**, $\text{Cl}_2\text{Pt}(\text{N}_2\text{H}_4)_2(\text{CNS})_2$,
 $+ \text{H}_2\text{O}$.

Ppt. (Cleve.)

Chloroplatin monodiamine chloride,

$\text{Cl}_2\text{Pt} \begin{smallmatrix} (\text{NH}_3)_2\text{Cl} \\ \text{NH}_3\text{Cl} \end{smallmatrix}$.

Quite easily sol. in H_2O . (Cleve.)

**Chloroplatin semidiamine carbonate chlo-
ride**, $2\text{Pt}(\text{NH}_3)_2\text{Cl}_2, \text{Pt}_2(\text{NH}_3)_4\text{Cl}_2(\text{CO}_3)_2$.

Sl. sol. in H_2O ; insol. in alcohol and ether.
Decomp. by cold $\text{HCl} + \text{Aq}$. (Schon, Z. anorg.
1897, 13. 37.)

Chloroplatin semidiamine chloride,

$\text{Cl}_2\text{Pt}(\text{NH}_3)_2\text{Cl}$.

Sol. in 300 pts. H_2O at 0° , and 65 pts. at
 100° . Not decomp. by conc. H_2SO_4 . Sol. in
 $\text{KOH} + \text{Aq}$ without decomp. (Cleve.)

Chloroplatinic acid, $\text{H}_2\text{PtCl}_6 + 6\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O , alcohol, or ether.
 $+ 4\text{H}_2\text{O}$. Deliquescent. (Pigeon, C. R.
112. 1218.)

$\text{PtCl}_4, \text{HCl} + 2\text{H}_2\text{O}$. (Pigeon.)

Aluminum chloroplatinate, $\text{AlCl}_3, \text{PtCl}_4 +$
 $15\text{H}_2\text{O}$.

Very sol. in H_2O and alcohol. (Welkow, B.
7. 304.)

Insol. in ether.

Ammonium chloroplatinate, $(\text{NH}_4)_2\text{PtCl}_6$.

Sl. sol. in cold, more easily in hot H_2O .
(Fresenius.)

100 pts. H_2O dissolve 0.666 pt. at ord.
temp. and 12.5 pts. at 100° . (Crookes, C. N.
9. 37.)

Insol. in cold HCl + Aq. Separates out on cooling from solution in hot HCl, HNO₃, or H₂SO₄. (Fischer.)

Very sl. sol. in cold, easily in hot NH₄OH + Aq. (Fresenius.)

Conc. NH₄Cl + Aq ppts. it almost completely from aqueous solution. (Böttger.)

Sol. in NH₄ succinate + Aq. (Döpping.)

Less sol. in H₂PtCl₆ + Aq than in H₂O. (Rogojski, A. ch. (3) 41. 452.)

Sol. in SnCl₂ + Aq. (Fischer.)

Very sol. with decomp. in KCNS + Aq. (Claus.)

At 15–20°, sol. in 26,535 pts. 97.5% alcohol, in 1476 pts. 76% alcohol, and in 665 pts. 55% alcohol. If free HCl is present, it is sol. in 672 pts. 76% alcohol. (Fresenius, A. 59. 118.)

Insol. in absolute alcohol or ether.

Barium chloroplatinate, BaPtCl₆ + 6H₂O.

Permanent; sol. in H₂O; decomp. by alcohol. (v. Bonsdorff, Pogg. 17. 250.)

Barium monochloroplatinate, PtCl(OH)₂Ba + H₂O.

Insol. in H₂O and in org. solvents. (Bellucci, C. C. 1903, I. 131.)

Barium pentachloroplatinate, OH.PtCl₅Ba + H₂O.

(Miolati, Chem. Soc. 1900, 78. (2) 732.)

Cadmium chloroplatinate, CdPtCl₆ + 6H₂O.

Deliquescent, and easily sol. in H₂O. (v. Bonsdorff.)

Cæsium chloroplatinate, Cs₂PtCl₆.

100 pts. H₂O dissolve at:

0°	10°	20°	30°
0.024	0.050	0.079	0.110
40°	50°	60°	70°
0.142	0.177	0.213	0.251
80°	90°	100°	
0.291	0.332	0.377	

pts. Cs₂PtCl₆. (Bunsen, Pogg. 113. 337.)

Sol. in 1308 pts. H₂O at 15°, and 261 pts. at 100°. (Crookes, C. N. 9. 205.)

Calcium chloroplatinate, CaPtCl₆ + 8H₂O.

Deliquescent; easily sol. in H₂O. (v. Bonsdorff.)

Calcium monochloroplatinate, PtCl(OH)₂Ca + H₂O.

Insol. in H₂O and in org. solvents. (Bellucci, C. C. 1903, I. 131.)

Cerium chloroplatinate, CeCl₃, PtCl₄ + 13H₂O.

Deliquescent; very sol. in H₂O or alcohol; insol. in ether. (Marignac.)

4CeCl₃, 3PtCl₄ + 8H₂O. Deliquescent; easily sol. in H₂O or alcohol; insol. in ether. (Holzmann, J. pr. 84. 80.)

Chromium chloroplatinate, CrCl₃, PtCl₄ + 10½H₂O.

Deliquescent. (Nilson, B. 9. 1056.)

+10H₂O. Very sol. in H₂O and alcohol. Nearly insol. in acetone. (Higley, J. Am. Chem. Soc. 1904, 26. 617.)

Cobalt chloroplatinate, CoPtCl₆ + 6H₂O.

Very deliquescent. (Jørgensen.)

Copper chloroplatinate, CuPtCl₆ + 6H₂O.

Deliquescent in moist air. (v. Bonsdorff.)

Didymium chloroplatinate, DiCl₃, PtCl₄ + 13H₂O.

Less deliquescent than the cerium salt. (Marignac.)

+10½H₂O. Deliquescent. (Cleve, Bull. Soc. (2) 43. 361.)

Erbium chloroplatinate, ErCl₃, PtCl₄ + 11H₂O.

Very deliquescent. (Cleve.)

Gadolinium chloroplatinate, GdCl₃, PtCl₄ + 10H₂O.

Ppt. (Benedicko, Z. anorg. 1900, 22. 204.)

Glucinum chloroplatinate, GlPtCl₆ + 8H₂O.

Deliquescent in moist air. Very sol. in H₂O, moderately in alcohol. Insol. in ether. (Welkow, B. 6. 1288.)

Indium chloroplatinate, 2InCl₃, 5PtCl₄ + 36H₂O.

Deliquescent. (Nilson.)

Iron (ferrous) chloroplatinate, FePtCl₆ + 6H₂O.

Deliquescent. (Topsoë.)

Iron (ferric) chloroplatinate, FeCl₃, PtCl₄ + 10½H₂O.

Deliquescent. (Nilson.)

Lanthanum chloroplatinate, LaCl₃, PtCl₄ + 13H₂O.

Deliquescent; extremely sol. in H₂O (Cleve.)

Lead chloroplatinate, PbPtCl₆ + 3H₂O.

Easily sol. in H₂O and alcohol (Topsoë), with decomp. (Birnbäum, Zeit. Ch. 1867. 520.)

Lead monochloroplatinate, [PtCl(OH)₂]₂Pb, Pb(OH)₂.

Ppt. (Bellucci, Chem. Soc. 1902, 82, II. 155.)

Lead pentachloroplatinate, basic, PtCl₅(OH)Pb, Pb(OH)₂.

(Miolati, Chem. Soc. 1900, 78. (2) 732.)

Lithium chloroplatinate, Li₂PtCl₆ + 6H₂O.

Extremely deliquescent (Jørgensen); efflorescent. Easily sol. in H₂O, alcohol, or ether-alcohol; insol. in ether. (Scheibler.)

Lithium pentachloroplatinate, OH.PtCl₅Li.

Very hygroscopic. (Miolati, Chem. Soc. 1900, 78 (2) 732.)

cesium chloroplatinate, $MgPtCl_6 + 6H_2O$.
in H_2O and abs. alcohol.
 $12H_2O$. Sol. in H_2O .

ganese chloroplatinate, $MnPtCl_6 + 6H_2O$.
t deliquescent; sol. in H_2O .
 $12H_2O$. Sl. efflorescent.

al chloroplatinate, $NiPtCl_6 + 6H_2O$.
in H_2O .

assium chloroplatinate, K_2PtCl_6 .
pts. H_2O dissolve at:
10° 20° 30° 40° 50°
0.90 1.12 1.41 1.76 2.17 pts. K_2PtCl_6 ,
70° 80° 90° 100°
3.19 3.79 4.45 5.18 pts. K_2PtCl_6 .
(Bunsen, Pogg. 113. 337.)

pts. H_2O dissolve 0.926 pt. at 15°, and
pts. at 100°. (Crookes, C. N. 9. 205.)

g. H_2O dissolve at:
16° 25° 35° 48°
2 0.6718 0.8641 1.132 1.745 g. K_2PtCl_6 ,
69° 68° 78° 92°
396 2.913 3.589 4.484 g. K_2PtCl_6 .
Archibald, J. Am. Chem. Soc. 1908, 30. 752.)
not attacked by cold conc. H_2SO_4 . (Las-
pe.)
sol. in cold, more easily in hot dil. acids.
sol. in $KCl + Aq$ than in H_2O , and nearly
in sat. $KCl + Aq$. (Schrötter, W. A. B.
268.)

Solubility in $KCl + Aq$ at 20°.

mol. KCl per l. of $KCl + Aq$.	G. K_2PtCl_6 in 100 g. of solution.
0.00	0.7742
0.20	0.0236
0.25	0.0207
0.50	0.0109
1.00	0.0046
2.00	0.0045
3.00	0.0043
4.00	0.0042
sat.	0.0034

Archibald, J. Am. Chem. Soc. 1908, 30. 757.)

Solubility in $NaCl + Aq$ at 16°.

mol. $NaCl$ per litre of $NaCl + Aq$.	G. K_2PtCl_6 in 100 g. of solution
0.00	0.672
0.05	0.700
0.10	0.729
0.25	0.758
0.50	0.775
0.75	0.791
1.00	0.805
2.00	0.834

Archibald, J. Am. Chem. Soc. 1908, 30. 757.)

Sol. in $KOH + Aq$. Insol. in cold or hot
alkali carbonates or bicarbonates + Aq . (Rose.
Easily sol. in warm $Na_2S_2O_3 + Aq$. (Himly.)
Sol. in $NH_4Cl + Aq$. (Brett.)
Sol. in NH_4 succinate + Aq . (Döpping.)
At 15–20°, sol. in 12,083 pts. absolute al-
cohol, in 3775 pts. 76% absolute alcohol, and
in 1053 pts. 55% absolute alcohol. (Fresen-
ius.)
Sol. in 1835 pts. 76% alcohol containing
 HCl at 15–20°. (Fresenius.)
Nearly absolutely insol. in alcohol con-
taining ether.
Sol. in 42,600 pts. absolute alcohol. (Precht,
Z. anal. 18. 509.)
1 l. methyl alcohol dissolves 0.072 g. at 20°.
(Peligot, Monit. Sci. 1892, (4) 6. I, 873.)

Solubility in methyl alcohol + Aq at 20°.

% alcohol by wt.	G. K_2PtCl_6 in 100 g. of solution
0	0.7742
5	0.5350
10	0.4120
20	0.2642
30	0.1831
40	0.1165
50	0.0625
60	0.0325
70	0.0182
80	0.0124
90	0.0038
100	0.0027

(Archibald, J. Am. Chem. Soc. 1908, 30. 755.)

Solubility in ethyl alcohol + Aq at 20°.

% alcohol by wt.	G. K_2PtCl_6 in 100 g. of solution
0	0.7742
5	0.4910
10	0.3720
20	0.2180
30	0.1340
40	0.0760
50	0.0491
60	0.0265
70	0.0128
80	0.0085
90	0.0025
100	0.0009

(Archibald, J. Am. Chem. Soc. 1908, 30. 755.)

Solubility in isobutyl alcohol + Aq at 20°.

% alcohol	g. K_2PtCl_6 in 100 g. of solution
0	0.7742
8.20	0.6250
sat.	0.3180

(Archibald, J. Am. Chem. Soc. 1908, 30. 755.)

Potassium pentachlorohydroplatinat, $K_2(PtCl_5OH)$.Easily sol. in H_2O . (Ruff, B. 1913. 46. 925.)**Praseodymium chloroplatinate**, $PrCl_3, PtCl_4 + 12H_2O$.Very sol. in H_2O . Sol. in conc. HCl . (von Scheele, Z. anorg. 1898, 18. 353.)**Rubidium chloroplatinate**, Rb_2PtCl_6 .100 pts. H_2O dissolve at:

0°	10°	20°
0.184	0.154	0.141 pts. Rb_2PtCl_6 ,
30°	40°	50°
0.145	0.166	0.203 pts. Rb_2PtCl_6 ,
60°	70°	80°
0.253	0.329	0.417 pts. Rb_2PtCl_6 ,
90°	100°	
0.521	0.634	pts. Rb_2PtCl_6 .

(Bunsen, Pogg. 113. 337.)

Sol. in 740 pts. H_2O at 15°, and 157 pts. at 100°. (Crookes, C. N. 9. 205.)

Insol. in alcohol.

Samarium chloroplatinate, $SmCl_3, PtCl_4 + 10\frac{1}{2}H_2O$.Deliquescent. Very sol. in H_2O . (Cleve, Bull. Soc. (2) 43. 165.)**Silver chloroplatinate**, Ag_2PtCl_6 .Ppt. Gradually decomp. by H_2O into $AgCl$ and $PtCl_4$. (Jørgensen, J. pr. (2) 16. 345.) $Ag_2PtCl_6(OH)_2$. Ppt.**Silver monochloroplatinate**, $[PtCl(OH)_2]Ag_2$.

Ppt. (Bellucci, Chem. Soc. 1902, 82. (2). 155.)

Silver pentachloroplatinate, $(OH)PtCl_5Ag_2$.Ppt.; stable in boiling H_2O . (Miolati, Chem. Soc. 1900, 78 (2). 732.)**Silver chloroplatinate ammonia**, $Ag_2PtCl_6, 2NH_3$.Insol. in H_2O . (Birnbbaum.)**Sodium chloroplatinate**, $Na_2PtCl_6 + 6H_2O$.Easily sol. in H_2O . Sat. solution at 15° contains 39.77 g. Na_2PtCl_6 and has sp. gr. of 1.368. Sol. in $NaCl + Aq$. More sol. in absolute alcohol than in 95% alcohol. Sat. solution in abs. alcohol contains 11.90%; 95% alcohol, 6.34%. Mixture of equal parts of alcohol and ether dissolve 2.43%. Insol. in ether. (Precht, Z. anal. 18. 502.)**Sodium pentachloroplatinate**, $(OH)PtCl_5Na_2$.

Exists only in solution. (Miolati, l.c.)

Strontium chloroplatinate, $SrPtCl_6 + 8H_2O$.Very sol. in H_2O .**Strontium monochloroplatinate**, $PtCl(OH) + H_2O$.Insol. in H_2O and org. solvents. (B. C. C. 1903, I. 131.)**Thallium chloroplatinate**, Tl_2PtCl_6 .Very sl. sol. in H_2O . Sol. in 15.5 H_2O at 15°, and 1948 pts. at 100°. (Cr**Thallium monochloroplatinate**, $[PtCl(OH)]Tl$. Ppt. (Bellucci, Chem. Soc. 1902, 82. 155.)**Thallium pentachloroplatinate**, $(OH)PtCl_5$ (Miolati, Chem. Soc. 1900, 78 (2). 732.)**Thorium chloroplatinate**, $ThCl_4, 12H_2O$.

Very deliquescent. (Cleve, Bull. S. 21. 118.)

Tin (stannic) chloroplatinate, $SnCl_4, 12H_2O$.

(Nilson, B. 9. 1142.)

Ytterbium chloroplatinate, $2YbCl_3, 12H_2O$, and $+35H_2O$.

Ppt. (Cleve, Z. anorg. 1902, 32. 13)

Vanadyl chloroplatinate, $(VO)PtCl_4 + 10\frac{1}{2}H_2O$.Sol. in H_2O ; cryst. from $PtCl_4$ (Brauner, M. 3. 58.)**Yttrium chloroplatinate**, $4YCl_3, 52H_2O$.Very deliquescent. (Cleve.) $2YCl_3, 3PtCl_4 + 30H_2O$. (Nilson, 1059.) $2YCl_3, PtCl_4 + 21H_2O$. (Nilson.)**Zinc chloroplatinate**, $ZnPtCl_6 + 6H_2O$.Deliquescent; sol. in H_2O and alcohol.**Zinc tetrachloroplatinate**, $ZnPt(OH)_4 + 5H_2O$.Extremely sol. in H_2O and alcohol. (Miolati, Z. anorg. 1900, 22. 458.)**Zirconyl chloroplatinate**, $(ZrO)PtCl_6 + 6H_2O$ (Nilson.)**Chloroplatinocyanhydric acid**, $ClPtP_2O_6H_4 = ClPt \begin{matrix} \nearrow P(OH)_2 \\ \searrow PO(CN)_2 \end{matrix}$ Not deliquescent. Sol. in H_2O . (Schäfer, Bull. Soc. (2) 18. 154.)**Chloroplatinocyanhydric acid**, $H_2Pt(CN)_4Cl_2$.

See Perchloroplatinocyanhydric acid.

ium chloroplatinocyanide, $5K_2Pt(CN)_4 \cdot Pt(CN)_4Cl_2 + 21H_2O$.
in H_2O ; insol. in alcohol.

chloroplatinocyanide,
 $K_2(PtCl_2(CN)_4)_2$.
(Miolati, C. C. 1901, I. 500.)

oplatinophosphoric acid,
 $H_2PtP(OH)_3$.
deliquescent, and sol. in H_2O .
zenberger, Bull. Soc. (2) 17. 493.)

chloroplatinophosphate, $Pb_2(Cl_2PtPO_3)_2 \cdot 8H_2O$.

$Cl_2PtPO_3)_2 \cdot 2PbO + 4H_2O$. Ppt.
zenberger, Bull. Soc. (2) 17. 494.)

chloroplatinophosphate, $Ag_2HPO_3 \cdot PtCl_2$.
(Schützenberger, Bull. Soc. (2) 17.

oplatinodiphosphoric acid, $PtCl_2 \cdot (OH)_6$.
deliquescent, and easily sol. in H_2O .
zenberger, Bull. Soc. (2) 18. 153.)

oplatinopyrophosphoric acid,
$$IPt \begin{array}{c} \nearrow P(OH)_2 \\ \searrow O \\ \searrow P(OH)_2 \end{array}$$

deliquescent than chloroplatinodiphosphoric acid.

oplatinous acid, H_2PtCl_4 .
own only in solution.

ium chloroplatinite, $AlPtCl_4 + 10\frac{1}{2}H_2O$.
y deliquescent; sol. in H_2O . (Nilson, (2) 15. 260.)

ium chloroplatinite, $(NH_4)_2PtCl_4$.
sol. in cold, easily in hot H_2O . Insol. in alcohol. (Peyrone, A. 55. 206.)

n chloroplatinite, $BaPtCl_4 + 3H_2O$.
deliquescent; sol. in H_2O . Very sl. sol. in 93% alcohol.

ium chloroplatinite ammonia, $CdPtCl_4 \cdot NH_3$.
sol. in H_2O or $NH_4OH + Aq$. Sol. in Aq . (Thomsen, B. 2. 668.)

m chloroplatinite, Cs_2PtCl_4 .
sol. in cold, easily in hot H_2O .
pts. H_2O dissolve 3.4 pts. salt at 20°
" " 6.73 " " 40°
" " 8.68 " " 60°
" " 10.92 " " 80°
" " 12.10 " " 100°.
(Godeffroy, A. 181. 176.)

Cs_2PtCl_4 . Ppt. Very sensitive to sunlight. Decomp. by H_2O into the higher and lower chlorides. (Wöhler, B. 1909, 42. 4104.)

Calcium chloroplatinite, $CaPtCl_4 + 8H_2O$.
Deliquescent; sol. in H_2O .

Cerium chloroplatinite, $CeCl_3 \cdot 2PtCl_2 + 10\frac{1}{2}H_2O$.
Deliquescent; easily sol. in H_2O . (Nilson, B. 9. 1847.)

Chromium chloroplatinite, $Cr_2Pt_2Cl_{12} + 18H_2O$.
Deliquescent.

Cobalt chloroplatinite, $CoPtCl_4 + 6H_2O$.
Sl. deliquescent in moist, efflorescent in dry air.

Copper chloroplatinite, $CuPtCl_4 + 6H_2O$.
Extremely deliquescent. (Topsøe.)

Copper chloroplatinite ammonia (cuprammonium chloroplatinite),
 $Cu(NH_3)_4PtCl_4$.
Insol. in H_2O or $NH_4OH + Aq$; easily sol. in $H_2SO_4 + Aq$. (Millon and Commaille, C. R. 57. 822.)

Didymium chloroplatinite, $DiCl_3 \cdot 2PtCl_2 + 10H_2O$.
Deliquescent; very sol. in H_2O . (Nilson.)
 $2DiCl_3 \cdot 3PtCl_2 + 18H_2O$. As above. (Nilson.)

Erbium chloroplatinite, $ErPtCl_4 + 13\frac{1}{2}H_2O$.
Deliquescent.
 $Er_2Pt_2Cl_{12} + 24H_2O$. Deliquescent in moist air.

Glucinum chloroplatinite, $GlPtCl_4 + 5H_2O$.
Deliquescent in moist air. Sol. in H_2O in all proportions.

Iron (ferrous) chloroplatinite, $FePtCl_4 + 7H_2O$.
Deliquescent. Rather sl. sol. in cold, very sol. in hot H_2O . (Nilson.)

Lanthanum chloroplatinite, $La_2Pt_2Cl_{12} + 18$,
and $27H_2O$.
Deliquescent.

Lead chloroplatinite, $PbPtCl_4$.
Insol. in cold H_2O .

Lithium chloroplatinite, $Li_2PtCl_4 + 6H_2O$.
Sol. in H_2O .

Magnesium chloroplatinite, $MgPtCl_4 + 6H_2O$.
Not very deliquescent; very sol. in H_2O .

Manganese chloroplatinite, $\text{MnPtCl}_4 + 6\text{H}_2\text{O}$.

As the Mg salt.

Mercurous chloroplatinite.

Ppt.

Nickel chloroplatinite, $\text{NiPtCl}_4 + 6\text{H}_2\text{O}$.

As the Co salt.

Potassium chloroplatinite, K_2PtCl_4 .

Moderately sol. in H_2O ; insol. in alcohol.

Rubidium chloroplatinite, Rb_2PtCl_4 .

Sl. sol. in cold; easily in hot H_2O .

Silver chloroplatinite, Ag_2PtCl_4 .

Insol. in H_2O . $\text{NH}_4\text{OH} + \text{Aq}$ dissolves out AgCl . (Lang.)

AgCl , $\text{PtCl}_2(?)$. As above. (Commaille, Bull. Soc. (2) 6. 262.)

Silver chloroplatinite ammonia, $\text{Ag}_2\text{PtCl}_4, 4\text{NH}_3$.

(Thomsen.)

Sodium chloroplatinite, $\text{Na}_2\text{PtCl}_4 + 4\text{H}_2\text{O}$.

Deliquescent; very sol. in H_2O .

Strontium chloroplatinite, $\text{SrPtCl}_4 + 6\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O .

Thallium chloroplatinite, Tl_2PtCl_4 .

Very sl. sol. even in boiling H_2O .

Thorium chloroplatinite, $\text{Th}_2\text{Pt}_2\text{Cl}_{14} + 24\text{H}_2\text{O}$.

Very deliquescent.

Yttrium chloroplatinite, $\text{Y}_2\text{Pt}_2\text{Cl}_{12} + 24\text{H}_2\text{O}$.

Deliquescent.

Zinc chloroplatinite, $\text{ZnPtCl}_4 + 6\text{H}_2\text{O}$.

Sl. sol. in cold, more easily in hot H_2O ; insol. in alcohol.

Zinc chloroplatinite ammonia, $\text{ZnPtCl}_4, 4\text{NH}_3$.

Sl. sol. in H_2O ; easily sol. in $\text{HCl} + \text{Aq}$. Insol. in alcohol. (Thomsen, J. B. 1868. 278.)

Zirconyl chloroplatinite, $(\text{ZrO})\text{PtCl}_4 + 8\text{H}_2\text{O}$.

(Nilson.)

Trichloroplatinous acid, $\text{H}_2\text{Pt}(\text{OH})\text{Cl}_3$.

Sol. in H_2O . (Miolati, Z. anorg. 1902, 33. 265.)

+ H_2O . (Nilson, J. pr. (2) 15. 260.)

Lead trichloroplatinite, $\text{PbPt}(\text{OH})\text{Cl}_3$.

Ppt. (Miolati.)

Silver trichloroplatinite, $\text{Ag}_2\text{Pt}(\text{OH})\text{Cl}_3$.

Ppt. (Miolati.)

Chloroplatosulphurous acid.

Ammonium chloroplatosulphite, acid, $\text{NH}_4\text{PtClSO}_3, \text{H}_2\text{SO}_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Birnbbaum, A. 152. 149.)

Ammonium chloroplatosulphite chloride sulphite, $\text{NH}_4\text{PtClSO}_3, (\text{NH}_4)_2\text{SO}_3, \text{NH}_4\text{Cl}$.

Very deliquescent. (Birnbbaum.)

Ammonium chloroplatosulphite sulphite, $\text{NH}_4\text{ClPtSO}_3, (\text{NH}_4)_2\text{SO}_3 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Birnbbaum.)

Barium chloroplatosulphite chloride ammonium chloride, $\text{Ba}(\text{ClPtSO}_3)_2, \text{Ba}(\text{PtClSO}_3)\text{Cl}, 6\text{NH}_4\text{Cl} + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Birnbbaum.)

Potassium chloroplatosulphite ammonium chloride, $\text{KPtClSO}_3, 2\text{NH}_4\text{Cl}$.

Very deliquescent. (Birnbbaum, A. 152. 142.)

Potassium chloroplatosulphite chloride, $\text{KPtClSO}_3, 2\text{KCl}$.

Deliquescent; sol. in H_2O . (Birnbbaum, A. 152. 145.)

Potassium chloroplatosulphite ammonium potassium sulphite, $\text{KPtClSO}_3, (\text{NH}_4)\text{KSO}_3 + 3\text{H}_2\text{O}$.

Very deliquescent. (Birnbbaum, A. 152. 120.)

Sodium chloroplatosulphite ammonium chloride, $\text{NaPtClSO}_3, 2\text{NH}_4\text{Cl}$.

Very deliquescent. (Birnbbaum, A. 152. 117.)

Chloroplumbic acid, H_2PbCl_4 .

Decomp. in solution on standing. (Gutbier, J. pr. 1914, (2) 90. 497.)

Ammonium chloroplumbate, $(\text{NH}_4)_2\text{PbCl}_4$.

Ppt. Difficultly sol. in a small amount of H_2O . Solution decomp. slowly when cold, more rapidly when warmed.

Decomp. by a large amount of H_2O . Sol. without decomp. in 20% HCl . Decomp. by dil. acids and alkalis. (Elbs, Z. Elektrochem. 1903, 9. 778.)

Difficultly sol. in small amount of H_2O and solution decomp. slowly in the cold, more rapidly when warmed. Decomp. by a large amount of H_2O . (Gutbier, J. pr. 1914, (2) 90. 498.)

Sol. in cold HNO_3 without decomp. (Friedrich, M. 1893, 14. 511.)

Insol. in conc. $\text{NH}_4\text{Cl} + \text{Aq}$. (Nikoljukin, B. 18. 370 R.)

$5\text{NH}_4\text{Cl}, 2\text{PbCl}_4$. Not hygroscopic. Decomp. by H_2O with pptn. of PbO_2 . Sol. in $\text{HCl} + \text{Aq}$ and in cold $\text{HNO}_3 + \text{Aq}$ without

(Classen and Zahorski, Z. anorg. 4. 335.)
 ition is $2\text{NH}_4\text{Cl}$, PbCl_4 . (Fried-
 l. B. 102, 2b. 527.)

chloroplumbate, Cs_2PbCl_4 .

absolutely insol. in conc. $\text{CsCl} + \text{Aq}$
 e of Cl . (Wells, Z. anorg. 4. 335.)
 conc. $\text{HCl} + \text{Aq}$ containing PbCl_4 dis-
 10049 g. Cs_2PbCl_4 . (Wells, Z. anorg.

with H_2O as the corresponding am-
 salt. (Gutbier, J. pr. 1914, (2)

chloroplumbate, K_2PbCl_4 .

p. by H_2O ; sol. in $\text{KCl} + \text{Aq}$. (Wells,
 4. 335.)
 decomp. in the air. (Gutbier, J.
 (2) 90. 499.)

chloroplumbate, Rb_2PbCl_4 .

p. by H_2O ; sl. sol. in conc. $\text{RbCl} +$
 lls, Z. anorg. 4. 335.)
 conc. $\text{HCl} + \text{Aq}$ containing PbCl_4
 0.003 g. Rb_2PbCl_4 . (Wells, Z.
 341.)
 with H_2O as the corresponding am-
 salt. (Gutbier, J. pr. 1914, (2)

p. by conc. H_2SO_4 . Insol. in 96%
 (Erdmann, A. 1896, 294. 76.)

purpleochromium bromide,
 $(\text{NH}_4)_2\text{Br}_2$.

hat more easily sol. in H_2O than the
 (Jørgensen, J. pr. (2) 20. 105.)

ride, $\text{CrCl}(\text{NH}_4)_2\text{Cl}_2$.

tly sol. in cold, and decomp. by hot

lissolves in 154 pts. H_2O at 16° .
 conc. $\text{HCl} + \text{Aq}$. More sol. in dil.
 aq than in H_2O . Sol. in $\text{NH}_4\text{OH} +$
 it decomp. (Jørgensen, J. pr. (2) 20.

curic chloride, $\text{CrCl}(\text{NH}_4)_2\text{Cl}_2$,
 Cl_2 .

fficultly sol. in H_2O . (Jørgensen.)

roplatinate, $\text{CrCl}(\text{NH}_4)_2(\text{PtCl}_4)$.

ely difficultly sol. in H_2O . (Jørgen-

mate, $\text{CrCl}(\text{NH}_4)_2(\text{CrO}_4)$.

in H_2O ; sl. more sol. than chloro-
 cobalt chromate. (Jørgensen.)

ionate, $\text{CrCl}(\text{NH}_4)_2(\text{S}_2\text{O}_8)$.

. sol. in cold, but much more easily
 O. (Jørgensen.)

Chloropurpleochromium ferrocyanide,
 $[\text{CrCl}(\text{NH}_4)_2]_2\text{Fe}(\text{CN})_6 + 4\text{H}_2\text{O}$.

Very difficultly sol. in cold H_2O . (Jørgen-
 sen.)

— **fluosilicate**, $\text{CrCl}(\text{NH}_4)_2(\text{SiF}_6)$.

Very difficultly sol. in H_2O . Insol. in
 $\text{H}_2\text{SiF}_6 + \text{Aq}$. (Jørgensen, J. pr. (2) 20. 105.)

— **mercuric iodide**, $\text{CrCl}(\text{NH}_4)_2\text{I}_2$, 2HgI_2 .

Decomp. by H_2O ; sol. in alcohol and warm
 $\text{KCN} + \text{Aq}$.

$\text{CrCl}(\text{NH}_4)_2\text{I}_2$, HgI_2 . Very difficultly sol.
 in cold H_2O ; easily sol. in $\text{KCN} + \text{Aq}$. (Jør-
 gensen, l.c.)

— **nitrate**, $\text{CrCl}(\text{NH}_4)_2(\text{NO}_3)_2$.

Sol. in 71 pts. H_2O at 17.5° . Insol. in
 $\text{HNO}_3 + \text{Aq}$. (Jørgensen.)

— **oxalate**, $\text{CrCl}(\text{NH}_4)_2\text{C}_2\text{O}_4$.

Very sl. sol. in cold H_2O . (Jørgensen, l.c.)

— **sulphate**, $\text{CrCl}(\text{NH}_4)_2\text{SO}_4 + 2\text{H}_2\text{O}$.

Sol. in H_2O ; precipitated by alcohol. (Jør-
 gensen.)

— **sulphate, acid**, $[\text{CrCl}(\text{NH}_4)_2]_2\text{SO}_4(\text{HSO}_4)_6$.

Quite sol. in H_2O . (Jørgensen, J. pr. (2)
 20. 185.)

— **pentasulphide**, $\text{CrCl}(\text{NH}_4)_2\text{S}_5$.

Very sl. sol. in cold, easily sol. in warm
 H_2O . Decomp. by dil. $\text{HCl} + \text{Aq}$. Insol. in
 alcohol. (Jørgensen.)

Chloropurpleocobaltic bromide,
 $\text{CoCl}(\text{NH}_4)_2\text{Br}_2$.

Properties resemble the chloride very
 closely. Sol. in 214 pts. H_2O at 14.3° (Jør-
 gensen, J. pr. (2) 18. 205.)

— **mercuric bromide**,

$4\text{CoCl}(\text{NH}_4)_2\text{Br}_2$, 9HgBr_2 .

Ppt. (J.)

— **bromoplatinate**, $\text{CoCl}(\text{NH}_4)_2\text{Br}_2$, PtBr_4 .

Very sl. sol. in H_2O . (J.)

— **carbonate**, $\text{CoCl}(\text{NH}_4)_2\text{CO}_3 + 4\frac{1}{2}\text{H}_2\text{O}$.

Efflorescent; very easily sol. in H_2O . (J.)

— **chloride**, $\text{CoCl}(\text{NH}_4)_2\text{Cl}_2$.

Very sl. sol. in cold, more easily in hot H_2O .
 Sol. in 244 pts. H_2O at 15.5° . (Claudet, Phil.
 Mag. J. (4) 2. 253.) In 287 pts. H_2O at 10.2°
 and 255 pts. at 11.5° . (Rose, Pogg. 20. 152.)
 100 pts. H_2O dissolve 0.232 pt. CoCl_2 , 5NH_3 ,
 at 0° , and 1.031 pts. at 46.6° . (Kurnakoff, J.
 Russ. Soc. 24. 629.)

Sl. decomp. by cold, completely by boiling
 H_2O ; decomp. prevented by a little HCl .
 Pptd. from aqueous solution by alcohol, HCl ,

or sat. KCl or NaCl + Aq; not decomp. by boiling HCl + Aq. (Claudet, *l.c.*) Nearly insol. in cold, but sol. in hot H₂O, to which a few drops of HCl have been added. Less sol. in dil. HCl + Aq than luteocobaltic chloride. (Rogojski, A. ch. (3) 41. 447.)

Insol. in alcohol. (Gibbs and Genth.)

Chloropurpureocobaltic antimony chloride,
2CoCl(NH₃)₅Cl₂, SbCl₃.

Ppt. Decomp. by H₂O. (Gibbs.)

— **bismuth chloride.**

Insol. in conc. HCl. Easily decomp. by H₂O. (Gibbs.)

— **mercuric chloride,** CoCl(NH₃)₅Cl₂, 3HgCl₂.

Insol. in cold, less sol. in hot H₂O than chloropurpureocobaltic chloride. Insol. in cold fuming HCl + Aq; sl. sol. in hot HCl + Aq, separating on cooling; sl. sol. in hot aqua regia; moderately sol. in hot HNO₃ + Aq; partly sol. in cold conc. H₂SO₄, wholly on warming. Easily sol. in warm H₂C₂O₄ + Aq. Insol. in HgCl₂ + Aq.

Moderately sol. in NH₄OH + Aq or (NH₄)₂CO₃ + Aq. (Carstanjen.)

CoCl(NH₃)₅Cl₂, 2HgCl₂. Sl. sol. in cold, but much more easily in hot H₂O. (Gibbs, Proc. Am. Acad. 10. 33.)

— **chloropalladite,** CoCl(NH₃)₅Cl₂, PdCl₂.

Sl. sol. in cold, moderately sol. in hot H₂O. (Carstanjen.)

— **chloroplatinate,** CoCl(NH₃)₅Cl₂, PtCl₄.

Nearly insol. in cold. Very sl. sol. in hot H₂O. (Gibbs and Genth, Sill. Am. J. (2) 23. 319.)

— **chromate,** CoCl(NH₃)₅CrO₄.

Very sl. sol. in H₂O. (J.)

— **dichromate,** CoCl(NH₃)₅Cr₂O₇.

Much more easily sol. in H₂O than the neutral salt. (J.)

— **dithionate,** CoCl(NH₃)₅S₂O₆.

Very sl. sol. in cold, more easily in hot H₂O. (J.)

— **manganic fluoride.**

Ppt. Sl. sol. in dil. HF + Aq. (Christensen, J. pr. (2) 35. 161.)

— **fluosilicate,** CoCl(NH₃)₅SiF₆.

Very sl. sol. in HF + Aq.

— **iodide,** CoCl(NH₃)₅I₂.

Much more sol. in H₂O than bromide or chloride. Sol. in 54.5 pts. H₂O at 15.6°, and 50 pts. at 19.3°. (J.)

Chloropurpureocobaltic mercuric
CoCl(NH₃)₅I₂, 2HgI₂.

Sl. sol. in H₂O. (J.)

CoCl(NH₃)₅I₂, HgI₂. Very sl. sol. H₂O. (J.)

— **nitrate,** CoCl(NH₃)₅(NO₃)₂.

Sol. in 80 pts. H₂O at 15°. Rather sol. in hot H₂O. (Jørgensen, J. pr. 209.)

— **oxalate,** CoCl(NH₃)₅C₂O₄.

Sl. sol. in H₂O. (J.)

— **pyrophosphate,** CoCl(NH₃)₅(H₂P₂O₇).

Sl. and very slowly sol. in cold, much more easily in warm H₂O. (J.)

[CoCl(NH₃)₅]₂P₂O₇ + xH₂O. Quite sol. in H₂O.

— **diphosphopentamolybdate,**

[CoCl(NH₃)₅]₂(5MoO₃, 2HPO₄).

Ppt. Nearly insol. in pure H₂O; n in dil. H₂SO₄ + Aq without decomp.

[CoCl(NH₃)₅]₂(5MoO₃, 2NH₄PO₄)
As above.

— **sulphate,** CoCl(NH₃)₅SO₄.

Anhydrous. Slowly sol. in 128–130 H₂O at 16°.

+ 2H₂O. Sol. in 133.4 pts. H₂O. Rather easily sol. in hot H₂O, and much more rapidly than the anhydrous salt. (J.)

[CoCl(NH₃)₅]₂SO₄(SO₄H)₄. Decomposes in H₂O into neutral sulphate. Sol. in E

— **tartrate,** CoCl(NH₃)₅(C₄H₄O₆)₂ · 2½H₂O.

Moderately sol. in H₂O; insol. in al

— **thiosulphate,** CoCl(NH₃)₅S₂O₃.

Nearly insol. in cold H₂O; very sl. sol. in boiling H₂O with partial decomp. (J.)

Chloropurpureoiridium comps.

See Iridopentamine comps.

Chloropurpureorhodium carbonate

ClRh(NH₃)₅CO₃ + H₂O.

Easily sol. in H₂O. (Jørgensen.)

— **chloride,** ClRh(NH₃)₅Cl₂.

Sol. in 179 pts. H₂O at 17°, and more in hot H₂O. Sol. in conc. H₂SO₄ or NaOH + Aq without decomp. Very sl. sol. in cold dil. HCl + Aq (1 : 1). Sl. sol. in H₂O + Aq. Insol. in alcohol. (Jørgensen (2) 27. 433; 34. 394.)

— **rhodium chloride,**
3ClRh(NH₃)₅Cl₂, 2RhCl₃.

Ppt. (Jørgensen, Z. anorg. 5. 75.)

pureorhodium chloroplatinate,
(NH₄)₂PtCl₆.

1 cold H₂O. (J.)

silicate, ClRh(NH₄)₂SiF₆.

sol. in cold H₂O. Sol. in NaOH +
no salt. (J.)

oxide, ClRh(NH₄)₂(OH)₂.

only in solution. (J.)

ite, ClRh(NH₄)₂(NO₃)₂.

in cold H₂O, but more easily than
de. Sol. in boiling NaOH + Aq as
(J.)

hate, ClRh(NH₄)₂SO₄ + 2H₂O.

in cold, more easily in hot H₂O. (J.)
NH₄)₂SO₄, 3H₂SO₄. Sl. sol. in cold,
ly in hot H₂O. (J.)

iodous acid.

m chlororhodite, (NH₄)₂RhCl₅ +

er, B. 1908, 41. 213.)

H₂O; insol. in alcohol. (Wollaston.)

nable. (Leidié, A. ch. (6) 17. 275.)

RhCl₅ + 1½H₂O. Sol. in H₂O, but

than Na salt; insol. in alcohol. Sol.

I₂Cl + Aq. (Claus, J. B. 1855. 423.)

l. c.)

m chlororhodite nitrate,

(NH₄)₂Rh₂Cl₅, 2NH₄NO₃.

sol. in H₂O. Decomp. by boiling with

sol. in HNO₃ + Aq. (Leidié, C. R.

)

chlororhodite, Ba₂(RhCl₅)₂.

ples the Na salt. (Bunsen, A. 146.

chlororhodite, Cs₂RhCl₅ + H₂O.

lty sol. in H₂O. (Gutbier, B. 1908,

rorhodite, Pb₂(RhCl₅)₂.

nsol. in H₂O. (Claus.) Not obtain-
eidié.)

is chlororhodite, Hg₂RhCl₅.

nsol. in H₂O. (Claus.) Not obtain-
eidié.)

n chlororhodite, K₂RhCl₅ + H₂O.

lorescent. Sl. sol. in H₂O. Sl. sol.

Aq. (Gibbs.) Insol. or sl. sol. in

(Berzelius.)

anhydrous. (Leidié.)

ns 1H₂O. (Seubert and Kobbé, B.

)

Can be cryst. from H₂O containing a little
HCl. (Gutbier, B. 1908, 41. 212.

K₂RhCl₅ + 3H₂O. Efflorescent. Sl. sol. in
H₂O. Aqueous solution decomp. to above on
standing. (Claus.)

Not obtainable. (Leidié.)

Also obtained by Seubert and Kobbé. (B.
23. 2556.)

+1½H₂O. (Leidié, C. R. 111. 106.)

Rubidium chlororhodite, Rb₂RhCl₅ + H₂O.

Difficulty sol. in H₂O. (Gutbier, B. 1908,
41, 214.

Silver chlororhodite, Ag₂RhCl₅.

Ppt. Insol. in H₂O. (Claus.)

Not obtainable. (Leidié.)

Sodium chlororhodite, Na₂Rh₂Cl₁₁ + 18H₂O.

Na₂RhCl₅ + 9H₂O.

Efflorescent. Sol. in 1.5 pts. H₂O. Melts
in crystal H₂O at 50°. Insol. in alcohol.
(Claus.)

+12H₂O. (Gutbier, B. 1908, 41. 213.)

Chlororuthenic acid.

Ammonium chlororuthenate, (NH₄)₂RuCl₆.

Easily sol. in H₂O. (Claus.)

Formula is (NH₄)₂Ru(NO)Cl₅. (Joly, C.
R. 107. 994.)

Sol. in H₂O with decomp. Sol. in HCl.
(Howe, J. Am. Chem. Soc. 1904, 26. 549.)

Ammonium aquochlororuthenate,

(NH₄)₂Ru(H₂O)Cl₅.

Ppt. (Howe, J. Am. Chem. Soc. 1904, 26.
548.)

Cæsium chlororuthenate, Cs₂RuCl₆.

Sl. sol. in H₂O. Sol. in hot dil. HCl + Aq.
(Howe, J. Am. Chem. Soc. 1901, 23. 784.)

Potassium chlororuthenate, K₂RuCl₆.

Very sol. in H₂O. Very sl. sol. in conc.
NH₄Cl + Aq. Insol. in 70% alcohol. (Claus.)

Formula is K₂Ru(NO)Cl₅. (Joly.)

Very sl. sol. in cold H₂O. Insol. in pres-
ence of KCl.

Partially decomp. in hot aqueous solution.
(Antony, Gazz. ch. it. 1899, 29. (2), 82.)

Easily sol. in H₂O with rapid decomp.
Sol. in HCl. (Howe, J. Am. Chem. Soc.
1904, 26. 548.)

Potassium aquochlororuthenate,

K₂Ru(OH₂)Cl₅.

Sol. in H₂O. (Howe, J. Am. Chem. Soc.
1904, 26. 547.)

Rubidium chlororuthenate, Rb₂RuCl₆.

Sl. sol. in H₂O; sol. in hot dil. HCl + Aq.
(Howe, J. Am. Chem. Soc. 1901, 23. 784.)

Chlororuthenious acid.

Ammonium chlororuthenite, $(\text{NH}_4)_4\text{Ru}_2\text{Cl}_{10}$.
Sl. sol. in H_2O . Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$ or alcohol. (Claus, J. pr. 80. 282.)

Cæsium chlororuthenite, $\text{Cs}_2\text{RuCl}_6 + \text{H}_2\text{O}$.
Sl. sol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$. (Howe, J. Am. Chem. Soc. 1901, 23. 785.)

Potassium chlororuthenite, $\text{K}_4\text{Ru}_2\text{Cl}_{10}$.
Moderately sol. in cold, more easily in hot H_2O . Decomp. easily by heating. Insol. in conc. $\text{NH}_4\text{Cl} + \text{Aq}$. Insol. in 80% alcohol.

Rubidium chlororuthenite, $\text{Rb}_2\text{RuCl}_6 + \text{H}_2\text{O}$.
Sl. sol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$. (Howe, J. Am. Chem. Soc. 1901, 23. 786.)

Sodium chlororuthenite, $\text{Na}_4\text{Ru}_2\text{Cl}_{10}$.
Deliquescent. Sol. in H_2O or alcohol.

Trichlorosilicomercaptane.

See Silicon chlorohydrosulphide.

Chlorosmic acid.

Ammonium chlorosmate, $(\text{NH}_4)_2\text{OsCl}_4$.
Sl. sol. in H_2O . Insol. in alcohol and H_2O containing HCl .

Potassium chlorosmate, K_2OsCl_4 .
Properties as the NH_4 salt.

Potassium amino chlorosmate,
 $(\text{NH}_2)\text{OsCl}_3 \cdot 2\text{KCl}$.
Ppt. (Brizard, A. ch. 1900, (7) 21. 375.)

Potassium amino, chlorosmate hydrogen chloride, $(\text{NH}_2)\text{OsCl}_3 \cdot 2\text{KCl} \cdot \text{HCl}$.
Ppt. (Brizard, A. ch. 1900, (7) 21. 378.)

Silver chlorosmate, Ag_2OsCl_6 .
Insol. in H_2O or $\text{HNO}_3 + \text{Aq}$. (Claus and Jacoby.)

Silver chlorosmate ammonia, $\text{Ag}_2\text{OsCl}_6 \cdot 2\text{NH}_3$.
Sol. in much H_2O . Sl. sol. in $\text{KOH} + \text{Aq}$. Easily sol. in $\text{KCN} + \text{Aq}$. (C. and J.)

Sodium chlorosmate, $\text{Na}_2\text{OsCl}_6 + 2\text{H}_2\text{O}$.
Easily sol. in H_2O or alcohol.

Chlorosmious acid.

Ammonium chlorosmite,
 $(\text{NH}_4)_4(\text{Os}_2\text{Cl}_{10}) + 3\text{H}_2\text{O}$.
Easily sol. in H_2O and alcohol; insol. in ether. (Claus and Jacoby, J. pr. 90. 65.)

Potassium chlorosmite, $\text{K}_6\text{Os}_2\text{Cl}_{12} + 6\text{H}_2\text{O}$.
Very easily sol. in H_2O or alcohol. Insol. in ether. (C. and J.)

Chlorosmisulphurous acid.

Potassium hydrogen chlorosmisulphite,
 $\text{OsCl}_4(\text{SO}_3)_2 \cdot \text{K}_2\text{H}_2$.
Ppt. (Rosenheim, Z. anorg. 1900, 24. 422.)

Sodium chlorosmisulphite,
 $\text{OsCl}_4(\text{SO}_3)_2 \cdot \text{Na}_2 + 10\text{H}_2\text{O}$.
Ppt. (Rosenheim, Z. anorg. 1900, 24. 420.)

Chloropyroselenious acid.

Ammonium chloropyroselenite, $\text{NH}_4\text{Cl} \cdot 2\text{SeO}_2 + 2\text{H}_2\text{O}$.
Sol. in H_2O . (Muthmann and Schäfer, B. 26. 1008.)

Potassium chloropyroselenite, $\text{KCl} \cdot 2\text{SeO}_2 + \text{H}_2\text{O}$.
As NH_4 salt. (M. and S.)

Rubidium chloropyroselenite, $\text{RbCl} \cdot 2\text{SeO}_2 + 2\text{H}_2\text{O}$.
As NH_4 salt. (M. and S.)

Chlorostannic acid, $\text{SnO}(\text{OH})\text{Cl}$.
(Mallet, Chem. Soc. 35. 524.)
 $\text{H}_2\text{SnCl}_6 + 6\text{H}_2\text{O}$. Extremely deliquescent; sol. in H_2O . (Seubert, B. 20. 793.)

Ammonium chlorostannate, $(\text{NH}_4)_2\text{SnCl}_6$ (pink salt).
Sol. in 3 pts. H_2O at 14.5° . Solution decomp. on boiling when dilute, but not when conc. (Bolley.)

Barium chlorostannate, $\text{BaSnCl}_6 + 5\text{H}_2\text{O}$.
Sol. in H_2O . (Lewy, A. ch. (3) 16. 308.)

Cæsium chlorostannate, Cs_2SnCl_6 .
Nearly insol. in conc. $\text{HCl} + \text{Aq}$. (Sharples, Sill. Am. J. (2) 47. 178.)

Calcium chlorostannate, $\text{CaSnCl}_6 + 5\text{H}_2\text{O}$.
Very deliquescent. (Lewy, A. ch. (3) 16. 308.)

Cerium chlorostannate, $\text{CeSnCl}_6 + 9\text{H}_2\text{O}$.
Deliquescent. Sol. in H_2O . (Cleve, Bull. Soc. (2) 31. 197.)

Cobalt chlorostannate, $\text{CoSnCl}_6 + 6\text{H}_2\text{O}$.
Sol. in H_2O . (Jørgensen.)

Didymium chlorostannate, $\text{DiCl}_2 \cdot \text{SnCl}_6 + 10\frac{1}{2}\text{H}_2\text{O}$.
Sol. in H_2O . (Cleve.)

Glucinum chlorostannate, $\text{GlSnCl}_6 + 8\text{H}_2\text{O}$.
Deliquescent. Sol. in H_2O . (Atterberg, Sv. V. A. Handl. 12. No. 4. 14.)

Lanthanum chlorostannate, $4\text{LaCl}_3, 5\text{SnCl}_4 + 45\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . (Cleve.)

Lithium chlorostannate, $\text{Li}_2\text{SnCl}_6 + 8\text{H}_2\text{O}$.

Sol. in little H_2O without decomp., but decomp. by dilution. (Chassevant, A. ch. (6) 80. 42.)

Magnesium chlorostannate, $\text{MgSnCl}_4 + 6\text{H}_2\text{O}$.

Very deliquescent. (Lewy.)

Manganous chlorostannate, $\text{MnSnCl}_4 + 6\text{H}_2\text{O}$.

Deliquescent in moist, efflorescent in dry air. (Jørgensen.)

Nickel chlorostannate, $\text{NiSnCl}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Jørgensen.)

Potassium chlorostannate, K_2SnCl_6 .

Sol. in H_2O .

Sodium chlorostannate, $\text{Na}_2\text{SnCl}_6 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Topsoë, Gm. K. Handb. 6^{te} Aufl. III. 149.)

Strontium chlorostannate, $\text{SrSnCl}_4 + 8\text{H}_2\text{O}$.

Sl. deliquescent, and easily sol. in H_2O . (Topsoë.)

Yttrium chlorostannate, $\text{YCl}_3, \text{SnCl}_4 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve, Bull. Soc. (2) 31.197.)

Zinc chlorostannate, $\text{ZnSnCl}_4 + 6\text{H}_2\text{O}$.

(Biron, C. C. 1904, II. 410.)

Chlorosulphobismuthous acid.

Cuprous chlorosulphobismuthite, $2\text{Cu}_2\text{S}, \text{Bi}_2\text{S}_3, 2\text{BiSCl}$.

Stable in air and insol. in H_2O at ord. temp.

Decomp. by boiling H_2O .

Decomp. by mineral acids with evolution of H_2S . (Ducatte, C. R. 1902, 134. 1212.)

Lead chlorosulphobismuthite, $\text{PbS}, \text{Bi}_2\text{S}_3, 2\text{BiSCl}$.

Stable in the air. Insol. in H_2O ; decomp. by boiling H_2O ; sol. in dil. acids with decomp. and evolution of H_2S . (Ducatte.)

Chlorosulphonic acid, HClSO_3 .

See Sulphuryl hydroxyl chloride.

Chloropyrosulphonic acid.

Ammonium chloropyrosulphonate, $\text{ClS}_2\text{O}_6\text{NH}_4$.

Fumes in the air.

Decomp. by H_2O and alcohol. (Traube, B. 1913, 48. 2519.)

Sodium chloropyrosulphonate, $\text{ClS}_2\text{O}_6\text{Na}$.

Fumes in the air.

Decomp. by H_2O and alcohol. (Traube.)

Chlorosulphuric acid, HSO_3Cl .

See Sulphuryl hydroxyl chloride.

SO_2Cl_2 . See Sulphuryl chloride.

Aluminum chlorosulphate, $\text{Al}(\text{SO}_4)\text{Cl} + 6\text{H}_2\text{O}$.

Very sol. in H_2O . Nearly insol. in abs. alcohol. (Recoura, Bull. Soc. 1902, (3) 27. 1155.)

Chromium chlorosulphate, $\text{CrClSO}_4 + 5\text{H}_2\text{O}$.

Green. (Weinland, Z. anorg. 1905, 48. 253.) (Recoura, C. R. 1902, 135. 164.)

Violet. (Weinland, Z. anorg. 1905, 48. 254.)

Very sol. in H_2O . Insol. in a mixture of alcohol and acetone. (Recoura, C. R. 1902, 135. 164.)

+ $8\text{H}_2\text{O}$. Two isomeric modifications:

(a) *Green needles*. Easily sol. in H_2O . (Weinland, Z. anorg. 1906, 48. 251.)

(b) *Violet plates*. Easily sol. in H_2O . (Weinland.)

Chlorosulphurous acid.

Ammonium palladious trichlorosulphite, $(\text{NH}_4)_2\text{PdCl}_3\text{SO}_3 + \text{H}_2\text{O}$.

Easily sol. in H_2O . (Rosenheim, Z. anorg. 1900, 23. 30.)

Chlorotelluric acid.

Ammonium chlorotellurate, $(\text{NH}_4)_2\text{TeCl}_6$.

Sol. without decomp. in a small amt. of H_2O , but decomp. by much H_2O or alcohol.

Cæsium chlorotellurate, Cs_2TeCl_6 .

Decomp. by H_2O . Sol. in dil. $\text{HCl} + \text{Aq}$. 100 pts. $\text{HCl} + \text{Aq}$ (sp. gr. 1.2) dissolve 0.05 pt. at 22° .

100 pts. $\text{HCl} + \text{Aq}$ (sp. gr. 1.05) dissolve 0.78 pt. at 22° .

Insol. in alcohol. (Wheeler, Sill. Am. J. 145. 267.)

Potassium chlorotellurate, K_2TeCl_6 .

Deliquescent; decomp. by H_2O and absolute alcohol. (Berzelius.)

The most sol. in H_2O of the chloro- or bromo-tellurates. Easily sol. in dil. $\text{HCl} + \text{Aq}$; conc. $\text{HCl} + \text{Aq}$ ppts. KCl . (Wheeler, Sill. Am. J. 145. 267.)

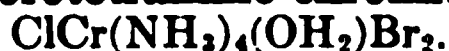
Rubidium chlorotellurate, Rb_2TeCl_6 .

Decomp. by H_2O . Much more sol. in dil. $\text{HCl} + \text{Aq}$ than Cs_2TeCl_6 .

100 pts. $\text{HCl} + \text{Aq}$ (sp. gr. 1.2) dissolve 0.34 pt. at 22° .

100 pts. $\text{HCl} + \text{Aq}$ (sp. gr. 1.05) dissolve, 13.99 pts. at 22° .

Sl. sol. in alcohol. (Wheeler.)

Chlorotetramine chromium bromide,

Very easily sol. in H_2O . (Cleve, 1861, (Jørgensen, J. pr. (2) 42. 210.)

— **chloride**, $\text{ClCr}(\text{NH}_3)_4(\text{OH}_2)\text{Cl}_2$.

Sol. in H_2O , but decomp. by boiling. Sol. in $\text{HCl} + \text{Aq}$, and this solution may be boiled without decomp. (Cleve.)

Sol. in 15.7 pts. H_2O at 15° . (Jørgensen, J. pr. 42. 208.)

— **chromate**, $\text{ClCr}(\text{NH}_3)_4(\text{OH}_2)\text{CrO}_4$.

Precipitate. (Cleve.)

— **fluosilicate**, $\text{ClCr}(\text{NH}_3)_4(\text{OH}_2)\text{SiF}_6$.

Sl. sol. in H_2O . (Jørgensen, J. pr. (2) 42. 218.)

— **hydroxide**, $\text{ClCr}(\text{NH}_3)_4(\text{OH})_2$.

Known only in solution. (Cleve.)

— **iodide**, $\text{ClCr}(\text{NH}_3)_4(\text{OH}_2)\text{I}_2$.

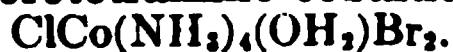
Easily sol. in H_2O . (Cleve.)

— **nitrate**, $\text{ClCr}(\text{NH}_3)_4(\text{OH}_2)(\text{NO}_3)_2$.

Very easily sol. in H_2O . (Cleve); (Jørgensen, J. pr. (2) 42. 209.)

— **sulphate**, $\text{ClCr}(\text{NH}_3)_4(\text{OH}_2)\text{SO}_4$.

Very difficulty sol. in cold, more easily in hot H_2O . (Cleve.)

Chlorotetramine cobaltic bromide,

More sol. in H_2O than chloride. Nearly insol. in $\text{HBr} + \text{Aq}$ (1:1). (Jørgensen, J. pr. (2) 42. 215.)

— **chloride**, $\text{ClCo}(\text{NH}_3)_4(\text{OH}_2)\text{Cl}_2$.

Sol. in about 40 pts. H_2O , and is identical with octamine cobaltic purpleochloride of Vortmann. (Jørgensen, J. pr. (2) 42. 211.)

— **chloroplatinate**, $\text{ClCo}(\text{NH}_3)_4(\text{OH}_2)\text{PtCl}_6 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Jørgensen.)

— **chromate**, $\text{ClCo}(\text{NH}_3)_4(\text{OH}_2)\text{CrO}_4$.

Easily sol. in cold H_2O . (Jørgensen, J. pr. (2) 42. 216.)

— **fluosilicate**, $\text{ClCo}(\text{NH}_3)_4(\text{OH}_2)\text{SiF}_6$.

Sl. sol. in H_2O . Nearly insol. in $\text{H}_2\text{SiF}_6 + \text{Aq}$. (Jørgensen, J. pr. (2) 42. 219.)

— **sulphate**, $\text{ClCo}(\text{NH}_3)_4(\text{OH}_2)\text{SO}_4$.

Sol. in H_2O . (Jørgensen, J. pr. (2) 42. 214.)

Chlorotitanic acid, $\text{TiCl}_4 \cdot 2\text{HCl} = \text{H}_2\text{TiCl}_6$.

Known only in solution. (Kowalewsky, Z. anorg. 1900, 25. 192.)

Chlorous acid, HClO_2 .

Known only in aqueous solution. 100 g. H_2O at 8.5° and 753 mm. pressure dissolve 4.7 g. Cl_2O_3 . Hydrate with 50.07–67.43% H_2O , perhaps $\text{HClO}_2 + \text{H}_2\text{O}$, separates out at 0° . (Brandan, A. 151. 340.)

Pure HClO_2 is not known even in solution. (Garzarolli-Thurnlakh, A. 209. 184.)

Chlorites.

All chlorites are easily sol. in H_2O and alcohol, with gradual decomp.

Ammonium chlorite.

Known only in aqueous solution, which decomposes on evaporation or long standing.

Barium chlorite, $\text{Ba}(\text{ClO}_2)_2$.

Deliquescent; easily sol. in H_2O . Solution decomp. on evaporation. Easily sol. in alcohol. (Millon, A. ch. (3) 7. 298.)

Lead chlorite, $\text{Pb}(\text{ClO}_2)_2$.

Nearly insol. in cold H_2O , and only sl. sol. in hot H_2O . Sol. in $\text{KOH} + \text{Aq}$. (Garzarolli and Hayn, A. 209. 203.)

Lead chlorite chloride, $6\text{Pb}(\text{ClO}_2)_2 \cdot 4\text{PbCl}_2 \cdot \text{PbO}$.

Rather difficulty sol. in H_2O . (Schiel, A. 109. 317.)

Potassium chlorite, KClO_2 .

Very deliquescent and sol. in H_2O . Sol. in alcohol of 38° . (Millon, A. ch. (3) 7. 323.) Sol. in $\text{HClO}_2 + \text{Aq}$.

Silver chlorite, AgClO_2 .

Sol. in hot, less in cold H_2O . Easily decomp. by heating above 100° . Decomp. by weakest acids. (Millon, A. ch. (3) 7. 329.)

Sodium chlorite, NaClO_2 .

Very deliquescent, and sol. in H_2O .

Strontium chlorite, $\text{Sr}(\text{ClO}_2)_2$.

Deliquescent and sol. in H_2O . Decomp. by slow evaporation. (Millon, A. ch. (3) 7. 327.)

Chloroxyfulminoplatinum,

Insol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$. (v. Meyer, J. pr. (2) 18. 305.)

Chloruranic acid, $\text{HUO}_2\text{Cl} + 2\text{H}_2\text{O}$.

Sol. in H_2O ; sl. sol. in alcohol. (Mylus, B. 1901, 34. 2776.)

Chromacichloride, CrO_2Cl_2 .

See Chromyl chloride.

Chromatiodic acid.

See Chromiodic acid.

Chromic acid, H_2CrO_4 .

Very sol. in H_2O . (Moissan, C. R. 98. 51.)

Does not exist except in solution. (Field, Chem. Soc. 61. 405.)

The composition of the hydrates formed by H_2CrO_4 at different dilutions is calculated from determinations of the lowering of the pt. produced by H_2CrO_4 and of the conductivity and sp. gr. of $H_2CrO_4 + Aq$. (Jones, Am. Ch. J. 1905, 34. 333.)

See also Chromium trioxide.

Chromates.

Chromates of the alkali metals and of Ca, Mg, and Sr are sol. in H_2O ; the others are generally insol. or sl. sol. in H_2O , but sol. in $HNO_3 + Aq$.

Aluminum chromate, basic, $Al_2O_3, CrO_3 + 7H_2O$.

Easily sol. in $NH_4OH + Aq$, alum, or acetic acid + Aq. Insol. in $NH_4Cl + Aq$. (Farrie, Chem. Soc. 4. 300.)

Insol. as such as H_2O , but easily decomp. into H_2CrO_4 and a basic insol. comp. Sol. in alkaline solutions and acids. Decomp. by many salts. (Eliot and Storer, Proc. Am. Acad. 5. 214.)

Aluminum sodium chromate silicate, $4Al_2O_3, 5Na_2O, CrO_3, 7SiO_2$.

(Weyberg, C. B. Miner, 1904. 727.)

Ammonium chromate, basic, $5(NH_4)_2O, 4CrO_3(?)$.

Easily sol. in cold H_2O . (Pohl, W. A. B. 6. 592.)

Ammonium chromate, $(NH_4)_2CrO_4$.

Very sol. in H_2O ; pptd. from aqueous solution by alcohol. (Malaguti and Sarzeau.) 100 g. H_2O dissolve 40.46 g. at 30° . (Schreinemakers, Chem. Weekbl. 1905, 1. 395.)

Sol. in H_2O without decomp. (Schreinemakers, C. C. 1905, II. 1067.)

Sp. gr. of $(NH_4)_2CrO_4 + Aq$ at $t^\circ/4^\circ$.

t°	13°	13.7°	19.6°
% $(NH_4)_2CrO_4$	10.52	19.75	28.04
Sp. gr.	1.0633	1.1197	1.1727

(Slotte, W. Ann. 1881, 14. 18.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.)

Difficulty sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Ammonium dichromate, $(NH_4)_2Cr_2O_7$.

Less sol. in H_2O than $(NH_4)_2CrO_4$. (Moser.)

100 g. H_2O dissolve 47.17g at 30° .

(Schreinemakers, Chem. Weekbl. 1905, 1. 395.)

Sp. gr. of $(NH_4)_2Cr_2O_7 + Aq$ at $t^\circ/4^\circ$.

t°	12°	10.5°	12°
% $(NH_4)_2Cr_2O_7$	6.85	13.00	19.93
Sp. gr.	1.0393	1.0782	1.1258

(Slotte, W. Ann. 1881, 14. 18.)

Sol. in alcohol. (Ranitzer, Zeit. angew. ch. 1913, 26. 456.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in acetone. (Naumann, B. 1904, 37. 4328.)

Ammonium trichromate, $(NH_4)_2Cr_3O_{10}$.

Not deliquescent, but very sol. in H_2O . (Siewert.)

Decomp. by H_2O into chromic acid and dichromate. (Jäger and Krüss, B. 22. 2036.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Ammonium tetrachromate, $(NH_4)_2Cr_4O_{13}$.

Deliquescent. Decomp. by H_2O . (Jäger and Krüss, B. 22. 2037.)

Ammonium hexachromate, $(NH_4)_2Cr_6O_{19} + 10H_2O(?)$

Very efflorescent. (Rammelsberg, Pogg. 94. 516.)

Ammonium barium chromate, $BaCrO_4, (NH_4)_2CrO_4$.

Ppt. Decomp. by H_2O . (Gröger, Z. anorg. 1908, 58. 414.)

Ammonium cadmium chromate, $(NH_4)_2O, 4CdO, 4CrO_3 + 3H_2O$.

Ppt. Decomp. by boiling H_2O . (Gröger, M. 1904, 25. 533.)

Ammonium cadmium chromate ammonia, $(NH_4)_2CrO_4, CdCrO_4, \frac{1}{2}NH_3 + 1\frac{1}{2}H_2O$.

Decomp. by H_2O . (Gröger, Z. anorg. 1908, 58. 418.)

$(NH_4)_2Cd(CrO_4)_2, 2NH_3$. Insol. in cold, decomp. by hot H_2O .

Sol. in dil. acids or in $NH_4OH + Aq$. (Briggs, Chem. Soc. 1903, 83. 395.)

Ammonium chromous chromate(?),

$(NH_4)_2CrO_4, CrCrO_4 = (NH_4)_2Cr(CrO_4)_2$.

Difficultly sol. in H_2O . Insol. in alcohol, ether, chloroform, or glacial acetic acid. Easily sol. in conc. acids, from which it is separated on dilution. Decomp. by $NaOH + Aq$. (Heintze, J. pr. (2) 4. 220.)

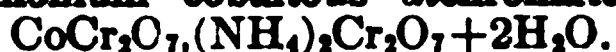
Ammonium chromyl chromate, $(3(NH_4)_2O, 2CrO_2, 3CrO_3$.

Nearly insol. in H_2O . (Pascal, C. R. 1909, 148. 1465.)

Ammonium cobaltous chromate.

Ppt. Easily decomp. (Briggs, Z. anorg. 1907, 56. 247.)

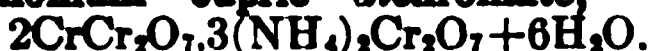
$(\text{NH}_4)_2\text{O}$, 4CoO , $4\text{CrO}_3 + 3\text{H}_2\text{O}$. Insol. in H_2O . Sol. in cold dil. H_2SO_4 . (Gröger, Z. anorg. 1906, 49. 202.)

Ammonium cobaltous dichromate,

Sl. hygroscopic; sol. in H_2O , insol. in alcohol. (Krüss, Z. anorg. 1895, 8. 454.)

Ammonium cobaltous chromate ammonia,

Ppt.; decomp. by H_2O . (Gröger, Z. anorg. 1908, 58. 422.)

Ammonium cupric dichromate,

Sol. in H_2O . (Krüss, Z. anorg. 1895, 8. 455.)

Ammonium cupric chromate ammonia,

Decomp. by H_2O . (Gröger, Z. anorg. 1908, 58. 420.)

Insol. in cold, decomp. by hot H_2O . Sol. in dil. acids or in $\text{NH}_4\text{OH} + \text{Aq}$. (Briggs, Chem. Soc. 1903, 83. 394.)

Ammonium iron (ferric) chromate,

More easily decomp. by H_2O than K_2CrO_4 , $\text{Fe}_2(\text{CrO}_4)_3 + 4\text{H}_2\text{O}$. (Hensgen, B. 12. 1300.)

6CrO_3 , $5\text{Fe}_2\text{O}_3$, $6(\text{NH}_4)_2\text{O}$, and 4CrO_3 , $\text{Fe}_2\text{O}_3, (\text{NH}_4)_2\text{O} + 4\text{H}_2\text{O}$. Ppts. (Lepierre, C. R. 1894, 119. 1217.)

Ammonium lithium chromate, $\text{NH}_4\text{LiCrO}_4 + 2\text{H}_2\text{O}$.

Not deliquescent. (Rammelsberg.)

Ammonium lead chromate, $(\text{NH}_4)_2\text{CrO}_4$, PbCrO_4 .

Ppt. Decomp. by H_2O . (Gröger, Z. anorg. 1908, 58. 424.)

Ammonium magnesium chromate,

Much more sol. in H_2O than the corresponding sulphate. (v. Hauer.)

Sol. in H_2O . (Gröger, Z. anorg. 1908, 58. 416.)

Ammonium manganous chromate,

Sol. in H_2O . (Hensgen, R. t. c. 3. 433.)

Ammonium nickel chromate, $(\text{NH}_4)_2\text{CrO}_4$, $\text{NiCrO}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Gröger, Z. anorg. 1906, 51. 353.)

Can be cryst. from H_2O under 40° . (Chem. Soc. 1903, 83. 392.)

Ammonium nickel chromate amm

Decomp. by H_2O . (Gröger, Z. 1906, 51. 354.)

$(\text{NH}_4)_2\text{Ni}(\text{CrO}_4)_2$, 2NH_3 . Insol. in H_2O . Decomp. by hot H_2O . Sol. acids or in $\text{NH}_4\text{OH} + \text{Aq}$. (Briggs, Soc. 1903, 83. 393.)

$+6\text{H}_2\text{O}$. (Briggs, Proc. Chem. Soc. 18. 254.)

Ammonium potassium chromate, NH_4

Sol. in H_2O . (E. Kopp, C. N. 1 $+ \text{H}_2\text{O}$. (Étard, C. R. 85. 443.)

$2(\text{NH}_4)_2\text{CrO}_4$, $3\text{K}_2\text{CrO}_4$. Very sol. in (Zehenter, M. 1897, 18. 51.)

Ammonium silver chromate, $(\text{NH}_4$ 

Decomp. by H_2O . (Gröger, Z. 1908, 58. 423.)

Ammonium sodium chromate, NH_4Na 

Very sol. in H_2O . (Zehenter, M. 18 54.)

Ammonium strontium chromate, $(\text{NH}_4$ 

Ppt. Decomp. by H_2O . (Gröger, anorg. 1908, 58. 415.)

Ammonium uranyl chromate, $(\text{NH}_4$ 

Decomp. by boiling with H_2O . acidulated H_2O . (Formánek, A. 257 $+3\text{H}_2\text{O}$. (Formánek.)

Ammonium zinc chromate, $(\text{NH}_4)_2\text{O}$, $2\text{CrO}_3 + \text{H}_2\text{O}$.

Decomp. by hot H_2O . (Gröger, M. 25. 520.)

Ammonium zinc chromate ammon

Insol. in cold, decomp. by hot H_2O in dil. acids or in $\text{NH}_4\text{OH} + \text{Aq}$. (Chem. Soc. 1903, 83. 394.)

4ZnCrO_4 , $2(\text{NH}_4)_2\text{CrO}_4$, $3\text{NH}_3 + 3\text{H}_2\text{O}$. Ppt. Decomp. by H_2O . (Gröger, Z. 1908, 58. 416.)

Ammonium dichromate chloride m chloride, $(\text{NH}_4)_2\text{Cr}_2\text{O}_7, 2\text{NH}_4\text{Cl}, 4\text{H}_2\text{O}$.

Ppt. Sol. in cold, more sol. in warm (Strömholm, Z. anorg. 1912, 75. 280.)

Ammonium dichromate chloride m cyanide, $(\text{NH}_4)_2\text{Cr}_2\text{O}_7$, $4\text{NH}_4\text{CN}$, $6\text{Hg}(\text{CN})_2 + 4\text{H}_2\text{O}$.

(Strömholm, Z. anorg. 1913, 80. 15)

um chromate chromyl fluoride,
 $(\text{CrO}_4)_2\text{CrO}_4, \text{CrO}_4\text{F}_2$.

H_2O . (Varenne, C. R. 91. 989.)

um chromate iodate.

romoiodate, ammonium.

um dichromate mercuric chloride,
 $(\text{Cr}_2\text{O}_7)_2\text{Cr}_2\text{O}_7, \text{HgCl}_2$.

be recryst. from H_2O or $\text{HgCl}_2 + \text{Aq}$,
 $(\text{NH}_4)_2\text{Cr}_2\text{O}_7 + \text{Aq}$. (Jäger and
 22. 2044.)

(Richmond and Abel, Chem. Soc.
 99.)

be made to crystallize with H_2O .
 d Krüss.)

$(\text{Cr}_2\text{O}_7)_2\text{Cr}_2\text{O}_7, \text{HgCl}_2$. Decomp. by H_2O .
 K.)

$(\text{Cr}_2\text{O}_7)_2\text{Cr}_2\text{O}_7, \text{HgCl}_2$. Decomp. by H_2O .
 ..)

$\text{Cr}_2\text{O}_7, 3\text{HgCl}_2$. (J. and K.)

$\text{Cr}_2\text{O}_7, 4\text{HgCl}_2$. (J. and K.)

um chromate phosphate.

osphochromate, ammonium.

um chromate tellurate.

romotellurate, ammonium.

chromate, BaCrO_4 .

ely sl. sol. in H_2O .

ted from electrical conductivity of
 Aq , 1 l. H_2O dissolves 3.8 mg.
 at 18° . (Kohlrausch and Rose, Z.
 12. 241.)

not ignited, BaCrO_4 is sol. in 86,957
 3; 22,988 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (0.5%
 3670 pts. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ (5%
); 1986 pts. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ (10%
); 1813 pts. $\text{H}_2\text{CrO}_4 + \text{Aq}$ (10%
) When ignited, 160,000 pts. H_2O are
 for solution. (Schweitzer, by Fre-
 anal. 29. 414.)

23,000 pts. boiling H_2O . (Mescher-
 anal. 21. 399.)

BaCrO_4 are dissolved in 1 l. of sat.
 at 18° . (Kohlrausch, Z. phys. Ch.
 168.)

sol. in HNO_3, HCl , or chromic acid +
 which it is precipitated by NH_4OH ,
 tion with H_2O . (Bahr.)

n $\text{K}_2\text{Cr}_2\text{O}_7 + \text{Aq}$. (Schweitzer.)

49,381 pts. $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$ (0.75
 ; 15° ; in 23,355 pts. $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$
 lt) at 15° ; in 45,162 pts. NH_4NO_3
 % salt) at 15° . (Fresenius, Z. anal.

sol. in alkali tartrates, or citrates +
 ischer, J. pr. (2) 5. 326.)

10^{-4} g. equiv. BaCrO_4 are dissolved
 15% alcohol at ord. temp. (Guerini,
 912.)

n acetic acid and in $\text{M}_2\text{Cr}_2\text{O}_7 + \text{Aq}$.

sol. in a mixture of the two, except

in presence of $\text{MC}_2\text{H}_3\text{O}_2$. (Caron and Raquet,
 Bull. Soc. 1906, (3) 35. 1064.)

Not completely insol. in acetic acid. (Bau-
 bigny, Bull. Soc. 1907, (4) 1. 58.)

Insol. in acetone. (Naumann, B. 1904,
 37. 4329.)

Insol. in methyl acetate. (Naumann, B.
 1909, 42. 3790.)

Barium dichromate, $\text{BaCr}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Decomp. by H_2O with separation of
 BaCrO_4 . Sol. in $\text{H}_2\text{CrO}_4 + \text{Aq}$. (Bahr, J. B.
 1853. 358.)

Sol. in cold H_2O with formation of BaCrO_4
 and CrO_3 .

Insol. in glacial acetic acid. (Mayer, B.
 1903, 36. 1742.)

Barium calcium chromate, $\text{BaCa}(\text{CrO}_4)_2$.

(Bourgeois Bull. Soc. Min. 1879, 2. 124.)

Barium potassium chromate, $\text{BaK}_2(\text{CrO}_4)_2$.

Decomp. by H_2O . (Gröger, Z. anorg.
 1907, 54. 186.)

Decomp. by H_2O . Stable in $\text{K}_2\text{CrO}_4 + \text{Aq}$,
 containing:

2.181	pts.	K_2CrO_4	per	100	pts.	H_2O	at	11.5°
3.395	"	"	"	"	"	"	"	27.5°
5.120	"	"	"	"	"	"	"	50.0°
7.119	"	"	"	"	"	"	"	76.0°
9.036	"	"	"	"	"	"	"	100.0°

(Barre, C. R. 1914, 158. 497.)

Barium potassium trichromate,

$\text{Ba}_2\text{K}_2(\text{Cr}_2\text{O}_{10})_3 + 3\text{H}_2\text{O}$.

Extremely deliquescent. (Bahr.)

Bismuth chromates, basic.

These comps. are insol. in H_2O even in
 presence of H_2CrO_4 ; sol. in HCl or $\text{HNO}_3 +$
 Aq . (Löwe, J. pr. 67. 288.)

100 pts. H_2O dissolve 0.00008 pt. "bis-
 muth chromate"; 100 pts. acetic acid dis-
 solve 0.00021 pt. "bismuth chromate"; 100
 pts. $\text{HNO}_3 + \text{Aq}$ (sp. gr. = 1.038) dissolve
 0.00024 pt. "bismuth chromate"; 100 pts.
 $\text{KOH} + \text{Aq}$ (sp. gr. = 1.33) dissolve 0.00016
 pt. "bismuth chromate." (Pearson, Phil.
 Mag. (4) 11. 206.)

Not insol. in dil. $\text{HNO}_3 + \text{Aq}$ unless K_2CrO_4
 is present. Less sol. in hot $\text{NaOH} + \text{Aq}$ than
 PbCrO_4 . (Storer.)

"Bismuth chromate" is insol. in acetone.
 (Naumann, B. 1904, 37. 4329.)

$3\text{Bi}_2\text{O}_3, 2\text{CrO}_3 = 2(\text{BiO})_2\text{CrO}_4, \text{Bi}_2\text{O}_3$. Insol.
 in H_2O ; sol. in $\text{HNO}_3 + \text{Aq}$.

$\text{Bi}_2\text{O}_3, \text{CrO}_3 = (\text{BiO})_2\text{CrO}_4$. Insol. in H_2O ;
 easily sol. in dil. $\text{HCl} + \text{Aq}$, less in dil. HNO_3
 or $\text{H}_2\text{SO}_4 + \text{Aq}$. (Muir.)

$\text{Bi}_2\text{O}_3, 2\text{CrO}_3 = (\text{BiO})_2\text{Cr}_2\text{O}_7$. Insol. in
 H_2O .

+ H_2O .

$5\text{Bi}_2\text{O}_3, 11\text{CrO}_3 + 6\text{H}_2\text{O}$. (Muir, Chem.
 Soc. 31. 24.)

$3\text{Bi}_2\text{O}_3, 7\text{CrO}_3$. Insol. in H_2O ; easily sol. in mineral acids, especially $\text{HCl} + \text{Aq}$. Partly sol. in $\text{KOH} + \text{Aq}$.

Bismuth chromate, acid, $\text{Bi}_2\text{O}_3, 4\text{CrO}_3 + \text{H}_2\text{O}$.

Insol. in hot or cold H_2O . Sol. in dil. HCl or $\text{HNO}_3 + \text{Aq}$. (Muir, Chem. Soc. **30**. 17.)

Bismuth potassium chromate, $\text{Bi}_2(\text{CrO}_4)_3, \text{K}_2\text{CrO}_4$.

Insol. in H_2O . Decomp. with hot H_2O . $\text{Bi}_2\text{O}_3, \text{K}_2\text{O}, 6\text{CrO}_3 + \text{H}_2\text{O}$. (Preis and Raymann, J. B. **1880**. 336.)

Bromomolybdenum chromate.
(Atterberg.)

Cadmium chromate, basic, $2\text{CdO}, \text{CrO}_3 + \text{H}_2\text{O}$.

Very sl. sol. in H_2O ; very slowly sol. in $\text{NH}_4\text{OH} + \text{Aq}$ with combination. (Malaguti and Sarzeau, A. ch. (3) **9**. 431.)

Composition as above. (Freese, B. **2**. 478.)

Cadmium chromate, CdCrO_4 .

Insol. in H_2O ; sol. in acids; decomp. by heating with H_2O . (Schulz, Z. anorg. **1895**, **10**. 153.)

Sol. in hot conc. $\text{CdSO}_4 + \text{Aq}$. (Briggs, Z. anorg. **1907**, **56**. 253.)

$+ 2\text{H}_2\text{O}$. Decomp. by boiling H_2O . (Schulz, Z. anorg. **1895**, **10**. 153.)

Cadmium dichromate, $\text{CdO}, 2\text{CrO}_3 + \text{H}_2\text{O}$.

Easily sol. in H_2O without decomp; hygroscopic. (Schulz, Z. anorg. **1895**, **10**. 152.)

Easily sol. in H_2O but decomp. on evaporation. (Gröger, Z. anorg. **1910**, **66**. 11.)

Cadmium trichromate, $\text{CdCr}_2\text{O}_7 + \text{H}_2\text{O}$.

Deliquescent. (Gröger, Z. anorg. **1910**, **66**. 12.)

Cadmium chromate ammonia, $\text{CdCrO}_4, 4\text{NH}_3 + 3\text{H}_2\text{O}$.

Efflorescent. Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; insol. in alcohol and ether. (Malaguti and Sarzeau.)

Cadmium potassium chromate, $\text{CdK}_2(\text{CrO}_4)_2 + 2\text{H}_2\text{O}$.

Ppt. Decomp. by H_2O . (Gröger, Z. anorg. **1907**, **54**. 189.)

$3\text{CdO}, \text{K}_2\text{O}, 3\text{CrO}_3 + 3\text{H}_2\text{O}$. Ppt. (Preis and Raymann, Sitzungsber. böhm. Gesell. **1880**.)

$4\text{CdO}, \text{K}_2\text{O}, 4\text{CrO}_3 + 3\text{H}_2\text{O}$. Ppt. Slowly decomp. by H_2O . (Groger, M. **1904**, **25**. 533.)

Cadmium potassium dichromate,

$\text{CdCr}_2\text{O}_7, \text{K}_2\text{Cr}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Sol. in H_2O ; sl. hygroscopic. (Krüss, Z. anorg. **1895**, **8**. 454.)

Cadmium dichromate mercuric cyanide, $\text{CdCr}_2\text{O}_7, 2\text{Hg}(\text{CN})_2 + 7\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (Z. anorg. **1895**, **8**. 460.)

Cæsium chromate, Cs_2CrO_4 .

(Chabrié, C. R. **1901**, **132**. 680.)

Aq. solution sat. at 30° contains (Schreinemakers, C. C. **1909**, I. 11.)

Cæsium dichromate, $\text{Cs}_2\text{Cr}_2\text{O}_7$.

(Chabrié, C. R. **1901**, **132**. 680.)

Much more sol. in hot H_2O , than in (Fraprie, Am. J. Sci. **1906**, (4) **21**. 309.)

Aq. solution sat. at 30° contains (Schreinemakers, C. C. **1909**, I. 11.)

Cæsium trichromate, $\text{Cs}_2\text{Cr}_3\text{O}_{10}$.

Decomp. by H_2O . (Schreinemakers, Weekbl. **1908**, **5**. 811.)

Sol. in H_2O . (Fraprie, Am. J. Sci. (4) **21**. 315.)

Cæsium tetrachromate, $\text{Cs}_2\text{Cr}_4\text{O}_{13}$.

Sol. in H_2O with decomp. (Schreinemakers, Chem. Weekbl. **1908**, **5**. 811.)

Cæsium cobaltous chromate,

$\text{Cs}_2\text{Co}(\text{CrO}_4)_2 + 6\text{H}_2\text{O}$.

(Briggs, Z. anorg. **1907**, **56**. 248.)

Cæsium magnesium chromate,

$\text{Cs}_2\text{Mg}(\text{CrO}_4)_2 + 6\text{H}_2\text{O}$.

(Briggs, Chem. Soc. **1904**, **85**. 680.)

Cæsium nickel chromate, $\text{Cs}_2\text{Ni}(\text{CrO}_4)_2 + 6\text{H}_2\text{O}$.

Sol. in cold H_2O without much change; decomp. by warm H_2O . (Briggs, Soc. **1904**, **85**. 679.)

Calcium chromate basic, $\text{Ca}_2\text{CrO}_6 + 3\text{H}_2\text{O}$.

Sol. in 230 pts. H_2O without decomp. (Mylius and Wrochem, Gm. K. **3**. I, 1)

Calcium chromate, CaCrO_4 .

Anhydrous. Very sl. sol. in H_2O . (Mylius, J. B. **1862**. 148.)

Aq. solution sat. at 18° contains CaCrO_4 ; sp. gr. = 1.023. (Mylius and Wrochem, B. **1900**, **33**. 3688.)

Insol. in acetone. (Naumann, B. **1843**, **29**.)

$+ \frac{1}{2}\text{H}_2\text{O}$. Aq. solution sat. at 18° contains 4.4% CaCrO_4 ; sp. gr. = 1.044. (Mylius and Wrochem, B. **1900**, **33**. 3688.)

$+ \text{H}_2\text{O}$. Solubility in H_2O at t° .

t°	0°	8°	13°	18°
% CaCrO_4	11.5	10.8	10.3	9.6

t°	40°	60°	75°	90°
% CaCrO_4	7.8	5.7	4.6	3.6

(Mylius and Wrochem, Gm.-K. **3**. I,

Sp. gr. of solution containing 9.6% by wt. CaCrO_4 at $18^\circ = 1.096$. (Mylius and Wrochem, B. 1900, **33**, 3688.)

+ $2\text{H}_2\text{O}$. Sol. in 241.3 pts. H_2O at 14° . (Siewert.)

Sol. in 34 pts. H_2O . (Schwarz, Dingl. **198**, 59.)

Solubility of two modifications in H_2O at t° .

α modification.

t°	0°	20°	30°	45°
% CaCrO_4	14.75	14.22	13.89	12.53

β modification.

t°	0°	14°	18°	19.5°	30°	40°
% CaCrO_4	9.8	10	10.3	10.4	10.4	10.4

(Mylius and Wrochem, Gm.-K. **3**, I, 1387.)

α modification. Sp. gr. of the solution containing 14.3% by wt. CaCrO_4 at $18^\circ = 1.149$. (Mylius and Wrochem, B. 1900, **33**, 3688.)

β modification. Sp. gr. of the solution containing 10.3% by wt. CaCrO_4 at $18^\circ = 1.105$. (Mylius and Wrochem, B. 1900, **33**, 3688.)

Easily sol. in H_2O containing CrO_3 . Insol. in absolute alcohol.

50 cc. of alcohol (29%) dissolve 0.608 g. CaCrO_4 ; 50 cc. of alcohol (53%) dissolve 0.44 g. CaCrO_4 . (Fresenius, Z. anal. **30**, 672.)

Sol. in acids and in dilute alcohol. (Caron and Raquet, Bull. Soc. 1906, (3) **35**, 1064.)

Calcium dichromate, $\text{CaCr}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Very deliquescent. (Bahr, J. pr. **60**, 60.)

In sat. solution at 18° , 61% CaCr_2O_7 is present. (Mylius and Wrochem, Gm.-K. **3**, 1387.)

Sol. in acetone. (Naumann, B. 1904, **37**, 28.)

Calcium potassium chromate, $\text{CaCrO}_4, \text{K}_2\text{CrO}_4$.

(Barre, C. R. 1914, **158**, 495.)

+ H_2O . Easily sol. in H_2O . (Duncan.)

sol. in H_2O when ignited.

+ $2\text{H}_2\text{O}$. Easily sol. in H_2O , even after ignition. Insol. in alcohol. (Duncan, J. B. **10**, 313.)

Formed below 45° . (Barre, C. R. 1914, **1**, 495.)

sol. in cold H_2O . Sl. sol. in sat. K_2CrO_4 + H_2O . (Gröger, Z. anorg. 1907, **54**, 187.)

Two modifications. Solubility of α modification is somewhat less than that of the β modification. (Wyrouboff, Bull. Soc. **18**, 1891, 14, 255.)

Solubility of two modifications in H_2O at t° .

t°	0°	15°
Solubility of α	23.06	25.06
" " β	23.01	24.45

(Rakowski, C. C. **1909**, I, 133.)

$4\text{CaCrO}_4, \text{K}_2\text{CrO}_4$.

$5\text{CaCrO}_4, \text{K}_2\text{CrO}_4$. Sol. in much H_2O . (Bahr.)

Calcium chromate potassium sulphate,

$\text{CaCrO}_4, \text{K}_2\text{SO}_4 + \text{H}_2\text{O}$.

Decomp. by H_2O . (Hannay, Chem. Soc. **32**, 399.)

$\text{CaCrO}_4, \text{K}_2\text{SO}_4, \text{K}_2\text{CrO}_4$. As above. (H.)

Cerous chromate.

Insol. in H_2O .

Calcium strontium chromate, $\text{CaSr}(\text{CrO}_4)_2$.

(Bourgeois, Bull. Soc. Min. 1879, **2**, 123.)

Ceric dichromate, $\text{CeO}_2, 2\text{CrO}_3 + 2\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in acids; decomp. completely by boiling H_2O . (Bricout, C. R. 1894, **118**, 145.)

Chromic chromate, $\text{CrO}_2 = \text{Cr}_2\text{O}_3, \text{CrO}_3$.

Insol. as such in H_2O , but decomp. thereby into CrO_3 and Cr_2O_3 ; decomp. by alkaline and many saline solutions. Easily sol. in dil. acids if recently pptd, but with difficulty if dried at a high temp. (Eliot and Storer, Proc. Am. Acad. **5**, 207.)

$\text{Cr}_2\text{O}_{12} = \text{Cr}_2\text{O}_3, 3\text{CrO}_3$. Sol. in $\text{HCl} + \text{Aq}$. Very slowly sol. in $\text{HNO}_3 + \text{Aq}$. Slowly decomp. by H_2SO_4 or $\text{NH}_4\text{OH} + \text{Aq}$. Easily decomp. by $\text{KOH} + \text{Aq}$.

Does not exist. (Eliot and Storer, *l.c.*)

$\text{Cr}_2\text{O}_{18} = 3\text{Cr}_2\text{O}_3, 2\text{CrO}_3$. Easily sol. in HCl or $\text{HNO}_3 + \text{Aq}$; difficulty sol. in acetic acid. Easily sol. in $\text{KOH} + \text{Aq}$. (Traube, A. **66**, 108.)

Existence doubtful.

$\text{Cr}_2\text{O}_9 = 2\text{Cr}_2\text{O}_3, \text{CrO}_3$. Insol. in all acids, even aqua regia; slowly attacked by a boiling conc. solution of alkali hydroxides. (Geuther and Merz, A. **118**, 62.) Cr_2O_5 according to Wöhler.

Chromic cupric chromate, $\text{CuCr}_4\text{O}_9, \text{Cr}_2\text{O}_3 + 12\text{H}_2\text{O}$.

Insol. in H_2O and H_2SO_4 . Sol. in HCl and HNO_3 . (Rosenfeld, B. 1879, **12**, 957.)

$6\text{CuO}, \text{Cr}_2\text{O}_3, \text{CrO}_3 + 9\text{H}_2\text{O}$. Insol. in H_2O . Sol. in acids. (Rosenfeld, B. 1879, **12**, 958.)

Chromic potassium chromate, $\text{Cr}_2\text{H}_2(\text{CrO}_4)_2, \text{K}_2\text{CrO}_4(?)$.

Insol. in H_2O , alcohol, or acetic acid. Not attacked by cold $\text{HNO}_3 + \text{Aq}$; sl. oxidized when hot. Insol. in cold, easily sol. in hot H_2SO_4 . Sl. sol. in $\text{SO}_2 + \text{Aq}$. Sol. in conc. $\text{HCl} + \text{Aq}$. (Tommasi, Bull. Soc. (2) **17**, 396.)

Chromous potassium chromate,

$\text{K}_2\text{CrO}_4(\text{CrO}_2)_2 = \text{K}_2\text{Cr}(\text{CrO}_4)_2(?)$.

Sat. cold solution in H_2O contains 9% of the salt. Insol. in alcohol and ether. (Heintze, J. pr. (2) **4**, 212.)

Cobaltous chromate, basic, $3\text{CoO}, \text{CrO}_3 + 4\text{H}_2\text{O}$.

Ppt. Decomp. by H_2O . (Malaguti and Sarzeau, A. ch. (3) 9. 431.)

True formula is $2\text{CoO}, \text{CrO}_3 + 2\text{H}_2\text{O}$. (Freese, Pogg. 140. 252.)

$4\text{CoO}, 3\text{CrO}_3 + 2\text{H}_2\text{O}$.
Decomp. by H_2O . (Gröger, Z. anorg. 1906, 49. 203.)

Cobaltous chromate, CoCrO_4 .

Much more sol. in H_2O than NiCrO_4 . Easily sol. in hot dil. $\text{HNO}_3 + \text{Aq}$. (Briggs, Z. anorg. 1909, 63. 327.)

$+ 2\text{H}_2\text{O}$. Ppt. (Briggs, Z. anorg. 1909, 63. 328.)

Cobaltous dichromate, $\text{CoCr}_2\text{O}_7 + \text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O . (Briggs, Z. anorg. 1907, 56. 247.)

Cobaltous potassium chromate, basic.

$\text{K}_2\text{O}, 4\text{CoO}, 4\text{CrO}_3 + 3\text{H}_2\text{O}$.

Sol. in cold dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Gröger, Z. anorg. 1906, 49. 199.)

Cobaltous potassium chromate, $\text{K}_2\text{Co}(\text{CrO}_4)_2 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Gröger, Z. anorg. 1906, 49. 200.)

Cupric chromate, basic, $3\text{CuO}, \text{CrO}_3 + 2\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in dil. $\text{HNO}_3 + \text{Aq}$ and in $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$. (Malaguti and Sarzeau, A. ch. (3) 9. 434.)

$7\text{CuO}, 2\text{CrO}_3 + 5\text{H}_2\text{O}$. Ppt. (Rosenfeld, B. 13. 1469.)

$7\text{CuO}, \text{CrO}_3 + 5\text{H}_2\text{O}$. Ppt. (R.)

Cobaltous dichromate mercuric cyanide, $\text{CoCr}_2\text{O}_7, 2\text{Hg}(\text{CN})_2 + 7\text{H}_2\text{O}$.

Very stable. Sol. in H_2O . (Krüss, Z. anorg. 1895, 8. 458.)

Cupric chromate, $\text{Cu}(\text{CrO}_4)$.

Insol. in H_2O ; very sol. in chromic acid and in other acids; decomp. by boiling with H_2O . (Schulz, Z. anorg. 1895, 10. 152.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

Cupric dichromate, basic, $\text{CuCr}_2\text{O}_7, 2\text{CuO}$.
(Stanley, C. N. 54. 194.)

Cupric dichromate, $\text{CuCr}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Deliquescent. Very easily sol. in H_2O , $\text{NH}_4\text{OH} + \text{Aq}$, and alcohol. (Drüge, A. 101. 39.)

Aqueous solution is decomp. by boiling. (Malaguti and Sarzeau, A. ch. (3) 9. 456.)

Very hygroscopic. Very sol. in H_2O without decomp. (Schulz, Z. anorg. 1895, 10. 150.)

Cupric tetrachromate, $\text{CuCr}_4\text{O}_{13} + 2\text{H}_2\text{O}$.

Deliquescent. Decomp. when its sol. in H_2O is concentrated. (Gröger, Z. 1910, 66. 15.)

Cupric lead chromate, $2(\text{PbCrO}_4, 2\text{CuCrO}_4, \text{CuO})$.

Min. *Vauquelinite*. Sol. in acids.

Cupric potassium chromate, basic, $\text{KCu}_2(\text{OH})(\text{CrO}_4)_2 + \text{H}_2\text{O}$.

Ppt. (Gröger, M. 1903, 24. 485.)

$3\text{CuO}, \text{K}_2\text{O}, 3\text{CrO}_3 + 2\text{H}_2\text{O}$. Nearly in H_2O . Sol. in NH_4OH or $(\text{NH}_4)_2\text{CO}_3$ (Knop, A. 70. 52.)

Does not exist. (Rosenfeld, B. 13. 4CuO, $\text{K}_2\text{O}, 4\text{CrO}_3 + \text{H}_2\text{O}$. Decom. by boiling H_2O . (Gerhardt.)
 $+ 3\text{H}_2\text{O}$. Decomp. by boiling H_2O . ger, Dissert. 1880.)

Cupric potassium chromate ammonia, $\text{K}_2\text{Cu}(\text{CrO}_4)_2, 2\text{NH}_3$.

Very sol. in dil. $\text{NH}_3 + \text{Aq}$; decomp. (Briggs, Chem. Soc. 1904, 85. 672.)

Cupric chromate ammonia, CuCrO_4 .

Decomp. by H_2O . Sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Parravano and Pasta, Gazz. 1907, 37. (2), 255.)

$4\text{CuCrO}_4, 3\text{NH}_3 + 5\text{H}_2\text{O}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; insol. in organic solvent sol. in $\text{AgNO}_3 + \text{Aq}$. (Schuyten, C. (I. 399.)

$2\text{CuCrO}_4, 7\text{NH}_3 + \text{H}_2\text{O}$. Decomp. l. Very sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Briggs, Soc. 1904, 85. 673.)

$3\text{CuO}, 2\text{CrO}_3, 10\text{NH}_3 + 2\text{H}_2\text{O}$. l. by H_2O ; sl. sol. or insol. in alcohol, e. $\text{NH}_4\text{OH} + \text{Aq}$. (Malaguti and Sarzeau, Dissert. 1880.)
Decomp. by hot H_2O ; insol. in (Büttger.)

Cupric dichromate ammonia, $\text{CuCr}_2\text{O}_7, 4\text{NH}_3 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Parravano and Pasta, Gazz. 1907, 37. (2) 255.)

Cupric dichromate mercuric cyanide, $\text{CuCr}_2\text{O}_7, \text{Hg}(\text{CN})_2 + 5\text{H}_2\text{O}$.

Not hygroscopic. Sol. in H_2O . (Krüss, Z. anorg. 1895, 8. 461.)

Didymium chromate, $\text{Di}_2(\text{CrO}_4)_3$.

Sl. sol. in H_2O , easily in dil. acids. (Richs and Smith, A. 191. 353.)
 $+ 7\text{H}_2\text{O}$. (Cleve.)

Didymium potassium chromate, $\text{Di}_2(\text{CrO}_4)_3, \text{K}_2\text{CrO}_4$.

Precipitate. Decomp. by H_2O . (C

chromate, $Dy_2(CrO_4)_3$

sol. in H_2O . 1.0002 pt. is sol. in O at 25° . (Jantsch, B. 1911, 44.)

chromate, basic, $GlCrO_4$, $13GlO +$

sol. in H_2O . (Creuzberg, Dingl.

$6Gl(OH_2)$. Ppt. Insol. in H_2O . (B. 1907, 40. 2603.)

chromate, $GlCrO_4 + H_2O$.

by H_2O with separation of the
nate. (Glassmann, B. 1907, 40.

) chromate, $Au_2(CrO_4)_3, CrO_3$.

loff, Ch. Z. 1907, 31. 1182.)

romate.

yer.)

chromate.

in H_2O . Known only in solution.

) chromate, basic.

by H_2O . (Maus.)

CrO_3 . Insol. in H_2O , but decomp.

by saline solutions; easily sol. in
l. in $H_2CrO_4 + Aq$. (Eliot and
c. Am. Acad. 5. 216.)

) dichromate.

H_2O and alcohol. (Maus, Pogg. 9.

ic) potassium chromate, basic,

$Fe_2O_3, 3K_2O$.

$Fe_2O_3, 4K_2O$.

$3Fe_2O_3, 7K_2O$.

$3Fe_2O_3, 4K_2O + 9H_2O$.

$Fe_2O_3, 6K_2O + 6H_2O$.

$Fe_2O_3, 6K_2O + 10H_2O$.

$3Fe_2O_3, 6K_2O + 5H_2O$.

$Fe_2O_3, 2K_2O + 7H_2O$.

$Fe_2O_3, K_2O + 4H_2O$.

$Fe_2O_3, 3K_2O$.

$4Fe_2O_3, 5K_2O + 8H_2O$.

compounds are ppts., insol. in H_2O ,
ether. (Lepierre, C. R. 1894, 119,

ic) potassium chromate,

$O_4)_3, K_2CrO_4 + 4H_2O$.

by much H_2O , conc. HCl , or
aq. Not decomp. by alcohol.
B. 12. 130C.)

c) sodium chromate, basic,

$7Fe_2O_3, 4Na_2O$.

Lepierre, C. R. 1894, 119, 1217.)

Lanthanum chromate, $La_2(CrO_4)_3$.

Sl. sol. in cold, more easily in hot H_2O ;
easily sol. in acids. (Frerichs and Smith, A.
191. 355.)

$+8H_2O$. Ppt. (Cleve.)

Lanthanum potassium chromate.

(Cleve.)

**Lead chromate, basic, $2PbO, CrO_3$ (chrome
red).**

Insol. in H_2O ; acetic acid dissolves out $\frac{1}{2}$
the PbC . Sol. in $KOH + Aq$. (Badams,
Pogg. 3. 221.)

Insol. in acetone. (Naumann, B. 1904, 37.
4329.)

$3PbO, CrO_3$. (Hermann, Pogg. 28. 162.)
 $+H_2O$. Ppt. (Strömholm, Z. anorg. 1904,
38. 443.)

Min. *Melanochroite*, *Phenicocroite*. Sol.
in acids.

$PbO, PbCrO_4$. Ppt. (S.)

Lead chromate, $PbCrO_4$.

Insol. in H_2O . Pptd. from $Pb(NO_3)_2$ in
presence of 70,000 pts. H_2O . (Harting.)

Calculated from electrical conductivity of
 $PbCrO_4 + Aq$, 1 l. H_2O dissolves 0.2 mg.
 $PbCrO_4$ at 18° . (Kohlrausch and Rose, Z.
phys. Ch. 12. 241.)

1 l. H_2O dissolves 1.2×10^{-4} g. $PbCrO_4$ at
 25° . (Hevesy, Z. anorg. 1913, 82. 328.)

Sol. in dil. $H_2SO_4 + Aq$ (Storer); sl. sol. in
dil. $HNO_3 + Aq$.

Sol. in 560 pts. $HNO_3 + Aq$ of 1.12 sp. gr.;
in 150 pts. $HNO_3 + Aq$ of 1.225 sp. gr.; in
130 pts. $HNO_3 + Aq$ of 1.265 sp. gr.; in 80
pts. $HNO_3 + Aq$ of 1.395 sp. gr. (Storer's
Dict.)

Solubility of $PbCrO_4$ in $HNO_3 + Aq$ at 18° .

(Millimols. per l.)

0.1N	0.2N	0.3N	0.4N
0.506	0.844	1.13	1.44

(Beck and Stegmüller, l.c.)

Easily decomp. by hot $HCl + Aq$. (Frese-
inus.)

Solubility of $PbCrO_4$ in $HCl + Aq$.

(Millimols. per l.)

t°	0.1N	0.2N	0.3N	0.4N	0.5N	0.6N
18	0.186	0.393	0.654	1.07	1.56	2.25
25	0.239	0.485	0.839	1.32	4.06	2.95
37	0.357	0.744	1.31	2.10	3.28	4.69

(Beck and Stegmüller, Arb. K. Gesund.
Amt. 1910, 34. 446.)

Insol. in $HC_2H_3O_2 + Aq$.

Easily sol. in KOH , or $NaOH + Aq$. 1 l.
 $KOH + Aq$ ($\frac{1}{2}$ normal) dissolves 11.9 g.
 $PbCrO_4$ at 15° ; 16.2 g. at 60° ; 26.1 g. at 80° ;

38.5 g. at 102. (Lachaud and Lepierre, Bull. Soc. (3) 6. 230.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq.}$ (Brett, 1837.)

Sol. in $\text{K}_2\text{Cr}_2\text{O}_7 + \text{Aq.}$; almost completely insol. in $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$, or $\text{NH}_4\text{NO}_3 + \text{Aq.}$

Not pptd. in presence of Na citrate. (Spiller.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Min. *Crocoite*. Sol. in hot $\text{HCl} + \text{Aq.}$; difficultly sol. in $\text{HNO}_3 + \text{Aq.}$; sol. in $\text{KOH} + \text{Aq.}$

Lead dichromate, PbCr_2O_7 .

Decomp. by H_2O .

+ $2\text{H}_2\text{O}$. As above. (Preis and Raymann, B. 13. 340.)

Lead lithium chromate, PbCrO_4 , Li_2CrO_4 .

(Lachaud and Lepierre, C. R. 110. 1035.)

Lead potassium chromate, PbCrO_4 , K_2CrO_4 .

Insol. in hot or cold H_2O or in alcohol. Dil. acids dissolve out K_2CrO_4 . (Lachaud and Lepierre, C. R. 110. 1035.)

Decomp. by H_2O . Stable in contact with solutions containing:

8.950 pts.	K_2CrO_4	per 100 pts.	H_2O	at	10°
8.077	"	"	"	"	27.5°
7.629	"	"	"	"	37.5°
7.150	"	"	"	"	50.0°
6.145	"	"	"	"	76.0°
4.940	"	"	"	"	100.0°

(Barre, C. R. 1914, 158. 497.)

Lead sodium chromate, PbCrO_4 , NaCrO_4 .

Sol. in $\text{H}_2\text{O} (?)$. (Lachaud and Lepierre.)

PbCrO_4 , 2PbO , Na_2CrO_4 . (L. and L.)

Lithium chromate, Li_2CrO_4 .

100 cc. of solution sat. at 18° contain 85 g. anhydrous salt. (Kohlrausch, B. A. B. 1897. 90.)

99.94 pts. are sol. in 100 pts. H_2O at 30°. (Schreinemakers, C. C. 1905. II, 1486.)

+ $2\text{H}_2\text{O}$. Very easily sol. in H_2O . (Rammelsberg, Pogg. 128. 323.)

100 g. H_2O dissolve 111 g. salt at 20°. (Von Weimarn, C. C. 1911. II, 1300.)

Sp. gr. of solution sat. at 18° = 1.574, and contains 52.6% LiCrO_4 . (Mylius and Wrochem, B. 1897, 30. 1718.)

Lithium dichromate, $\text{Li}_2\text{Cr}_2\text{O}_7$.

130.4 pts. are sol. in 100 pts. H_2O at 30°. (Schreinemakers, C. C. 1905. II, 1486.)

+ $2\text{H}_2\text{O}$. Deliquescent. Sol. in H_2O . (Rammelsberg.)

Lithium potassium chromate, K_2CrO_4 ,

$\text{Li}_2\text{CrO}_4 + \frac{1}{2}\text{H}_2\text{O}$.

Hydroscopic. (Zehenter, M. 1897, 18. 54.)

Magnesium chromate, MgCrO_4 .

Sol. in H_2SO_4 , and HCl ; insol. in 1 (Dufau, C. R. 1896, 123. 888.)

Sp. gr. of $\text{MgCrO}_4 + \text{Aq sat. at } t^\circ$			
t°	13.6°	14.5°	1
% MgCrO_4	12.31	21.86	:
Sp. gr.	1.0886	1.1641	:

(Slotte, W. Ann. 1881, 14. 19.)

Sol. in acetone. (Naumann, B. 19 4328.)

+ $7\text{H}_2\text{O}$. Easily sol. in H_2O . (Vau 100 cm. of solution sat. at 18° conta MgCrO_4 . (Kohlrausch, B. A. B. 1897

Sp. gr. of solution sat. at 18 = 1.4 contains 42% MgCrO_4 . (Mylius an chem, B. 1897, 30. 1718.)

+ $5\text{H}_2\text{O}$. Very sol. in H_2O . (Wyn Bull. Soc. Min. 12. 60.)

Magnesium dichromate, $\text{Mg}_2\text{Cr}_2\text{O}_7$.

Sol. in H_2O .

Sl. sol. in alcohol. (Reinitzer, Zeit. 1913, 26. 456.)

Magnesium potassium chromate, $\text{MgK}_2\text{CrO}_4 + 2\text{H}_2\text{O}$.

100 pts. H_2O dissolve 28.2 pts. at 2 pts. at 60°. (Schweitzer.)

Sol. in H_2O . Sl. sol. in sat. K_2CrC (Gröger, Z. anorg. 1907, 54. 188.)

Insol. in alcohol.

+ $6\text{H}_2\text{O}$. Efflorescent. (Briggs, Soc. 1904, 85. 679.)

Magnesium rubidium chromate, $\text{MgRb}_2(\text{CrO}_4)_2 + 6\text{H}_2\text{O}$.

(Briggs, Chem. Soc. 1904, 85. 679.) (Chem. Soc. 1911, 99. 1327.)

Magnesium sodium chromate.

(Stanley, C. N. 54. 194.)

Manganous chromate, 2MnO , CrO_3 .

Ppt. Sol. in dil. H_2SO_4 , or HNO_3 (Warrington and Reinsch, Schw. J.

Manganous potassium chromate, $\text{MgK}_2\text{CrO}_4 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in dil. (Gröger, Z. anorg. 1905, 44. 459.)

2MnCrO_4 , $\text{K}_2\text{CrO}_4 + 4\text{H}_2\text{O}$. Sol. i (Hensgen, R. t. c. 3. 433.)

Mercurous chromate, basic, $4\text{Hg}_2\text{O}$,

Very sl. sol. in cold, more in boiling Sl. sol. in $\text{HNO}_3 + \text{Aq.}$ Decomp. by Aq. Sl. sol. in $\text{NH}_4\text{Cl} + \text{Aq}$ or NH_4NO_3 (Brett.)

Does not exist. (Richter, B. 15. 1 3 Hg_2O , CrO_3 . Sol. in $\text{HNO}_3 + \text{Aq.}$ ter.)

CrO_3 . Ppt. (Fichter, Z. anorg. 50.)

chromate, Hg_2CrO_4 .

sol. in cold, more readily in hot. l. in dil. HNO_3 + Aq; sol. in conc. in KCN + Aq; insol. in $\text{Hg}_2(\text{NO}_3)_2$; see, Pogg. 53. 124.)

in K_2CrO_4 + Aq. than in H_2O . anorg. 1912, 76. 349.)

acetone. (Naumann, B. 1904, 37.)

chromate, basic, 2HgO , CrO_3 .

l. and in HNO_3 + Aq. (Geuther.)

CrO_3 . Sl. sol. in H_2O . (Millon.)

true compound. All others are

HgO or HgCrO_4 with this com-
p. Z. anorg. 1904, 40. 155.)

CrO_3 . Sl. sol. in H_2O . (Millon, A. 365.)

CrO_3 . Easily sol. in warm HNO_3 ,
ly precipitated. Easily sol. in
(Geuther, A. 106. 247.)

exist. (Freese, B. 2. 477.)

CrO_3 . Easily sol. in HCl + Aq.
l. in HNO_3 + Aq. Decomp. by

O_2 . Insol. in H_2O . (Jäger and
2. 2049.)

chromate, HgCrO_4 .

by H_2O and acids into basic
her.)

acids. Sol. in warm NH_4Cl , or
l. in $\text{Hg}(\text{NO}_3)_2$, or HgCl_2 +

ethyl acetate. (Naumann, B.
4.)

acetone. (Naumann, B. 1904,

chromate, HgCr_2O_7 .

Walowski, C. C. 1906. II, 1307.)

potassium chromate,

CrO_4).

comp. by H_2O . (Gröger, Z. anorg.
1.)

chromate, basic, ammonia, 12HgO ,
 2NH_3 + $3\text{H}_2\text{O}$.

Z. anorg. 1908, 58. 420.)

chromate ammonia, HgCrO_4 ,
- H_2O .

Z. anorg. 1908, 58. 419.)

chromate sulphide, 2HgCrO_4 , HgS .

ked by weak acids. (Palm, C. C.

mate, basic, 4NiO , CrO_3 + $6\text{H}_2\text{O}$.

H_2O ; easily sol. in NH_4OH + Aq.
nd Sarzeau, A. ch. (3) 9. 451.)

3NiO , CrO_3 + $6\text{H}_2\text{O}$. Insol. in H_2O ; sol. in
 NH_4OH + Aq. (Freese, J. B. 1869. 271.)

2NiO , CrO_3 + $6\text{H}_2\text{O}$. As above. (Schmidt,
A. 156. 19.)

5NiO , 2CrO_3 + $12\text{H}_2\text{O}$. As above.
(Schmidt.)

Nickel chromate, NiCrO_4 .

Not attacked by boiling H_2O .

Nearly insol. in hot dil. HNO_3 . Slowly sol.
in conc. HNO_3 and aqua regia.

Somewhat sol. in NH_3 + Aq. (Briggs, Z.
anorg. 1909, 63. 326.)

Nickel dichromate, $2\text{NiCr}_2\text{O}_7$ + $3\text{H}_2\text{O}$.

Slowly sol. in cold, rapidly sol. in hot H_2O .

Deliquescent. (Briggs, Z. anorg. 1907, 56.
246.)

Nickel potassium chromate, NiCrO_4 , K_2CrO_4 ,
+ $2\text{H}_2\text{O}$.

Decomp. by H_2O . (Gröger, Z. anorg. 1906,
51. 353.)

+ $6\text{H}_2\text{O}$. Efflorescent. (Briggs, Chem.
Soc. 1904, 85. 678.)

Nickel rubidium chromate, $\text{NiRb}(\text{CrO}_4)_2$ +
 $6\text{H}_2\text{O}$.

Sl. efflorescent at ord. temp. (Briggs,
Chem. Soc. 1904, 85. 678.)

Nickel chromate ammonia, NiCrO_4 , 6NH_3 +
 $4\text{H}_2\text{O}$.

Decomp. by H_2O . Quite easily sol. in
 NH_4OH + Aq of 0.96 sp. gr. (Schmidt.) In-
sol. in alcohol or ether.

Potassium chromate, K_2CrO_4 , $\text{K}_2\text{Cr}_2\text{O}_7$, etc.

System: K_2O , CrO_3 , H_2O at 0°

100 g. of the sat. solu- tion contain		Solid phase
g. K_2O	g. CrO_3	
31.18	...	
26.06	0.54	K_2CrO_4
19.31	4.27	"
17.73	5.50	"
17.06	11.77	"
17.18	11.91	"
17.62	18.71	"
17.63	18.72	"
17.61	18.91	K_2CrO_4 + $\text{K}_2\text{Cr}_2\text{O}_7$
17.79	19.10	"
17.80	19.10	"
10.90	11.93	$\text{K}_2\text{Cr}_2\text{O}_7$
8.07	8.93	"
1.87	3.13	"
1.41	3.00	"
1.42	3.01	"
0.97	3.94	"
0.78	22.38	"
1.02	38.83	"
1.26	40.10	"
1.36	40.41	"
1.22	41.70	"

System: K_2O , CrO_3 , H_2O at 0° —Continued			System: K_2O , CrO_3 , H_2O at 30°		
100 g. of the sat. solution contain		Solid phase	100 g. of the sat. solution contain		Solid phase
g. K_2O	g. CrO_3		g. K_2O	g. CrO_3	
1.28	41.75	$K_2Cr_2O_7$	46.8	0.94	$KOH, 2H_2O$
1.40	42.10	"	26.89	3.06	K_2CrO_4
1.23	42.11	"	22.25	6.99	"
1.33	42.16	"	19.52	13.72	"
1.31	42.28	"	18.65	17.00	"
1.38	42.48	"	18.60	17.03	"
1.40	42.68	"	18.70	20.30	"
1.47	42.93	$K_2Cr_2O_7 + K_2Cr_2O_{10}$	19.12	21.00	$K_2CrO_4 + K_2Cr_2O_7$
1.47	42.95	"	19.35	16.85	$K_2Cr_2O_7$
1.47	43.09	"	15.04	16.51	"
1.25	44.52	$K_2Cr_2O_{10}$	14.77	14.57	"
1.27	44.95	"	12.28	13.11	"
1.18	45.84	"	11.20	10.48	"
1.17	46.84	"	4.98	19.34	"
1.36	47.22	$K_2Cr_2O_{10} + K_2Cr_2O_{12}$	3.07	28.21	"
1.36	47.31	"	2.42	33.77	"
1.40	47.67	"	2.35	36.78	"
1.24	48.23	$K_2Cr_2O_{12}$	2.30	40.41	"
1.35	51.66	"	2.30	44.50	$K_2Cr_2O_7 + K_2Cr_2O_{10}$
1.10	53.81	"	2.50	49.95	$K_2Cr_2O_{10} + K_2Cr_2O_{12}$
1.08	55.63	"	2.25	53.39	$K_2Cr_2O_{12}$
1.16	56.93	"	1.35	62.81	$K_2Cr_2O_{12} + CrO_3$
0.96	57.63	"	0.69	62.52	CrO_3
1.16	59.46	"	(Koppel and Blumenthal, Z. anorg. 1907, 235.)		
0.91	59.87	"			
0.81	60.16	"	System: K_2O , CrO_3 , H_2O at 60°		
0.70	61.76	$K_2Cr_2O_{12} + CrO_3$			
0.62	61.77	"	100 g. of the sat. solution contain		Solid phase
0.57	61.78	"	g. K_2O	g. CrO_3	
0.67	61.86	"	50.0	0.53	KOH, H_2O
	61.51	CrO_3	32.98	9.15	K_2CrO_4
	61.52	"	21.05	8.99	"
	61.55	"	20.70	14.43	"
	61.57	"	20.25	16.56	"
			20.32	21.94	"
			20.67	22.00	"
			20.72	23.40	$K_2CrO_4 + K_2Cr_2O_7$
			20.68	23.74	$K_2Cr_2O_7$
			20.55	20.82	"
			14.53	20.93	"
			13.36	21.24	"
			10.01	21.24	"
			10.01	26.95	"
			8.39	31.49	"
			7.65	32.92	"
			7.51	39.64	"
			6.86	49.84	$K_2Cr_2O_7 + K_2Cr_2O_{12}$
			7.06	50.40	$K_2Cr_2O_{12}$
			0.51	52.70	"
			5.33	52.79	"
			5.49	53.42	"
			5.06	53.58	"
			5.12	53.70	"
			5.30		"

(Koppel and Blumenthal, Z. anorg. 1907, 53, 245.)

System: K_2O , CrO_3 , H_2O at 20°

100 g. of the sat. solution contain		Solid phase
g. K_2O	g. CrO_3	
2.21	42.92	$K_2Cr_2O_7 + K_2Cr_2O_{10}$
2.20	43.28	"
2.10	44.02	$K_2Cr_2O_{10}$
2.02	45.28	"
2.01	46.24	"
2.00	48.46	$K_2Cr_2O_{10} + K_2Cr_2O_{12}$
1.94	48.62	$K_2Cr_2O_{12}$
1.62	49.01	"
0.62	62.80	$K_2Cr_2O_{12} + CrO_3$

(Koppel and Blumenthal, Z. anorg. 1907, 53, 243.)

: K_2O , CrO_3 , H_2O at 60° —Continued

the sat. solu- n contain		Solid phase
	g. CrO_3	
	54.09	$K_2Cr_2O_7 + K_2Cr_4O_{13}$
	54.73	
	54.91	$K_2Cr_4O_{13}$
	55.43	"
	56.41	"
	58.05	"
	58.69	"
	60.69	"
	61.25	"
	61.27	"
	61.29	"
	62.57	"
	65.77	$K_2Cr_4O_{13} + CrO_3$
	65.12	

and Blumenthal, Z. anorg. 1907, **53**.
240.)

ystem: K_2O , CrO_3 , H_2O at the
cryohydric pt.

100 g. of the solu- tion contain		Solid phase
g. K_2O	g. CrO_3	
17.18	18.11	$K_2CrO_4 + K_2Cr_2O_7$
1.18	42.51	
0.79	45.69	$K_2Cr_2O_7 + K_2Cr_3O_{10}$
		$K_2Cr_3O_{10} + K_2Cr_4O_{13}$

and Blumenthal, Z. anorg. 1907, **53**.
263-5.)

. of solutions of $CrO_3 + K_2O + Aq$.

100 g. of the solu- tion contain		Solid phase
g. K_2O	g. CrO	
30.01	11.92	K_2CrO_4
23.8	25.3	
24.3	30.5	$K_2CrO_4 + K_2Cr_2O_7$
16.4	35.6	
16.8	59.2	$K_2Cr_2O_7 + K_2Cr_3O_{10}$
...	71.2	
		CrO_3

and Blumenthal, Z. anorg. 1907, **53**.
255.)

m chromate, K_2CrO_4 .
sol. in H_2O .

pts. H_2O at 18.75° . (Ahl.)
 H_2O at 15° dissolve 43.857 pts. K_2CrO_4 and
sp. gr. of 1.3032. (Michel and Krafft, A.
478.)

dissolves in 2.07 pts. H_2O at 15.5° .
n.)
dissolves in 1.75 pts. H_2O at 17.5° ,
67 pts. H_2O at 100° . (Moser.)

100 pts. H_2O dissolve at—

0°	10°	20°	30°
58.90	60.92	62.94	64.96 pts. K_2CrO_4 ,

40°	50°	60°	70°
66.98	69.00	71.02	73.04 pts. K_2CrO_4 ,

80°	90°	100°	
75.06	77.08	79.10	pts. K_2CrO_4 .

(Alluard, C. R. **59**. 500.)

100 pts. H_2O dissolve at—

0°	10°	27.37°	42.1°
61.5	62.1	66.3	70.3 pts. K_2CrO_4 ,

63.6°	93.6°	106.1°	
74.9	79.7	81.8	pts. K_2CrO_4 .

(Nordenskjöld and Lindström, Pogg. **186**.
314.)

100 pts. $K_2CrO_4 + Aq$ sat. at $10-12^\circ$ con-
tain 37.14 pts. salt. (v. Hauer, J. pr. **103**.
114.)
100 pts. H_2O at 19.5° dissolve 62.3 pts.
 K_2CrO_4 , and solution has sp. gr. of 1.3787.
(Schiff, A. **109**. 326.)

Sat. $K_2CrO_4 + Aq$ contains at—

34°	53°	79°
39.7	40.3	41.8% K_2CrO_4 .

96°	120°	157°
42.6	44.0	45.4% K_2CrO_4 .

(Étard, A. ch. 1894, (7) **2**. 550.)

100 cc. sat. $K_2CrO_4 + Aq$ contain 53 g.
 K_2CrO_4 at 18° . (Kohlrausch, B. A. B. **1897**.
90.)
100 pts. H_2O dissolve 64.91 pts. K_2CrO_4
at 30° , or 100 g. of solution contain 39.36 g.
 K_2CrO_4 . (Schreinemakers, Chem. Weekbl.
1905, **1**. 837.)

100 g. H_2O dissolve:
54.57 g. K_2CrO_4 at -11.37° (cryohydric pt.)
57.11 g. " " 0°
65.13 g. " " 30°
74.60 g. " " 60°
88.80 g. " " 105.8° (b-pt. of sat. sol.)
(Koppel, Z. anorg. 1907, **53**. 262.)

64.62 g. K_2CrO_4 are sol. in 100 g. H_2O at
 25° . (Amadori, Real. Att. Line. 1912, (5) **21**,
I. 667.)

Sp. gr. of K_2CrO_4 +Aq at 19.5°.					
% K_2CrO_4	Sp. gr.	% K_2CrO_4	Sp. gr.	% K_2CrO_4	Sp. gr.
1	1.0080	15	1.1287	28	1.2592
2	1.0161	16	1.1380	29	1.2700
3	1.0243	17	1.1474	30	1.2808
4	1.0325	18	1.1570	31	1.2921
5	1.0408	19	1.1667	32	1.3035
6	1.0492	20	1.1765	33	1.3151
7	1.0576	21	1.1864	34	1.3268
8	1.0663	22	1.1964	35	1.3386
9	1.0750	23	1.2066	36	1.3505
10	1.0837	24	1.2169	37	1.3625
11	1.0925	25	1.2274	38	1.3746
12	1.1014	26	1.2379	39	1.3868
13	1.1104	27	1.2485	40	1.3991
14	1.1195

(Kremers, and Schiff, calculated by Gerlach, Z. anal. 8. 288.)

K_2CrO_4 dissolved in 2 pts. H_2O has sp. gr., 1.28; 3 pts., 1.21; 4 pts., 1.18; 5 pts., 1.15; 6 pts., 1.12; 7 pts., 1.11; 8 pts., 1.10. (Moser.)

Sp. gr. of sat. solution at 8° = 1.368. (Anthon, 1837.)

Sp. gr. of sat. K_2CrO_4 +Aq containing 24.26% K_2CrO_4 = 1.2335 at 18°/4°. (Slotte, W. Ann. 1881, 14. 18.)

Sp. gr. of K_2CrO_4 +Aq at 25°.

Concentration of K_2CrO_4 +Aq.	Sp. gr.
1-normal	1.0935
1/2- "	1.0475
1/4- "	1.0241
1/5- "	1.0121

(Wagner, Z. phys. Ch. 1890, 5. 36.)

Sat. K_2CrO_4 +Aq boils at 107°. (Kremers.)
Sat. K_2CrO_4 +Aq boils at 104.2° under 718 mm. pressure. (Alluard.)

Freezing point of sat. K_2CrO_4 +Aq = -12.5°. (Rüdorff.)

By dissolving K_2CrO_4 in 2 pts. H_2O , the temp. is lowered 10°. (Moser.)

100 pts. sat. solution of K_2CrO_4 and K_2SO_4 contain 37.14 pts. of the two salts at 10-12°. (v. Hauer, J. pr. 103. 114.)

Solubility of K_2CrO_4 + K_2SO_4 in H_2O at 25°. (G. per 100 g. H_2O .)

K_2CrO_4	K_2SO_4	K_2CrO_4	K_2SO_4
63.09	0.76	20.83	5.75
61.39	1.17	14.65	7.12
58.40	1.84	7.81	8.98
51.81	2.36	4.36	10.25
40.93	3.33	1.94	10.86
27.36	4.82

(Amadori, Real. Att. Line. 1912, (5) 21, I. 667.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

100 g. sat. solution in glycol at 15.4° contain 1.7 g. K_2CrO_4 . (de Coninck, C. C. 1906, II. 183.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899. II, 1014.)

+4 H_2O . Easily sol. in H_2O and in NH_4OH +Aq. (Wesch, Dissert. 1909.)

Potassium dichromate, $K_2Cr_2O_7$.

Sol. in H_2O , with slight absorption of heat. Less sol. in H_2O than K_2CrO_4 .

Sol. in 9.6 pts. H_2O at 17.3°. (Thompson.)
" 10 " " " 18.7°. (Moser.)

100 pts. H_2O at 15° dissolve 9.126 pts. $K_2Cr_2O_7$, and solution has sp. gr. = 1.0618. (Michel and Krafft, A. ch. (3) 41. 478.)

100 pts. H_2O dissolve pts. $K_2Cr_2O_7$. A = according to Alluard (C. R. 59. 500); K = according to Kremers (Pogg. 92. 497).

t°	A	K	t°	A	K
0	4.6	4.97	60	45.0	50.5
10	7.4	8.5	70	56.7
20	12.4	13.1	80	68.6	73.0
30	18.4	...	90	81.1
40	25.9	29.1	100	94.1	102.00
50	35.0

Solubility in H_2O at high temperatures.
100 pts. H_2O dissolve pts. $K_2Cr_2O_7$ at t°.

t°	Pts. $K_2Cr_2O_7$	t°	Pts. $K_2Cr_2O_7$
117	128.3	148	200.6
129	153.8	180	262.7

(Tilden and Shenstone, Phil. Trans. 1884. 23.)

Solubility of $K_2Cr_2O_7$ in H_2O at t°.

t°	% $K_2Cr_2O_7$	t°	% $K_2Cr_2O_7$
-1	4.1	92	42.8
+1	4.3	97	44.0
6	5.6	104	48.0
7	6.1	120	52.0
12	7.2	130	54.4
15	8.5	150	60.8
20	10.4	157	62.8
29	14.2	178	66.6
36	16.6	215	76.9
57	28.2	291	89.7
61	30.2	312	91.8
65	32.0	360	97.4
70	34.4

(Étard, A. ch. 1894, (7) 2. 550.)

100 g. H_2O dissolve 10.1 g. $K_2Cr_2O_7$ at 45° . (Greenish and Smith, Pharm. J. 1901, 774.)

100 pts. H_2O at 30° dissolve 18.12 pts. $K_2Cr_2O_7$. (Schreinemakers, Chem. Weekbl. 1905 1. 837.)

100 g. H_2O dissolve:
 4.50 g. $K_2Cr_2O_7$ at -0.63° (cryohydric pt.)
 4.64 g. " " 0°
 8.13 g. " " 30°
 5.44 g. " " 60°
 08.2 g. " " 104.8° (b-pt. of sat. sol.)
 (Koppel, Z. anorg. 1907, 53. 263.)

100 c.c. sat. solution contain 11.43 g. $K_2Cr_2O_7$ at 20° . (Sherrill and Eaton, J. Am. Chem. Soc. 1907, 29. 1643.)

100 g. sat. $K_2Cr_2O_7$ contain:
 5.52 g. $K_2Cr_2O_7$ at 4.81°
 15.17 " " 30.10°
 17.77 " " 35.33°
 Le Blanc and Schmandt, Z. phys. Ch. 1911, 77. 614.)

100 g. sat. $K_2Cr_2O_7 + Aq.$ at 35.03° contains 17.72 g. $K_2Cr_2O_7$. (Le Blanc, Z. phys. Ch. 1913, 86. 335.)

$K_2Cr_2O_7 + Aq$ sat. at 8° has sp. gr. 1.065. (Anthon, 337.)

Sp. gr. of $K_2Cr_2O_7 + Aq$ at 19.5° .

% $K_2Cr_2O_7$	Sp. gr.	% $K_2Cr_2O_7$	Sp. gr.
1	1.007	9	1.065
2	1.015	10	1.073
3	1.022	11	1.080
4	1.030	12	1.085
5	1.037	13	1.097
6	1.043	14	1.102
7	1.050	15	1.110
8	1.056

Kremers, calculated by Gerlach, Z. anal. 8. 288.)

Sp. gr. of $K_2Cr_2O_7 + Aq$ containing 4.71% $K_2Cr_2O_7 = 1.0325$ at $11^\circ/4^\circ$; containing 6.97% $K_2Cr_2O_7 = 1.0493$ at $10.6^\circ/4^\circ$. (Slotte, W. m. 1881, 14. 18.)

Sat. $K_2Cr_2O_7 + Aq$ boils at 104° (Kremers); 3.4° . (Alluard.)

Insol. in alcohol.

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. 1898, 20. 829.)

Insol. in alcohol. (Reinitzer, Zeit. angew. 1. 1913, 26. 456.)

100 g. sat. solution in glycol contain 6 g. $K_2Cr_2O_7$. (de Coninck, Bull. acad. roy. lg. 1906, 257.)

Insol. in benzonitrile. (Naumann, B. 14, 47. 1370.)

Insol. in acetone. (Naumann, B. 1904, 37. 29.)

Insol. in acetone and in methylal. (Eidmann, C. C. 1899. II, 1014.)

Potassium trichromate, $K_2Cr_2O_{10}$.

Easily sol. in H_2O and alcohol. (Bothe, J. pr. 46. 184.)

Not deliquescent; decomp. by H_2O in chromic acid and $K_2Cr_2O_7$. (Jäger and Krüss, B. 22. 2041.)

Potassium tetrachromate, $K_2Cr_2O_{12}$.

Very deliquescent, and easily sol. in H_2O . (Schwarz, Dingl. 186. 31.)

Not deliquescent. Decomp. by H_2O . (Jäger and Krüss, B. 22. 2042.)

Potassium samarium chromate,



Precipitate. (Cleve.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Potassium sodium chromate, $3K_2CrO_4$, Na_2CrO_4 .

Sol. in H_2O . (v. Hauer, J. pr. 83. 359.)
 64.2 pts. are sol. in 100 pts. H_2O at 14° .
 (Zehenter, M. 1897, 18. 49.)

Potassium strontium chromate, $K_2Sr(CrO_4)_2$.

Ppt. Decomp. by H_2O . (Gröger, Z. anorg. 1907, 54. 187.)

Decomp. by H_2O . Stable in contact with solutions containing:

at 11.5° , 2.914 pts. K_2CrO_4 per 100 pts. H_2O .
 at 27.5° , 4.123 " " " " " "
 at 50° , 5.942 " " " " " "
 at 76° , 7.920 " " " " " "
 at 100° , 9.784 " " " " " "

(Barre, C. R. 1914, 158. 496.)

Potassium thallium chromate, K_2CrO_4 , Tl_2CrO_4 .

(Lachaud and Lepierre, Bull. Soc. (3) 6. 232.)

+ $2H_2O$. Rapidly hydrolyzed by H_2O unless a large excess of the CrO_4 ion is present.

Readily sol. in dil. mineral acids.

Difficulty sol. in $K_2Cr_2O_7 + Aq$. (Hawley, J. Am. Chem. Soc. 1907, 29. 304.)

Potassium uranyl chromate, K_2CrO_4 , $2(UO_2)CrO_4 + 6H_2O$.

Decomp. by boiling with H_2O . Sol. in acidified H_2O . (Formánek, A. 257. 103.)

K_2CrO_4 , $(UO_2)CrO_4 + H_2O$; $2K_2CrO_4$, $3(UO_2)CrO_4 + 7H_2O$; $3K_2CrO_4$, $4(UO_2)CrO_4 + 7H_2O$; and K_2CrO_4 , $3(UO_2)CrO_4 + 14H_2O$.
 Precipitates. (Wiesner, C. C. 1882. 777.)

Potassium ytterbium chromate, basic, $2KYb(CrO_4)_2 + Yb(OH)_3 + 15\frac{1}{2}H_2O$.

Ppt. (Cleve, Z. anorg. 1902, 32. 151.)

Potassium yttrium chromate, K_2CrO_4 ,
 $Y_2(CrO_4)_3 + xH_2O$.

Ppt. (Cleve.)

Potassium zinc chromate, basic, K_2O , $5ZnO$,
 $4CrO_3 + 6H_2O$, or K_2O , $4ZnO$, $3CrO_3$,
 $+ 3H_2O$.

Slightly sol. in cold, decomp. by hot H_2O .
(Wöhler.)

K_2O , $4ZnO$, $3CrO_3 + 3H_2O$. Insol. in cold,
decomp. by hot H_2O . (Gröger, M. 1904,
25. 520.)

Potassium zinc chromate, $K_2Zn(CrO_4)_2 +$
 $2H_2O$.

Ppt. Decomp. by H_2O . (Gröger, Z.
anorg. 1907, 54. 189.)

Potassium dichromate chloride mercuric
chloride, $K_2Cr_2O_7, 2KCl, 4HgCl_2 + 2H_2O$.

Solution in H_2O sat. at 20.5° contains
6.78% salt. Salt is much more sol. in hot
 H_2O . (Strömholm, Z. anorg. 1912, 75. 278.)

Potassium chromate iodate.

See Chromoiodate, potassium.

Potassium chromate magnesium sulphate,
 $K_2CrO_4, MgSO_4 + 9H_2O$.

Sol. in H_2O . (Étard, C. R. 85. 443.)

Potassium chromate [mercuric chloride,
 $K_2CrO_4, 2HgCl_2$.

Easily sol. in H_2O . Sol. in dil. $HCl + Aq$.
(Darby.)

Potassium dichromate mercuric chloride,
 $K_2Cr_2O_7, HgCl_2$.

Ether or absolute alcohol dissolves out
 $HgCl_2$. (Millon, A. ch. (3) 18. 388.)

Can be crystallized from H_2O . (Jäger and
Krüss, B. 22. 2046.)

Potassium chromate mercuric cyanide,
 $2K_2CrO_4, 3Hg(CN)_2$.

Easily sol. in H_2O .

$+ H_2O$. (Dexter.)

Formula is $K_2CrO_4, 2Hg(CN)_2$. (Clarke
and Sterne, Am. Ch. J. 3. 352.)

Potassium dichromate mercuric cyanide,
 $K_2Cr_2O_7, Hg(CN)_2 + 2H_2O$.

Sol. in H_2O . (Wyrouboff, J. B. 1880. 309.)

Potassium chromate phosphate.

See Phosphochromate, potassium.

Potassium chromate sulphate, K_2CrO_4 ,
 $6K_2SO_4$.

Easily sol. in H_2O . (Boutron-Chalard.)

Potassium chromate tellurate.

See Chromotellurate, potassium.

Rubidium chromate, Rb_2CrO_4 .

Sol. in H_2O . (Piccard, J. pr. 86. 455.)

Solubility in H_2O at t° .

t°	% Rb_2CrO_4
— 7	36.65
0	38.27
10.3	40.22
20	42.42
30	44.11
40	46.13
50	47.44
60.4	48.90

(Schreinemakers and Filippo, Chem. Wee
1906, 3. 157.)

Rubidium dichromate, $Rb_2Cr_2O_7$.

Sol. in H_2O . (Grandeau, A. ch. (3)
227.)

Very sl. sol. in H_2O ; 5% at 10° , 8%
26°, 35% at 60° . (Wyrouboff, Bull.
Min. 1881, 4. 129.)

100 pts. H_2O dissolve 10.46 pts. Rb_2C
at 30° . The solution contains 9.47%
(Schreinemakers and Filippo, Chem. Wee
1906, 3. 157.)

Two forms of crystals. Figures denote
salt per 100 pts. H_2O .

t°	14°	26°	43°
Monoclinic form	4.45	8.00	16.52
Triclinic form	4.40	7.91	16.57

(Wyrouboff, Bull. Soc. 1908, (4) 3. 7.)

Solubility of monoclinic and triclinic for

Temp.	Pts. of salt in 100 pts. H_2O		Temp.	Pts. of salt 100 pts. H_2O	
	Mono- clinic	Tri- clinic		Mono- clinic	Tri- clinic
18°	5.42	4.96	40°	13.22	12
24°	6.94	6.55	50°	18.94	18
30°	9.08	8.70	60°	28.1	27

(Stortenbeker, C. C. 1907, II. 1588.)

**Rubidium dichromate chloride mer-
curic chloride**, $Rb_2Cr_2O_7, 2RbCl, 4Hg$
 $2H_2O$.

Sol. in H_2O .

Solution sat. at 20.5° contains 5.35%
(Strömholm, Z. anorg. 1912, 75. 284.)

Silver (argentous) chromate, Ag_2CrO_4 .

Sol. in dil. acids. (Wöhler and Rai-
berg.)

Existence very doubtful.

Silver chromate, Ag_2CrO_4 .

Absolutely insol. in H_2O . Sol. in a
ammonia, and alkali chromates + Aq . (C-
ington, A. 27. 12.)

Appreciably sol. in cold, and still mo
hot H_2O . (Meineke, A. 261. 341.)

100 ccm. H_2O dissolve 0.064 grain Ag_2
at 100° ; 100 ccm. H_2O containing 50 g

the following salts dissolve the given amts. Ag_2CrO_4 at 100° : NaNO_3 , 0.064 grain; KNO_3 , 0.192 grain; NH_4NO_3 , 0.320 grain; $(\text{NO}_3)_2$, 0.256 grain. (Carpenter, J. S. C. 5. 286.)

According to electrical conductivity of $\text{CrO}_4 + \text{Aq}$, 1 l. H_2O dissolves 28 mg. CrO_4 at 18° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

1. H_2O dissolves 25 mg. Ag_2CrO_4 at 18° . (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)

5 mg. are contained in 1 l. of sat. solution at 8° . Solubility increases unusually rapidly with temp. (Kohlrausch, Z. phys. Ch. 1908, 168.)

sol. in 26,378 pts. cold H_2O and 9116 pts. H_2O at 100° . (Koninck and Nihoul, Zeit. anorg. Ch. 1891, 5. 295.)

1. H_2O dissolves 1.2×10^{-4} gram. atoms silver at 25° . (Abegg and Cox, Z. phys. Ch. 1903, 46. 11.)

1. H_2O dissolves 0.029 g. Ag_2CrO_4 at 25° . (Jäger, Z. anorg. 1905, 45. 310.)

1. H_2O dissolves 0.0256 g. Ag_2CrO_4 at 0.0341 g. at 27° ; 0.0534 g. at 50° . (Whitby, Z. anorg. 1910, 67. 108.)

sol. in hot $\text{NH}_4\text{OH} + \text{Aq}$ of sp. gr. 0.94 (63% NH_3); sl. sol. in cold $\text{NH}_4\text{OH} + \text{Aq}$ of sp. gr. 0.91 (24.99% NH_3). (Margosches, Z. anorg. 1904, 41. 73.)

Solubility of Ag_2CrO_4 in $\text{NH}_4\text{OH} + \text{Aq}$ at 25° .

Mols. NH_4OH per l.	Mols. $\times 10^3$ Ag_2CrO_4 per l.
0.01	2.004
0.02	4.169
0.04	8.595
0.08	17.58

(Sherrill and Russ, J. Am. Chem. Soc. 1907, 29. 1662.)

sl. sol. in very conc. $\text{K}_2\text{CrO}_4 + \text{Aq}$. Practically insol. in $\text{AgNO}_3 + \text{Aq}$. (Margosches.)

Solubility of Ag_2CrO_4 in $\text{HNO}_3 + \text{Aq}$ at 25° .

Mols. HNO_3 per l.	Milliat. per l.		Solid Phase
	Cr	Ag	
0	32.20	5.390	$\text{Ag}_2\text{CrO}_4 + \text{Ag}_2\text{CrO}_7$
0.01	25.06	6.131	"
0.02	20.21	7.148	"
0.04	13.59	9.529	"
0.06	11.10	11.10	$\text{Ag}_2\text{Cr}_2\text{O}_7$
0.08	11.10	11.10	"
0.08 + 0.1 AgNO_3	6.624	"

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

1 l. 65% alcohol dissolves 0.0129 g. Ag_2CrO_4 at ord. temp. (Guerini, Dissert. 1912.)

Insol. in H_2O containing acetic acid in presence of large excess of AgNO_3 . (Gooch and Weed, Am. J. Sci. 1908, (4) 26. 85.)

Practically insol. in glacial acetic acid but somewhat sol. in dil. acetic acid. It behaves in a similar manner toward propionic, lactic and other organic acids. The red modification is more sol. than the greenish-black. (Margosches, Z. anorg. 1906, 51. 233.)

Silver dichromate, $\text{Ag}_2\text{Cr}_2\text{O}_7$.

Sl. sol. in H_2O . Easily sol. in HNO_3 , or $\text{NH}_4\text{OH} + \text{Aq}$. (Warrington.)

Decomp. by boiling with H_2O into CrO_3 and Ag_2CrO_4 . (Jäger and Krüss, B. 22. 2050.)

Decomp. by cold H_2O . (Autenrieth, B. 1902, 35. 2061.)

1 pt. is sol. in 12,000 pts. H_2O at 15° . (Mayer, B. 1903, 36. 1741.)

Solubility in H_2O at $25^\circ = 7.3 \times 10^{-3}$ atoms Ag per l. Decomp. by $\text{HNO}_3 + \text{Aq}$ (less than 0.06 N) with separation of Ag_2CrO_4 . (Sherrill and Russ, J. Am. Chem. Soc. 1907, 29. 1674.)

Solubility of Ag_2CrO_7 in $\text{HNO}_3 + \text{Aq}$ at 25° .

Mols. HNO_3 per l.	Milliat. per l.		Solid Phase
	Cr	Ag	
0	32.20	5.390	$\text{Ag}_2\text{CrO}_4 + \text{Ag}_2\text{CrO}_7$
0.01	25.06	6.131	"
0.02	20.21	7.148	"
0.04	13.59	9.529	"
0.06	11.10	11.10	$\text{Ag}_2\text{Cr}_2\text{O}_7$
0.08	11.10	11.10	"
0.08 + 0.1 AgNO_3	6.624	"

(Sherrill and Russ, J. Am. Chem. Soc. 1907, 29. 1664.)

Silver uranyl chromate, $2\text{Ag}_2\text{CrO}_4, \text{UO}_2\text{CrO}_4$. Ppt. (Formánek, A. 257. 110.)

Silver chromate ammonia, $\text{Ag}_2\text{CrO}_4, 4\text{NH}_3$.

Decomp. by H_2O . Sol. in warm conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Mitscherlich, Pogg. 12. 141.)

Silver dichromate mercuric cyanide, $\text{Ag}_2\text{Cr}_2\text{O}_7, \text{Hg}(\text{CN})_2$.

Sol. in cold H_2O ; very sol. in hot H_2O without decomp. (Krüss, Z. anorg. 1895, 8. 456.)

$\text{Ag}_2\text{Cr}_2\text{O}_7, 2\text{Hg}(\text{CN})_2$. Scarcely sol. in cold, more readily in hot H_2O . Sol. in hot $\text{HNO}_3 + \text{Aq}$, separating on cooling. (Darby, Chem. Soc. 1. 24.)

Sodium chromate, basic, $\text{Na}_4\text{CrO}_6 + 13\text{H}_2\text{O}$.

Sol. without decomp. in H_2O .

Sat. solution at 30° contains 41.3%

(Sherrill and Russ, J. Am. Chem. Soc. 1907, 29. 1663.)

Na_2CrO_4 . (Schreinemakers, Z. phys. Ch. 1906, **55**. 93.)

Deliquescent.

Solubility in H_2O at t° .

t°	0°	10°	20.5°
% Na_2CrO_4	33.87	35.58	38.05

t°	27.7°	35°	37°
% Na_2CrO_4	40.09	44.09	45.13

(Mylius and Funk, Gm.-K. **3**. I, 1379.)

$\text{Na}_2\text{CrO}_4 + \text{Aq}$ sat. at 18° contains 37.50% Na_2CrO_4 , and has sp. gr. = 1.446. (Mylius and Funk, B. 1900, **33**. 3688.)

Sodium chromate, Na_2CrO_4 .

100 ccm. of solution sat. at 18° contain 54 g. Na_2CrO_4 . (Kohlrausch, B. A. B. 1897. 90.)

Solubility in H_2O at t° .

t°	% Na_2CrO_4
70	55.15
80	55.53
100	55.74

(Mylius and Funk, Gm.-K. **3**. I, 1379.)

$\text{Na}_2\text{CrO}_4 + \text{Aq}$ sat. at 18° contains 40.10% Na_2CrO_4 , and has sp. gr. = 1.432. (Mylius and Funk, B. 1900, **33**. 3686.)

See also +4, 6, and $10\text{H}_2\text{O}$.

Sp. gr. of $\text{Na}_2\text{CrO}_4 + \text{Aq}$ at $t^\circ/4^\circ$.

t°	17.4°	17.1°	20.7°
% Na_2CrO_4	5.76	10.62	14.81
Sp. gr.	1.0576	1.1125	1.1644

(Slotte, W. Ann. 1881, **14**. 18.)

+ $4\text{H}_2\text{O}$. Sat. solution at 30° contains 46.62% Na_2CrO_4 . (Schreinemakers, Z. phys. Ch. 1906, **55**. 93.)

Solubility in H_2O at t° .

t°	% Na_2CrO_4	t°	% Na_2CrO_4
25.6	46.08	49.5	50.93
31.5	47.05	54.5	52.28
36	47.98	59.5	53.39
40	48.97	65	55.23
45	50.20		

(Mylius and Funk, Gm.-K. **3**. I, 1379.)

Solubility in H_2O at t° .

t°	% Na_2CrO_4	Mols. H_2O to 1 mol. anhydrous salt	Mols. anhydrous salt to 100 mols. H_2O
28.9	46.47	10.37	9.64
29.7	46.54	10.34	9.67
31.2	47.08	10.12	9.88

(Salkowski, B. 1901, **34**. 1948.)

+ $6\text{H}_2\text{O}$.

Solubility in H_2O at t° .

t°	% Na_2CrO_4	Mols. H_2O to 1 mol. anhydrous salt	Mols. anhydrous salt to 100 mols. H_2O
17.7	43.65	11.60	8.62
19.2	44.12	11.40	8.77
21.2	44.64	11.16	8.96
23.2	45.27	10.88	9.19
24.7	45.75	10.77	9.37
26.6	46.28	10.45	9.57

(Salkowski, B. 1901, **34**. 1948.)

+ $10\text{H}_2\text{O}$. Deliquescent. (Kopp, A. **42**. 99.) Easily sol. in H_2O . Melts in crystal H_2O at 23° . (Berthelot.)

Sp. gr. of solution sat. at 18° = 1.409, and contains 38.1% Na_2CrO_4 . (Mylius and Funk, B. 1897, **30**. 1718.)

Solubility in H_2O at t° .

t°	% Na_2CrO_4
0	24.04
10	33.41
18.5	41.65
19.5	44.78
21	47.40

(Mylius and Funk, Gm.-K. **3**. I, 1379.)

Sp. gr. of solution at 18° containing 40.1% Na_2CrO_4 = 1.432. (Mylius, B. 1900, **33**. 3688.)

Sl. sol. in alcohol. (Moser.)

100 g. absolute methyl alcohol dissolve 0.345 g. Na_2CrO_4 at 25° . (de Bruyn, Z. phys. Ch. 10. 783.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329.)

Sodium dichromate, $\text{Na}_2\text{Cr}_2\text{O}_7$.

More sol. in H_2O than Na_2CrO_4 .

Solubility in H_2O at t° .

t°	% $\text{Na}_2\text{Cr}_2\text{O}_7$
93°	81.19
98°	81.25

(Mylius and Funk, Gm.-K. **3**. I, 1380.)

Sp. gr. of aqueous solution containing—
1 5 10 15 20 25% $\text{Na}_2\text{Cr}_2\text{O}_7$,
1.007 1.035 1.071 1.105 1.141 1.171

30 35 40 45 50 % $\text{Na}_2\text{Cr}_2\text{O}_7$,
1.208 1.245 1.280 1.313 1.343

(Stanley, C. N. **54**. 194.)

Sp. gr. of sat. solution containing 63.92% $\text{Na}_2\text{Cr}_2\text{O}_7$ at 18° = 1.745. (Mylius and Funk, B. 1900, **33**. 3688.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 829.)

in acetone. (Naumann, B. 1904, 37.)

O. Deliquescent.

in H₂O dissolve at—

5° 30° 80° 100° 139°
9.2 116.6 142.8 162.8 209.7 pts. salt.

(Stanley, C. N. 54. 194.)

Solubility in H₂O at t°

t°	% Na ₂ Cr ₂ O ₇
0	61.98
17	63.82
34.5	67.36
52	71.76
72	76.90
81	79.80

us and Funk, Gm.-K. 3. I, 1380.)

H₂O at 30° dissolve 197.6 g. Na₂Cr₂O₇,
solution at 30° contains 66.4%
%. (Schreinemakers, Z. phys. Ch.
. 97.)

mm. of a solution of sodium dichro-
alcohol contain 5.133 g. Na₂Cr₂O₇ +
19.4°. The solution decomp. rapidly.
r, Zeit. angew. Ch. 1913, 26. 456.)

composition of the hydrates formed by
at different dilutions is calculated
terminations of the lowering of the
produced by Na₂Cr₂O₇ and of the
vity and sp. gr. of Na₂Cr₂O₇ + Aq.
lm. ch. J. 1905, 34. 317.)

trichromate, NaCr₂O₁₀.

escent. Very sol. in H₂O. (Stanley,
. 194.)

. Sat. solution at 30° contains
Cr₂O₁₀. (Schreinemakers, Z. phys.
, 55. 94.)

ity in H₂O at t°.

0° 15° 55° 99°
O₁₀ 80.03 80.44 82.68 85.78

us and Funk, Gm.-K. 3. I, 1380.)

of sat. solution containing 80.6%
at 18° = 2.059. (Mylius and Funk,
33. 3688.)

stochromate, Na₂Cr₂O₁₁ + 4H₂O.

ity in H₂O at t°.

0° 16° 22°
O₁₁ 72.19 74.19 76.01

us and Funk, Gm.-K. 3. I, 1380.)

escent.

olution at 18° contains 74.6%
and has sp. gr. = 1.926. (Mylius
, B. 1900, 33. 3688.)

Sodium uranyl chromate, Na₂CrO₄,
2(UO₂)CrO₄ + 10H₂O.

Easily sol. in H₂O. (Formánek, A. 257.
108.)

100 pts. of the solution in H₂O contain
52.52 pts. of the anhydrous salt at 20°. (Rim-
bach, B. 1904, 37. 482.)

Sodium chromate silicate, Na₂O, Cr₂O₃,
2SiO₂ + 14H₂O.

Not decomp. by HCl + Aq (Singer, Dis-
sert. 1910.)

2Na₂O, 3Cr₂O₃, 6SiO₂. Not decomp. by
boiling conc. acids except HF. (Weyberg,
C. B. Miner, 1908. 519.)

5Na₂O, 2Cr₂O₃, 11SiO₂. (Weyberg.)

3Na₂O, 2Cr₂O₃, 9.58SiO₂. (Weyberg.)

Strontium chromate, SrCrO₄.

Somewhat sol. in H₂O. Sol. in 840 pts.
H₂O (Meschevski, Z. anal. 21. 399); sol. in
831.8 pts. H₂O at 15°. (Fresenius, Z. anal. 29.
419.)

100 cc. H₂O dissolve 0.4651% at 10°;
1% at 20°; 2.417% at 50°; 3% at 100°. (Rei-
chard, Ch. Z. 1903, 27. 877.)

Easily sol. in HCl, HNO₃, or H₂CrO₄ + Aq.

Sol. in 512 pts. 0.5% NH₄Cl + Aq at 15°.

Sol. in 63.7 pts. 1% HC₂H₃O₂ + Aq at 15°.

Sol. in 348.8 pts. solution containing 0.75%
NH₄C₂H₃O₂, 4 drops HC₂H₃O₂, and 6 drops

(NH₄)₂CrO₄ + Aq. (Fresenius.)
100 ccm NH₄Cl + Aq sat. at bpt. dissolve
1 g. SrCrO₄. (Dumesnil, A. ch. 1900, (7) 20.
125.)

50 ccm. alcohol (29%) dissolve 0.0066 g.
SrCrO₄.

50 ccm. alcohol (53%) dissolve 0.001 g.
SrCrO₄. (Fresenius, Z. anal. 30. 672.)

Strontium dichromate, SrCr₂O₇.

Easily sol. in H₂O.

Strontium trichromate, SrCr₂O₁₁ + 3H₂O.

Very deliquescent, and sol. in H₂O. (Preis
and Raymann, B. 13. 340.)

Strontium chromate mercuric hydrogen
chloride, SrCrO₄, 2HgCl₂, HCl.

According to Strömholm is SrCl₂, SrCr₂O₇,
4HgCl₂ + H₂O.

Recryst. from H₂O. (Imbert and Belugon,
Bull. Soc., 1897, (3) 17. 471.)

2SrCrO₄, 6HgCl₂, HCl. (Imbert and Belu-
gon.)

Thallos chromate, TiCrO₄.

100 pts. H₂O dissolve 0.03 pt. at 60°.
(Rupp and Zimmer, Z. anorg. 1902, 33. 157.)

Ppt. Insol. in cold moderately conc.
HC₂H₃O₂ + Aq, or in very dil. HNO₃ + Aq,
and very sl. sol. on boiling therewith. Dil.
NH₄OH, and Na₂CO₃ + Aq have the same
action. Attacked by very dil. HCl + Aq.
Sol. in hot conc HCl + Aq. Decomp. by dil.
H₂SO₄ + Aq. (Carstanjen.)

1 l. KOH + Aq (112 g. per l.) dissolves about 3.5 g. Ti_2CrO_4 on boiling, which separates out on cooling.

Boiling conc. KOH + Aq (31% KOH) dissolves 18 g. Ti_2CrO_4 per litre. (Lepierre and Lachaud, C. R. 113. 196.)

Thalious dichromate, $\text{Ti}_2\text{Cr}_2\text{O}_7$.

Insol. in H_2O , etc. Has the same properties as Ti_2CrO_4 .

Thalious trichromate, $\text{Ti}_2\text{Cr}_3\text{O}_{10}$.

Sol. in 2814 pts. H_2O at 15° , and 438.7 pts. at 100° . (Crookes.)

Thallic chromate.

Ppt.

Thorium chromate, basic, $\text{Th}(\text{OH})_2\text{CrO}_4$.

Ppt.; unstable in solution. (Palmer, Am. Ch. J. 1895, 17. 278.)

Thorium chromate, $\text{Th}(\text{CrO}_4)_2 + \text{H}_2\text{O}$.

Ppt. Sol. in HCl and $\text{NH}_4\text{Cl} + \text{Aq}$. 1 pt. is sol. in 284 pts. H_2O at 22° . (Palmer, Am. Ch. J. 1895, 17. 375 and 278.)

+ $3\text{H}_2\text{O}$. Ppt. (Haber, M. 1897, 18. 689.)

+ $8\text{H}_2\text{O}$. Insol. in H_2O . (Chydenius, Pogg. 119. 54.)

Tin (stannous) chromate.

Ppt. Sol. in dil. acids. (Berzelius.)

Tin (stannic) chromate.

Ppt. (Leykauf, J. pr. 19. 127.)

Uranyl chromate, basic, $\text{UO}_3 \cdot 2(\text{UO}_2)\text{CrO}_4 + 8\text{H}_2\text{O}$.

Ppt. (Orloff, Ch. Z. 1907, 31. 375.)

$\text{UO}_3 \cdot (\text{UO}_2)\text{CrO}_4 + 6\text{H}_2\text{O}$. (Orloff.)

Uranyl chromate, $(\text{UO}_2)_2\text{CrO}_4 + 3\text{H}_2\text{O}$.

1 pt. is sol. in 13.3 pts. H_2O at 15° ; slowly sol. in alcohol to give a solution which is decomp. on boiling. (Orloff, Ch. Z. 1907, 31. 375.)

+ $11\text{H}_2\text{O}$. Very sol. in H_2O . (Formánek, A. 257. 108.)

Yttrium chromate.

Deliquescent. Easily sol. in H_2O . (Berlin.)

Zinc chromate, basic, $4\text{ZnO} \cdot \text{CrO}_3 + 3\text{H}_2\text{O}$.
(Gröger, Z. anorg. 1911, 70. 135.)

+ $5\text{H}_2\text{O}$. Insol. in H_2O ; sol. in hot $\text{H}_2\text{CrO}_4 + \text{Aq}$; slowly sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Malaguti and Sarzeau, A. ch. (3) 9. 431.)

$3\text{ZnO} \cdot \text{CrO}_3 + 2\text{H}_2\text{O}$. (Gröger.)

$2\text{ZnO} \cdot \text{CrO}_3 + \text{H}_2\text{O}$. (Briggs, Z. anorg. 1907, 56. 254.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. Ppt. Insol. in H_2O . Sol. in hot $\text{H}_2\text{CrO}_4 + \text{Aq}$. (Prüssen and Phillipona, A. 149. 92.)

+ $2\text{H}_2\text{O}$. Ppt. Not wholly insol. in H_2O . (Prüssen and Phillipona.)

$3\text{ZnO} \cdot 2\text{CrO}_3 + \text{H}_2\text{O}$. (Gröger.)

Zinc chromate, ZnCrO_4 .

Insol. in H_2O ; very sol. in acids; decomp. by boiling with H_2O . (Schulze, Z. anorg. 1895, 10. 154.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 830.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+ H_2O . (Gröger, Z. anorg. 1911, 70. 135.)

Zinc dichromate, $\text{ZnCr}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Hygroscopic.

Very sol. in H_2O and sl. decomp. by boiling. (Schulze, Z. anorg. 1895, 10. 153.)

Zinc trichromate, $\text{ZnCr}_3\text{O}_{10} + 3\text{H}_2\text{O}$.

Deliquescent; very sol. in H_2O . (Gröger, Z. anorg. 1910, 66. 10.)

Zinc chromate ammonia, $\text{ZnCrO}_4 \cdot \text{NH}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O . (Gröger, Z. anorg. 1908, 58. 417.)

$\text{ZnCrO}_4 \cdot 4\text{NH}_3 + 5\text{H}_2\text{O}$. Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in alcohol and ether. (Malaguti and Sarzeau, A. ch. (3) 9. 431.)

+ $3\text{H}_2\text{O}$. Efflorescent. Decomp. by H_2O . Easily sol. in dil. acids and $\text{NH}_4\text{OH} + \text{Aq}$. (Bieler, A. 151. 223.)

$2\text{ZnO} \cdot 3\text{CrO}_3 \cdot 10\text{NH}_3 + 10\text{H}_2\text{O}$. Ppt. (Malaguti and Sarzeau.)

Zinc dichromate mercuric cyanide, $\text{ZnCr}_2\text{O}_7 \cdot 2\text{Hg}(\text{CN})_2 + 7\text{H}_2\text{O}$.

Very sol. in H_2O . Stable in aqueous solution at 100° . (Krüss, Z. anorg. 1895, 8. 460.)

Perchromic acid

See Perchromic acid.

Chromicomolybdic acid, $\text{Cr}_2\text{O}_3 \cdot 12\text{MoO}_3 + 28\text{H}_2\text{O}$.

Slowly sol. in H_2O . (Hall, J. Am. Chem. Soc. 1907, 29. 708.)

Ammonium chromicomolybdate, $3(\text{NH}_4)_2\text{O} \cdot \text{Cr}_2\text{O}_3 \cdot 12\text{MoO}_3 + 20\text{H}_2\text{O}$.

Sol. in H_2O . (Strüve, J. pr. 61. 457; Hall, J. Am. Chem. Soc. 1907, 29. 695.)

+ $26\text{H}_2\text{O}$. (Marckwald, Dissert, 1896.)

Ammonium barium chromicomolybdate, $(\text{NH}_4)_2\text{O} \cdot 22\text{BaO} \cdot \text{Cr}_2\text{O}_3 \cdot 12\text{MoO}_3 + 20\text{H}_2\text{O}$.

(Hall, J. Am. Chem. Soc. 1907, 29. 707.)

Barium chromicomolybdate, 4BaO , Cr_2O_3 , $12\text{MoO}_3 + 15\text{H}_2\text{O}$; 4BaO , Cr_2O_3 , $12\text{MoO}_3 + 18\text{H}_2\text{O}$; 5BaO , Cr_2O_3 , $12\text{MoO}_3 + 16\text{H}_2\text{O}$.

Ppts. (Hall, J. Am. Chem. Soc. 1907, 29. 705.)

Lead chromicomolybdate, 4PbO , Cr_2O_3 , $12\text{MoO}_3 + 22\text{H}_2\text{O}$, and $+24\text{H}_2\text{O}$.

Ppts. (Hall, J. Am. Chem. Soc. 1907, 29. 706.)

Mercurous chromicomolybdate, $8\text{Hg}_2\text{O}$, Cr_2O_3 , $12\text{MoO}_3 + 16\text{H}_2\text{O}$.

Ppt. (Hall, J. Am. Chem. Soc. 1907, 29. 707.)

Potassium chromicomolybdate, K_2O , Cr_2O_3 , 3MoO_3 .

Sol. in $\text{HCl} + \text{Aq}$ with evolution of Cl . (Bradbury, Z. anorg. 1894, 7. 46.)

$3\text{K}_2\text{O}$, Cr_2O_3 , $12\text{MoO}_3 + 20\text{H}_2\text{O}$. Sol. in 38.51 pts. H_2O at 17° . (Strüve; Hall.) $+24\text{H}_2\text{O}$. (Hall.)

$4\text{K}_2\text{O}$, Cr_2O_3 , $12\text{MoO}_3 + 15\text{H}_2\text{O}$. (Hall, J. Am. Chem. Soc. 1907, 29. 709.)

$7\text{K}_2\text{O}$, $2\text{Cr}_2\text{O}_3$, $24\text{MoO}_3 + 32\text{H}_2\text{O}$. (Hall.)

Silver chromicomolybdate, $5\text{Ag}_2\text{O}$, Cr_2O_3 , $12\text{MoO}_3 + 17\text{H}_2\text{O}$.

Ppt. (Hall.)

Sodium chromicomolybdate, $3\text{Na}_2\text{O}$, Cr_2O_3 , $12\text{MoO}_3 + 21\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O . (Strüve.)

Chromic sulphuric acid.

See Sulphochromic acid.

Chromicyanhydric acid,

$\text{H}_2\text{Cr}(\text{CN})_6(?)$.

Insol. in H_2O . (Kaiser, A. Suppl. 3. 163.)

Ammonium chromicyanide, $(\text{NH}_4)_2\text{Cr}(\text{CN})_6$.

Easily sol. in H_2O . (Kaiser, A. Suppl. 3. 163.)

Cupric chromicyanide, $\text{Cu}_2[\text{Cr}(\text{CN})_6]_2$.

Ppt. Insol. in dil. or conc. acids, except on heating. Insol. in NH_4OH , or $\text{KOH} + \text{Aq}$. (Kaiser.)

Lead chromicyanide, basic, $3\text{Pb}(\text{CN})_2$, $2\text{Cr}(\text{CN})_6$, $\text{Pb}(\text{OH})_2$.

Ppt. Sol. in HNO_3 , $\text{NaOH} + \text{Aq}$, or Pb salts $+ \text{Aq}$. (Kaiser.)

Potassium chromicyanide, $\text{K}_2\text{Cr}(\text{CN})_6$.

Very sol. in H_2O .

100 pts. cold H_2O dissolve 30.9 pts. salt.

Insol. in absolute alcohol, but somewhat sol. in dil. alcohol.

Sol. in conc. H_2SO_4 without decomp. (Kaiser, A. Suppl. 3. 170.)

Silver chromicyanide, $\text{Ag}_2\text{Cr}(\text{CN})_6$.

Insol. in all solvents, excepting $\text{KCN} + \text{Aq}$. (Kaiser.)

Sol. in large excess of $\text{HCl} + \text{Aq}$. Sl. sol. in cold, easily sol. in hot conc. HNO_3 . Very sol. in conc. H_2SO_4 . Insol. in hot or cold acetic acid. (Cruser, Dissert. 1896.)

Chromisulphocyanhydric acid.

$\text{H}_2\text{Cr}(\text{SCN})_6$.

Known only in aqueous solution.

Ammonium chromisulphocyanide,

$(\text{NH}_4)_2\text{Cr}(\text{SCN})_6 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O . (Rössler, A. 141. 185.)

Barium chromisulphocyanide, $\text{Ba}_2[\text{Cr}(\text{SCN})_6]_2 + 16\text{H}_2\text{O}$.

Deliquescent, and sol. in H_2O . (R.)

Lead chromisulphocyanide, $\text{Pb}_2[\text{Cr}(\text{SCN})_6]_2$, $4\text{PbO}_2 \cdot \text{H}_2 + 8\text{H}_2\text{O}$.

Insol. in H_2O , but decomp. thereby into— $\text{Pb}_2[\text{Cr}(\text{SCN})_6]_2$, $4\text{PbO}_2 \cdot \text{H}_2 + 5\text{H}_2\text{O}$. Insol. in H_2O .

Potassium chromisulphocyanide, $\text{K}_2\text{Cr}(\text{SCN})_6 + 4\text{H}_2\text{O}$.

Sol. in 0.72 pt. H_2O , and 0.94 pt. alcohol.

Silver chromisulphocyanide, $\text{Ag}_2\text{Cr}(\text{SCN})_6$.

Insol. in H_2O or conc. $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in $\text{KCN} + \text{Aq}$.

Sodium chromisulphocyanide, $\text{Na}_2\text{Cr}(\text{SCN})_6 + 7\text{H}_2\text{O}$.

Deliquescent; sol. in H_2O .

Chromium,

Two modifications—(a) Not attacked by H_2O . Easily sol. in cold $\text{HCl} + \text{Aq}$. Sl. sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Deville.) Easily sol. in a hot mixture of 1 pt H_2SO_4 and 20 pts. H_2O . (Regnault, A. ch. 62. 357.) Easily sol. in warm conc. H_2SO_4 . (Gmelin.) Very slowly sol. in hot $\text{HNO}_3 + \text{Aq}$. (Vauquelin.) Insol. in dil. or conc. $\text{HNO}_3 + \text{Aq}$. (Deville.) Very slowly (Richter), not at all (Berzelius) sol. in hot aqua regia. Easily sol. in $\text{HF} + \text{Aq}$.

(β) Insol. in all acids, even aqua regia (Fremy); probably contains Si.

Pure Cr is sol. in conc. H_2SO_4 , HCl and dil. HNO_3 ; sol. in $\text{HgCl}_2 + \text{Aq}$.

Insol. in fuming HNO_3 and aqua regia. (Moissan, C. R. 1894. 119, 187.)

Cr prepared by aluminothermic method is sol. in haloid acids to form chromic and chromous salts, even in absence of air. (Doring, J. pr. 1902, (2) 66. 65; 1906, (2) 73. 393.)

Aluminothermic Cr is active in contact

with HCl, HBr, HI, HF, H_2SO_4 , $\text{H}_2\text{C}_2\text{O}_4$, i. e., sol. in cold conc. or warm dil. acids. Is inactive in contact with conc. HNO_3 , $\text{H}_2\text{Cr}_2\text{O}_4$, HClO_3 , HClO_4 , H_3PO_4 , KOH, citric, formic, acetic and tartaric acids. Cause attributed to a different electric state. (Hittorff, Z. phys. Ch. 1898, 25. 729.)

Chromium ammonia compounds.

See—

Bromotetramine chromium compounds,
 $\text{BrCr}(\text{NH}_3)_4\text{X}_2$.

Bromopurpureochromium compounds,
 $\text{BrCr}(\text{NH}_3)_4\text{X}_2$.

Chlorotetramine chromium compounds,
 $\text{ClCr}(\text{NH}_3)_4\text{X}_2$.

Chloropurpureochromium compounds,
 $\text{ClCr}(\text{NH}_3)_4\text{X}_2$.

Diamine chromium sulphocyanides,
 $\text{Cr}(\text{NH}_3)_2(\text{SCN})_4\text{M}$.

Erythrochromium compounds,
 $(\text{HO})\text{Cr}_2(\text{NH}_3)_{10}\text{X}_2$.

Iodopurpureochromium compounds,
 $\text{ICr}(\text{NH}_3)_4\text{X}_2$.

Iodotetramine chromium compounds,
 $\text{ICr}(\text{NH}_3)_4\text{X}_2$.

Luteochromium compounds, $\text{Cr}(\text{NH}_3)_6\text{X}_3$.

Rhodochromium compounds,
 $(\text{HO})\text{Cr}_2(\text{NH}_3)_{10}\text{X}_2$.

Rhodochromium compounds,
 $(\text{HO})_3\text{Cr}_2(\text{NH}_3)_6\text{X}_3$.

Roseochromium compounds,
 $\text{Cr}(\text{NH}_3)_6(\text{OH})_2\text{X}_4$.

Xanthochromium compounds,
 $(\text{HO}_2)\text{Cr}(\text{NH}_3)_6\text{X}_2$.

Chromium arsenide, CrAs .

Insol. in mineral acids. (Dieckmann, Z. anorg. 1914, 86. 294.)

Cr_2As_3 . Insol. in mineral acids. (Dieckmann.)

Chromium azoimide, CrN_3 .

Pptd. by addition of alcohol and ether. Insol. in H_2O . (Curtius, J. pr. 1900, (2) 61. 410.)

Chromium boride, CrB .

Insol. in HCl, dil. H_2SO_4 , HF, HF + HNO_3 . Sl. sol. in HNO_3 and in aqua regia. (Wedekind, B. 1907, 40. 299.)

Sol. in cold dil. or conc. HCl, HF, and H_2SO_4 . (Jassonneix, C. R. 1906, 143. 1151.)

Cr_2B_3 . Sol. in conc. or dil. HF, HCl, H_2SO_4 ; insol. in HNO_3 or alkalis + Aq. (Jassonneix.)

Chromous bromide, CrBr_2 .

Sol. in H_2O . Not deliquescent in dry air. (Moissan, C. R. 92. 1051.)

Chromic bromide, CrBr_3 .

Anhydrous. Insol. in H_2O , but d at once in presence of the least trace of (Bauck, A. 111. 382.)

+ $6\text{H}_2\text{O}$. Deliquescent. Very sol. H_2O dissolves more than 2 pts. cry ord. temp. Very sol. in alcohol. 1 ether. (Recoura, C. R. 110. 1029.)

Blue modification. Insol. in (Recoura, C. R. 110. 1193.)

Very hygroscopic. Easily sol. in and acetone. Insol. in ether. (Wer 1902, 322. 343.)

+ $8\text{H}_2\text{O}$. Sol. in H_2O . (Varenne, C 727.)

Chromium molybdenyl bromide, $\text{CrMo}_3\text{O}_4\text{Br}_4$.

Apparently wholly insol. in dil. Sol. in hot conc. HCl + Aq with d Insol. in M_2CrO_4 + Aq. (Atterberg, + $2\text{H}_2\text{O}$. Apparently wholly in dil. acids.

Sol. in hot conc. HCl + Aq with d Insol. in M_2CrO_4 + Aq. (Atterberg.)

Chromic rubidium bromide, CrBr_3 , + H_2O .

Sol. in H_2O with decomp. (Wer 1902, 322. 345.)

Chromic bromide ammonia.

See Bromotetramine chromium bro

Chromous bromide hydrazine, $\text{CrBr}_2, 2\text{N}_2\text{H}_4$.

Insol. in H_2O . Sol. in acids. In alcohol, ether and similar solvents. (B. 1913, 46. 1507.)

Chromium carbide, Cr_3C .

(Moissan, C. R. 1894, 119. 187.)

Cr_3C_2 . Does not decomp. H_2O at a temp. or at 100° ; insol. in conc. HCl, and aqua regia; sol. in dil. HCl (slowly) in fused KOH; sol. in fused KNO_3 . (M Bull. Soc. 1894, (3) 117. 1016.)

Chromium iron carbide, $3\text{Fe}_3\text{C}, 2\text{Cr}_3\text{C}$

Decomp. by H_2O ; sol. in gaseo dracids; insol. in HNO_3 and aqua regia. liams, C. R. 1898, 127. 484.)

Chromium tungsten carbide, CW_2 :

Not attacked by acids.

Slowly attacked by fused KOH or carbonates. Rapidly decomp. by alkali nitrates or KClO_3 . (Moissan, 1903, 137. 294.)

Chromous chloride, CrCl_2 .

Deliquescent. Very sol. in H_2O with tion of much heat. (Moberg, J. pr. 21

ally insol. in ether. Moderately solute alcohol, methyl alcohol, and yde. (Rohland, Z. anorg. 1899, 21.

,O. (Moissan, A. ch. (5) 25. 40.)
l. (Knight and Rich, Chem. Soc. 89.)
l. (Knight and Rich.)

hydrogen chloride, 3CrCl₃, 2HCl + O.

p. by H₂O. (Recoura, C. R. 100.

in liquid NH₃. (Gore, Am. Ch. J. 827.)

chloride, CrCl₃.

ous.—Peach-blossom-colored modi- Insol. in pure H₂O (Peligot), but continued boiling of the finely divided H₂O, traces are dissolved with Not decomp. by boiling conc. other acids, even aqua regia.

sol. with evolution of heat in H₂O g only 1/40,000 pt. CrCl₃. (Peligot, J.). Also sol. in presence of traces of mg. SnCl₂ cause 1 g. CrCl₃ to dis- eCl₂, Cu₂Cl₂, Na₂S₂O₃, and other substances; chlorides without re- operties have no effect. (Pelouze, A. . 251). TiCl₃ and SO₂ have similar rtion. (Ebelmen, A. ch. (3) 20. 390); dil. acids. (Moberg.)

in dil. alkalies + Aq; very slowly by boiling conc. alkalies or alkali s + Aq. (Fellenberg, Pogg. 50. 76.) lty sol. in methyl acetate. (Nau- 1909, 42. 3790.)

n CS₂. (Arctowski, Z. anorg. 1894,

n acetone. (Eidmann, C. C. 1899.

ally insol. in absolute ethyl alcohol, alcohol, acetaldehyde and ether. , Z. anorg. 1899, 21. 39.)

Sl. sol. in benzonitrile. (Nau- 1914, 47. 1369.)

modification. Very sol. in H₂O to reen solution. (Moberg, J. pr. 44.

iolet mod. is almost insol. in H₂O ,000 pt. chromous chloride is present, ly sol. (Rohland, Z. anorg. 1899, 21.

). Sl. deliquescent. Very sol. in hol, and ethyl acetate. (Godeffroy, . (2) 43. 229.)

). Deliquescent. Sol. in H₂O, but decomp. to CrOCl₂.

ally insol. in ether. Moderately solute ethyl alcohol, methyl alcohol taldehyde. (Rohland, Z. anorg. 39.)

chlorochromic chloride" is sol. in l fuming HCl(1:1). (Bjerrum, B. 1599.)

Green modification.

Solubility in H₂O at 25°.

25 g. green CrCl₃ + 6H₂O and 10 g. H₂O.

Time	Total Solu- bility in %	Composition of the sat. solution	
		% violet salt	% green salt
1/4 hr.	58.36	8.30	91.70
1/2 hr.	...	12.57	87.43
4 hrs.	63.27	24.80	75.20
1 day	68.50	37.64	62.36
2 days	...	40.90	59.10
3 "	68.95	42.78	57.22
11 "	...	42.84	57.16
13 "	...	42.39	57.61
19 "	68.58	42.62	57.38

(Olie, Z. anorg. 1906, 51. 55.)

Solubility of green CrCl₃ + 6H₂O in H₂O at 32°.

10 g. CrCl₃ + 6H₂O and 4 g. H₂O.

Time	Total solubility in %	Composition of the dissolved substance		Solid phase
		% violet salt	% green salt	
7'	63.69	12.87	87.13	Almost all dissolved
45'	66.24	21.43	78.57	
2h 5'	69.53	34.53	65.47	
48h	69.33	45.27	54.73	
* 11 dys.	70.81	45.27	54.73	

* First 8 days at 35°.

(Olie, Z. anorg. 1907, 53. 276.)

Solubility of green CrCl₃ + 6H₂O in H₂O at 35°.

10 g. CrCl₃ + 6H₂O and 3.3 g. H₂O.

Time	Total solubil- ity in %	Composition of the dissolved substance	
		% violet salt	% green salt
8'	65.85	16.47	83.53
38'	66.74	25.02	74.98
1h	66.21	25.45	74.55
2h 10'	68.90	31.47	68.53
4h	70.79	36.28	63.72
23h	71.34	42.95	57.05
72h	70.79	42.88	57.12

(Olie, l. c.)

If a solution saturated with the green hexa- hydrate below 32° is cooled, the decahydrate separates out; if the solution is saturated above 32°, both the decahydrate and hexa- hydrate separate out on cooling. (Olie, l. c.)

*Violet modification.*Solubility in H_2O at 25° .25 g. violet $CrCl_3 + 6H_2O$ and 10 ccn. of a 35% solution of green $CrCl_3 + 6H_2O$.

Time	Total Solubility in %	Composition of the sat. solution	
		% violet salt	% green salt
1 1/2 hr.	65.49	84.05	15.95
5 "	..	84.47	15.53
29 "	..	78.16	21.84
2 dys.	70.47	73.19	26.81
4 "	..	68.71	31.29
5 "	76.38	60.66	39.34
5 dys, 6 hrs.	..	60.36	39.64
6 dys.	..	65.10	34.90
8 "	73.26	65.80	34.20
10 "	..	58.08	41.92
12 "	71.14	41.40	58.60

(Olie, Z. anorg. 1906, 51. 57.)

Solubility in H_2O at 25° .25 g. violet $CrCl_3 + 6H_2O$ and 10 g. H_2O .

Time	Total Solubility in %	Composition of the sat. solution	
		% violet salt	% green salt
1/2 hr.	61.99	98.47	1.53
4 1/2 hrs.	..	96.70	3.30
1 dy.	63.88	91.54	8.46
2 "	..	83.37	16.63
4 "	70.68	69.11	30.89
5 "	..	62.20	37.80
7 "	72.11	62.72	37.28
8 "	..	54.03	45.97
12 "	..	46.39	53.61
13 "	..	47.66	52.34
26 "	70.62	48.55	51.45

(Olie, l. c.)

+6 1/2 H_2O . *Green modification.* 100 pts. H_2O dissolve 130 pts. salt at 15° . Sol. in alcohol. (Recours, C. R. 102. 518.)

Grayish-blue modification. Very sol. in H_2O (Recours, C. R. 102. 548.)

+10 H_2O . Very deliquescent; melts in crystal H_2O at $6-7^\circ$. Very sol. in H_2O , alcohol, and ethyl acetate. (Godeffroy.)

Easily sol. in H_2O ; can be recryst. from H_2O . Sol. in alcohol and ether. (Werner, B. 1906, 39. 1827.)

*Green modification.*Solubility of green $CrCl_3 + 10H_2O$ in H_2O at 29° .14.2 g. $CrCl_3 + 10H_2O$ and 2.5 g. E

Time	Total solubility in %	Composition of the dissolved substance		Solid
		% violet salt	% green salt	
7'	61.35	8.71	91.29	$CrCl_3$
20'	62.46	9.90	90.10	
1 ^h 55'	65.04	25.05	74.95	
4 ^h 30'	67.41	32.90	67.10	$CrCl_3$
24 ^h	69.44	42.93	57.07	all diss.
28 ^h	..	31.78	68.22	$CrCl_3$
29 ^h	67.59	33.65	66.36	$CrCl_3$
48 ^h	69.42	42.17	57.83	
72 ^h	68.69	43.80	56.20	

(Olie, Z. anorg. 1907, 53. 275.)

The composition of the hydrates by $CrCl_3$ at different dilutions is calculated from determinations of the lowering fr. pt. produced by $CrCl_3$ and of the conductivity and sp. gr. of $CrCl_3 + Aq$. (Am. Ch. J. 1905, 34. 310.)

Chromic glucinum chloride, $CrCl_3 + H_2O$.

Sol. in H_2O with decomp. (Neumann 244. 329.)

Chromic lithium chloride, $CrCl_3 + 4H_2O$.

Very hygroscopic.

Sol. in ice water but solution soon decomp.

Easily sol. in alcohol. (Werner, B. 34. 1603.)

$[Cr(OH_2)Cl_2]Li_2 + 4H_2O$.

Very hygroscopic. Sol. in ice and in alcohol (Werner, B. 1901, 34.)

Chromic magnesium chloride, $CrCl_3 + H_2O$.

Decomp. by H_2O . (Neumann.)

Chromic phosphoric chloride, $CrCl_3$.

Decomp. by H_2O . (Cronander.)

Chromium platinum chloride.

See chloroplatinate, chromium.

Chromic potassium chloride, $CrCl_3$.

Decomp. by H_2O .

$CrCl_3 + 2KCl + H_2O$. (Neumann, A. 329.)

$CrCl_3 + 3KCl$. Easily sol. in H_2O with decomp. (Fremy, A. ch. (3) 12. 361.)

rubidium chloride, CrCl_3 , $2\text{RbCl} +$
 p. by H_2O . (Neumann, A. 244.)

sol. in cold, rapidly sol. in hot H_2O
 comp. (Werner, B. 1901, 34. 1603.)
 $3\text{RbCl} + 8\text{H}_2\text{O}$. Unstable. Decomp.
 sol. (Werner, B. 1906, 39. 1830.)

sodium chloride, CrCl_3 , NaCl .
 H_2O . (Berzelius.)
 3NaCl . Sol. in H_2O . (Berzelius.)

thallium chloride, CrCl_3 , 3TlCl .
 with decomp. in H_2O . (Neumann, A.
)

chloride ammonia.

chlorotetramine chromium chloride.

is chloride hydrazine, CrCl_3 ,
 H_4 .
 in H_2O . Sol. in acids. Insol. in
 ether and similar solvents. (Traube,
 46. 1506.)

chloride ferric oxide.

is easily sol. in dil., difficultly sol.
 $\text{CrCl}_3 + \text{Aq}$. (Béchamp, A, ch. (3)

is fluoride, CrF_3 .

. in H_2O , hot H_2SO_4 or dil. HNO_3 .
 in boiling HCl . Insol. in alcohol.
 , C. R. 1893, 116. 254.)

fluoride, CrF_3 .

tly sol. in H_2O . (Berzelius.)
 in liquid NH_3 . (Gore, Am. Ch. J.
 827.)
 in methyl acetate. (Naumann, B.
 3790); ethyl acetate. (Naumann,
 43. 314.)

O. Insol. in H_2O . (Werner and
 escu, B. 1908, 41. 4243.)
 H_2O . Sol. in H_2O . (Poulenc, C. R.
 6. 255.)

O. Sl. sol. in H_2O . (Werner and
 escu, B. 1908, 41. 4242.)

O. *Violet modification*. Very sl. sol.
 Insol. in alcohol. Sol. in HCl , and
 Aq. (Fabris, Gazz. ch. it. 20. 582.)

um hexafluoride, CrF_6 .

ap. by H_2O with evolution of heat.
 is.)
 at composition is CrO_2F_2 . (Oliveri,
 i. it. 16. 218.)

cobaltous fluoride, CrF_3 , $\text{CoF}_2 +$
 O .

sol. in H_2O . (Petersen, J. pr. (2)

Chromic cupric fluoride, $\text{CrCuF}_4 + 5\text{H}_2\text{O}$.

Can be cryst. from $\text{HF} + \text{Aq}$. (Higley, J.
 Am. Chem. Soc. 1904, 26. 630.)

Chromic nickel fluoride, CrF_3 , $\text{NiF}_2 + 7\text{H}_2\text{O}$.

Somewhat more sol. in H_2O than CrF_3 ,
 $\text{CoF}_2 + 7\text{H}_2\text{O}$. (Petersen, J. pr. (2) 40. 61.)

Chromic potassium fluoride, CrF_3 , 3KF .

Nearly insol. in H_2O . (Christensen, J.
 pr. (2) 35. 161.)

CrF_3 , $2\text{KF} + \text{H}_2\text{O}$. Nearly insol. in H_2O .
 Sol. in conc. $\text{HCl} + \text{Aq}$. (Christensen.)

Chromic sodium fluoride, CrF_3 , $2\text{NaF} + \text{H}_2\text{O}$.

(Wagner, B. 19. 896.)

Chromic thalious fluoride, 2CrF_3 , 3TlF .

Sol. in hot H_2O , less sol. in cold. Sl. sol.
 in HF . (Ephraim, Z. anorg. 1909, 61. 242.)

Chromic zinc fluoride, CrF_3 , $\text{ZnF}_2 + 7\text{H}_2\text{O}$.

Can be cryst. from $\text{HF} + \text{Aq}$. (Higley, J.
 Am. Chem. Soc. 1904, 26. 630.)

Colloidal solution is perfectly clear.
 (Biltz, B. 1902, 35. 4433.)

Chromous hydroxide, CrO_2H_2 .

Decomp. by H_2O , especially if hot. (Pe-
 ligot, A. ch. (3) 12. 539.)

Slowly sol. in cold conc. acids, even aqua
 regia; almost insol. in dil. acids. (Moberg, J.
 pr. 43. 119.)

Chromic hydroxide, Cr_2O_3 , $x\text{H}_2\text{O}$, probably
 $\text{Cr}_2\text{O}_3\text{H}_6$.

Insol. in H_2O . Easily sol. in acids. Easily
 sol. in cold KOH , or $\text{NaOH} + \text{Aq}$; much less
 sol. in cold $\text{NH}_4\text{OH} + \text{Aq}$; the presence of
 NH_4Cl has no influence upon solubility
 in $\text{NH}_4\text{OH} + \text{Aq}$. (Fresenius.) Insol. in
 $\text{NH}_4\text{OH} + \text{Aq}$ if it has been thoroughly
 washed.

Insol. in $\text{KCN} + \text{Aq}$, but sl. sol. in $\text{KCN} +$
 $\text{HCN} + \text{Aq}$. (Rodgers, 1834.)

Gradually sol. in dil. $\text{FeCl}_3 + \text{Aq}$; after three
 months, 2 mols. $\text{Cr}_2\text{O}_3\text{H}_6$ are dissolved by 1
 mol. FeCl_3 without pptn. of $\text{Fe}_2\text{O}_3\text{H}_6$. (Bé-
 champ, A. ch. (3) 57. 296.)

Also sol. in $\text{CrCl}_3 + \text{Aq}$; in four months, $1\frac{1}{2}$
 mols. $\text{Cr}_2\text{O}_3\text{H}_6$ are dissolved by 1 mol. CrCl_3 .
 (Béchamp.)

Sol. in $\text{Cr}(\text{NO}_3)_3 + \text{Aq}$, and clear solution
 formed as long as 3 mols. HNO_3 are present
 for 8 mols. Cr_2O_3 . (Ordway, Sill. Am. J. (2)
 27. 197.)

Chromic hydroxide, pptd. by alkalis is
 easily sol. in excess of the reagent; after being
 dried in a vacuum, however, it is insol. in
 alkalis. (Herz, Z. anorg. 1901, 28. 344.)

Freshly pptd. it is sol. in aq. alkali, but it
 is readily changed into a modification which is
 insol. (Herz, Z. anorg. 1902, 31. 352.)

The solubility of chromic hydroxide in an

aqueous solution of red chromic chloride is not directly proportional to the concentration of the latter—a basic chloride is probably formed. (Fischer, Z. anorg. 1904, 40: 43.)

Not pptd. in presence of Na citrate. (Spiller.)

Insol. in amylamine+Aq; not pptd. in presence of alkali tartrates, sugar, etc.

$\text{Cr}_2\text{O}_3 \cdot \text{H}_2\text{O} + 4\text{H}_2\text{O}$. Difficultly sol. in acids.

$\text{Cr}_2\text{O}_3 \cdot \text{H}_2\text{O} + \text{H}_2\text{O}$. Extremely hygroscopic.

Exists in a soluble modification, obtained by dialysis; solution can be diluted with pure H_2O , but gelatinizes with traces of salts. (Graham, Roy. Soc. Trans. 1861. 183.)

$\text{Cr}_2\text{O}_3(\text{OH})_2$. Insol. in boiling dil. $\text{HCl} + \text{Aq}$.

$\text{Cr}_2\text{O}(\text{OH})_4$ (Guignet's green). Scarcely sol. in boiling $\text{HCl} + \text{Aq}$. (Salvétat, C. R. 48. 295.)

Guignet gave formula as $2\text{Cr}_2\text{O}_3 + 3\text{H}_2\text{O}$.

Chromochromic hydroxide, $\text{Cr}_2\text{O}_4, \text{H}_2\text{O}(?)$.

Slightly attacked by acids. (Peligot, A. ch. (3) 12. 539.)

Chromous iodide, CrI_2 .

Easily sol. in H_2O . (Moissan, A. ch. (5) 25. 401.)

Chromic iodide, $\text{CrI}_3(?)$.

Insol. in cold, sol. in hot H_2O , but no separation occurs on cooling. (Berlin.)

$+9\text{H}_2\text{O}$. Hygroscopic. Sol. in alcohol and acetone. Insol. in CHCl_3 . (Higley, J. Am. Chem. Soc. 1904, 26. 628.)

Chromous iodide hydrazine, $\text{CrI}_2, 2\text{N}_2\text{H}_4$. (Traube, B. 1913, 46. 1507.)

Chromium nitride, CrN .

Insol. in dil. acids and alkalis, conc. HNO_3 , HCl , or $\text{HF} + \text{Aq}$, even on heating. Slowly sol. in hot aqua regia or cold H_2SO_4 . Sol. in cold solutions of alkali hypochlorites. (Ufer, A. 112. 281.)

Insol. in HCl , HNO_3 , and aqua regia. (Férée, Bull. Soc. 1901, (3) 25. 618.)

Unacted upon by acids at ordinary temperatures. (Smits, Chem. Soc. 1897, 72. (2) 33.)

Cr_3N_2 . Slowly attacked by conc. HNO_3 , and by $\text{HNO}_3 + \text{HCl}$. All other reagents are without action. (Henderson and Galletly, J. Soc. Chem. Ind. 1908, 27. 387.)

CrN_3 . See Chromium azoimide.

Chromous oxide, CrO .

Insol. in HNO_3 and dil. $\text{H}_2\text{SO}_4 + \text{Aq}$.

Sol. in HCl . (Férée, Bull. Soc. 1901, (3) 25. 619.)

Chromic oxide, Cr_2O_3 .

When ignited is nearly insol. in a dissolves in H_2SO_4 by long boiling in liquid HCl . (Gore.)

Insol. in acetone. (Fidmann, C. II, 1014.)

Solubility in (calcium succate + Aq.

1 l. solution containing 418.6 g. 34.3 g. CaO dissolves 1.07 g. Cr_2O_3 ; tion containing 296.5 g. sugar and 2 dissolves 0.56 g. Cr_2O_3 ; 1 l. solution ing 174.4 g. sugar and 14.1 g. CaC 0.20 g. Cr_2O_3 . (Bodenbender, J. 600.)

See also Chromic hydroxide.

$+ \text{H}_2\text{O}$. The compound to which gave the formula Cr_2O_4 . Insol. in easily attacked by HNO_3 . (Férée, 1901, (3) 25. 620.)

Chromochromic oxide, $\text{Cr}_2\text{O}_4 = \text{CrI}$

Known only in form of hydrox see.

$+3\text{H}_2\text{O}$. Stable in dry air. D moist air. (Baugé, C. R. 1898, 127.

Cr_2O_4 , or Cr_2O_6 (?). Insol. in aqua regia. (Bunsen, Pogg. 91. 6

Not obtainable. (Geuther, A. 11

Formula is $\text{Cr}_2\text{O}_3 + \text{H}_2\text{O}$. (Férée

Chromium trioxide, CrO_3 .

Deliquescent, and very sol. in form solution of H_2CrO_4 or $\text{H}_2\text{Cr}_2\text{O}_7$

Solubility in H_2O at t° .

t°	0°	15°	50°
% CrO_3	62.08	62.38	64.

(Mylius and Funk, Gm. K. 3. 1

Sat. $\text{CrO}_3 + \text{Aq}$ contains at:

0°	20°	60°
61.54	62.52	65.12% (

(Koppel and Blumenthal, Z. anorg 228.)

The system $\text{CrO}_3 - \text{H}_2\text{O}$ has be at temp. from 0° to -74° . In th concentration investigated, from CrO_3 , no hydrate of CrO_3 cryst aq. solution. (Kremann, M. 1911,

Sat. $\text{CrO}_3 + \text{Aq}$ contains at:

82°	100°	115°
66	67.4	68.4% Cr

(Kremann, M. 1911, 32. 6

Solubility in H ₂ O at°.		
t°	% by wt. CrO ₃	Solid phase.
9°	3.6	Ice
9	7.8	"
7	11.5	"
8	14.1	"
95	24.9	"
7	25.2	"
75	33.5	"
25	39.2	"
5	49.1	"
	53.3	"
	61.7	CrO ₃
	62.24	"
8	62.88	"
	63.50	"
	64.83	"
	68.5	"
2	70.7	"
196	100	"

r and Prins, Z. phys. Ch. 1912, 81. 114.)

Sp. gr. of CrO ₃ +Aq at t°.		
	Sp. gr.	% CrO ₃
	1.0606	8.25
	1.0679	8.79
	1.0694	8.79
	1.0957	12.34
	1.1569	19.33
	1.20269	31.83
	1.20264	31.83
	1.20714	31.83
	1 20940	32.59
	1.21914	32.59
	1.22106	32.59
	1.22384	32.59
	1.3441	37.77
	1.3448	37.82
	1.34416	37.82
	1.7028	62.23

(Zettnow, Pogg. 143. 474.)

of CrO₃+Aq (H₂CrO₄+Aq). M=rding to Mendelejeff at 15°; Z=ac-ing to Zettnow, calculated by Ger-(Z. anal. 27. 300).

M	Z	% CrO ₃	M	Z
.036	1.037	35	1.324	1.312
.076	1.076	40	1.383	1.373
.119	1.118	45	1.445	1.440
.166	1.162	50	1.510	1.512
.215	1.208	55	1.579	1.587
.268	1.258	60	...	1.656

B.-pt. of CrO ₃ +Aq at ord. pressure.	
B.-pt.	G. CrO ₃ in 100 g. of the solution.
102°	10.81
104	24.08
107	36.47
110.5	45.15
116	54.56
120	61.54
127	71.24 sat. solution

(Koppel and Blumenthal, Z. anorg. 1907, 53. 254.)

Sol. in H₂SO₄; the solubility is least when the acid contains 66% H₂SO₄ (Schrötter); 84.5% H₂SO₄ (Bolley).

Very sol. in H₂SO₄ of 1.85 sp. gr. Sl. sol. in cold KHSO₄+Aq. (Fritzsche.)

The statement that CrO₃ is insol. in acids is incorrect. 2.85 g. (ignited) are sol. in HNO₃ to the extent of 2.58 g. 0.81 g. (ignited) are sol. in HNO₃ to the extent of 0.77 g. (Jovitschitsch, M. 1909, 30. 48.)

Practically insol. in POCl₃. (Walden, Z. anorg. 1910, 68. 312.)

Sl. sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

Sol. in alcohol with decomp.

Sol. in anhydrous ether.

Sol. in acetic anhydride. (Fry, J. Am. Chem. Soc. 1911, 33. 702.)

Sol. in acetone. (Naumann, B. 1904, 4328.)

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Chromium oxide, Cr₂O₃=2Cr₂O₃, CrO₃.

Cr₂O₁₅=3Cr₂O₃, 2CrO₃.

CrO₂=Cr₂O₃, CrO₃.

Cr₂O₁₂=Cr₂O₃, 3CrO₃.

Cr₂O₁₈=Cr₂O₃, 4CrO₃.

See Chromate, chromium.

Chromium peroxide, Cr₂O₇(?)

More sol. in ether than in H₂O. Ether solution is somewhat more stable than aqueous solution. (Aschoff, J. pr. 81. 401.)

Formula is CrO₃, H₂O₂. (Moissan, C. R. 97. 96.)

Chromium peroxide ammonia, CrO₄, 3NH₃.

Sol. in H₂O with partial decomp.

Sl. sol. in NH₄OH+Aq.

Sol. with decomp. in 20% acetic acid. (Hofmann, B. 1905, 38. 3060.)

Chromium tetroxide potassium cyanide, CrO₄, 3KCN.

Sol. in H₂O. Insol. in other ordinary solvents. (Wiede, B. 1899, 32, 381.)

Chromic oxychloride.

From Cr_2O_3 . Sol. in H_2O as long as 1 mol. CrCl_3 is present for $2\frac{1}{2}$ mols. $\text{Cr}_2\text{O}_3\cdot\text{H}_2\text{O}$. (Ordway, Sill. Am. J. (2) 27. 197.)

$\text{Cr}_2\text{O}_3, 2\text{CrCl}_3$. Sol. in H_2O . (Kletzinsky, Zeit. Ch. 1866. 277.)

$\text{Cr}_2\text{O}_3, \text{CrCl}_3 = \text{CrOCl}$. Anhydrous. Only partly sol. in H_2O .

+ $3\text{H}_2\text{O}$. Very deliquescent, and sol. in H_2O . (Peligot.)

$\text{Cr}_2\text{O}_3, 4\text{CrCl}_3 + 6\text{H}_2\text{O} = \text{Cr}_2\text{OCl}_4 + 2\text{H}_2\text{O}$. (Peligot, J. pr. 37. 38.)

+ $9\text{H}_2\text{O} = \text{Cr}_2\text{OCl}_4 + 3\text{H}_2\text{O}$. Sol. in H_2O (Moberg); = $\text{Cr}_2(\text{OH})_2\text{Cl}_4 + 2\text{H}_2\text{O}$. (Schiff, A. 124. 157.)

$\text{Cr}_2\text{O}_3, 7\text{CrCl}_3 = \text{Cr}_2\text{OCl}_7$. Very sol. in H_2O with decomp. (Besson and Fournier, C. R. 1909, 148. 1194.)

$\text{Cr}_2\text{O}_3, 8\text{CrCl}_3 + 24\text{H}_2\text{O}$. Sol. in H_2O (Moberg); = $\text{Cr}_2(\text{OH})\text{Cl}_6 + 4\text{H}_2\text{O}$. (Schiff, l. c.)

$(\text{CrO}_2)_2\text{Cl}_2$. (Pascal, C. R. 1909, 148. 1464.)

$\text{Cr}_2\text{O}_3\text{Cl}_4$. Insol. in H_2O . (Pascal, C. R. 1909, 148. 1464.)

From CrO_3 .

See Chromyl chloride.

Chromic oxychloride potassium chloride, $\text{CrOCl}_3, 2\text{KCl}$.

Decomp. in the air.

Sol. in conc. HCl without decomp. (Weinland, B. 1906, 39. 4043.)

Chromic oxychloride rubidium chloride, $\text{CrOCl}_3, 2\text{RbCl}$.

Decomp. in the air.

Sol. in conc. HCl without decomp. (Weinland, B. 1906, 39. 4045.)

Chromium oxyfluoride, CrO_2F_2 .

See Chromyl fluoride.

Chromium phosphide, CrP .

Insol. in acids, but a trace dissolves in aqua regia. Insol. in $\text{HF} + \text{Aq}$. (Berzelius.)

Not attacked by acids or by aqua regia. (Granger, C. N. 1898, 77. 228.)

Insol. in all acids except a mixture of HNO_3 and HF . (Maronneau, C. R. 1900, 130. 658.)

Insol. in mineral acids. (Dieckmann, Z. anorg. 1914, 86. 295.)

Insol. in aqua regia. (Granger, C. R. 1897, 124. 191.)

Cr_2P_3 . Insol. in mineral acids. (Dieckmann, Z. anorg. 1914, 86. 295.)

Chromous selenide, CrSe .

(Moissan, C. R. 90. 817.)

Chromic selenide, Cr_2Se_3 .

Insol. in H_2O . (Moissan, C. R. 90. 817.)

Chromic potassium selenide, $\text{K}_2\text{Cr}_2\text{Se}_4$.

Insol. in HCl . Easily sol. in conc. H . (Milbauer, Z. anorg. 1904, 42. 451.)

Chromium silicide, Cr_3Si .

Sol. in fused KNO_3 ; insol. in cold HCl aqua regia.

Insol. in $\text{HF} + \text{Aq}$. (Moissan, C. R. 121. 625.)

CrSi_2 . Sol. in HF . Insol. in HCl aqua regia. (Chalmot, Am. Ch. J. 189. 69.)

Cr_2Si_3 . Insol. in dil. HCl ; sol. in conc. HCl and in HF ; insol. in HNO_3 , H_2SO_4 . (Lebeau, C. R. 1903, 136. 133.)

Cr_3Si . Sol. in HF ; insol. in other sol. in fused KOH and fused alkali ni and carbonates. (Zettel, C. R. 1898, 834.)

Chromous sulphide, CrS .

Insol. in H_2O or $\text{K}_2\text{S} + \text{Aq}$. (Peligot. Easily sol. in acids. (Moissan, C. R. 817.)

Sol. in cold conc. acids.

Sol. in molten alkalis. (Mourlot, 1895, 121. 944.)

Min. Daubrelite.

Chromic sulphide, Cr_2S_3 .

Insol. in H_2O or alkali sulphides + Ac attacked by $\text{HCl} + \text{Aq}$. (W. Müller, 127. 404.)

$\text{HNO}_3 + \text{Aq}$ decomposes or not acco to method of preparation. Easily dec by aqua regia.

Insol. in caustic alkalis + Aq .

Insol. in $\text{K}_2\text{S} + \text{Aq}$. (Berzelius.)

Chromochromic sulphide, $\text{Cr}_2\text{S}_4 = \text{Cr}_2\text{S}_3$.

Insol. in H_2O , HCl , or dil. H_2SO_4 . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Gröger, W. 81. (2) 531.)

Chromic zinc sulphide, Cr_2ZnS_4 .

(Gröger, W. A. B. 1880, 81. 534.)

Chromicyanhydric acid.**Cadmium chromicyanide, $\text{Cd}_2[\text{Cr}(\text{CN})_6]$**

Readily sol. in an excess of KCN a $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by conc. HNO_3 or H_2SO_4 . Slowly decomp. by rapidly by hot dil. HCl , HNO_3 , or F . Quickly dissolved by aqua regia. De by boiling with Na_2O_2 , by $\text{NaOH} + \text{A}$ by $\text{Na}_2\text{CO}_3 + \text{Aq}$. Slowly decomp. by l acetic acid. (Cruser and Miller, J. Chem. Soc. 1906, 28. 1136.)

Cobaltous chromicyanide, $\text{Co}_2[\text{Cr}(\text{CN})_6]$

Sol. in cold, readily sol. in hot conc or H_2SO_4 . Sl. sol. even in boiling

Slowly sol. in cold dil. H_2SO_4 , HCl or HNO_3 . Readily sol. in boiling dil. HCl or HNO_3 . Decomp. but not entirely dissolved in aqua regia. Readily sol. in an excess of HCl . Decomp. by NH_4OH , NaOH or KOH . Insol. in cold or boiling acetic acid. and Miller.)

chromicyanide, $\text{Cu}_2\text{Cr}(\text{CN})_6$.

Insol. in KCN , cold conc. or boiling dil. HCl . Slowly sol. in cold conc. H_2SO_4 , but rapidly sol. in hot dil. and conc. H_2SO_4 . Readily sol. by aqua regia. Decomp. by dil. HCl , slowly going into solution in H_2O , but quickly on boiling. (Cruser and Miller.)

chromicyanide, $\text{Cu}_2[\text{Cr}(\text{CN})_6]$.

Insol. in cold dil. HCl , HNO_3 or H_2SO_4 . The first two readily dissolve it, H_2SO_4 dissolves it only slowly. Sol. in aqua regia or cold conc. H_2SO_4 . Readily sol. in cold or hot conc. HCl . Decomp. by HNO_3 or by boiling HNO_3 . Decomp. by $\text{Na}_2\text{O}_2 + \text{Aq.}$ or NH_4OH , or $\text{Na}_2\text{CO}_3 + \text{Aq.}$ Readily sol. in excess of $\text{KCN} + \text{Aq.}$ Insol. in cold acetic acid. (Cruser and Miller.)

chromicyanide, $\text{Ni}_2[\text{Cr}(\text{CN})_6]$.

Insol. in cold, readily sol. in hot dil. HCl , HNO_3 or H_2SO_4 . Slowly sol. in cold, rapidly sol. in hot conc. H_2SO_4 , HCl or HNO_3 . Slowly decomp. by cold, rapidly by boiling aqua regia. Readily sol. in excess of HCl . Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Decomp. by $\text{Na}_2\text{CO}_3 + \text{Aq.}$ or $\text{Na}_2\text{O}_2 + \text{Aq.}$ Insol. in cold or boiling acetic acid. (Cruser and Miller.)

ammonium thallous chromicyanide, $\text{TlCr}(\text{CN})_6$.

Prepared by Fischer and Benzian, Ch. Z. 1902, 26. 50.)

chromicyanide, $\text{Tl}_2\text{Cr}(\text{CN})_6$.

Insol. in H_2O . (Fischer and Benzian, 1902, 26. 50.)

chromicyanide, $\text{Zn}_2[\text{Cr}(\text{CN})_6]$.

Insol. in H_2O . Sol. in excess of NH_4OH , and $\text{KCN} + \text{Aq.}$ Decomp. by Na_2CO_3 . Sol. in cold dil. HCl . Slowly sol. in H_2SO_4 and in dil. HNO_3 . By boiling in dil. acids a clear solution is quickly obtained. (Cruser, Dissert. 1906.)

chromisulphocyanhydric acid.

lithium chromisulphocyanide, $\text{Li}_2\text{Cr}(\text{SCN})_6 + 2\text{H}_2\text{O}$.

Insol. in H_2O than K salt. (Osann, 1907.)

Lithium chromisulphocyanide, $\text{Li}_2\text{Cr}(\text{SCN})_6 + \text{H}_2\text{O}$.

Extremely deliquescent. (Osann.)

Rubidium chromisulphocyanide, $\text{Rb}_2\text{Cr}(\text{SCN})_6 + 4\text{H}_2\text{O}$.

Appreciably less sol. in H_2O and alcohol than the K salt. (Osann.)

Chromocyandric acid, $\text{H}_4\text{Cr}(\text{CN})_6$.

Decomp. rapidly on air. Sol. in H_2O (Moissan, A. ch. (6) 4. 144.)

Potassium chromocyanide, $\text{K}_4\text{Cr}(\text{CN})_6$.

Very sol. in H_2O ; 100 pts. H_2O dissolve 32.33 pts. at 20° . Much more sol. in hot H_2O . Insol. in alcohol, ether, benzene, or chloroform. (Moissan, A. ch. (6) 4. 136.)

Above salt was $\text{K}_2\text{Cr}(\text{CN})_6$. (Christensen.) $+3\text{H}_2\text{O}$. (Christensen, J. pr. (2) 31. 166.)

Chromiodic acid, $\text{CrO}_3, \text{HIO}_3 + 2\text{H}_2\text{O}$.

Deliquescent. (Berg, C. R. 104. 1514.)

Ammonium chromiodate, $\text{CrO}_3, \text{NH}_4\text{IO}_3 + \text{H}_2\text{O}$.

Moderately sol. in H_2O . (Berg.)

Lithium chromiodate, $\text{CrO}_3, \text{LiIO}_3 + \text{H}_2\text{O}$.

Very sol. in H_2O . (Berg.)

Magnesium chromiodate.

Sol. in H_2O . (Berg.)

Potassium chromiodate, $\text{CrO}_3, \text{KIO}_3$.

Sol. in H_2O . (Berg.)

$+ \text{H}_2\text{O} = \text{KCrIH}_2\text{O}_7$. Sl. decomp. by H_2O . (Blomstrand, J. pr. (2) 40. 331.)

Silver chromiodate, $\text{CrO}_3, \text{AgIO}_3$.

Sl. attacked by cold, rapidly decomp. by hot H_2O . (Berg, C. R. 111. 42.)

Sodium chromiodate, $\text{CrO}_3, \text{NaIO}_3 + \text{H}_2\text{O}$.

Very sol. in H_2O . (Berg.)

Chromosulphocyanhydric acid.

Sodium chromosulphocyanide,

$\text{Na}_2\text{Cr}(\text{SCN})_6 + 10, \text{ or } 11\text{H}_2\text{O}$.

Unstable.

Decomp. by H_2O . (Koppel, Z. anorg. 1905, 45. 360.)

Chromosulphuric acid, $\text{H}_2\text{Cr}_2(\text{SO}_4)_4$.

Sol. in H_2O in all proportions, but solution is easily decomp. on standing or boiling. (Recoura, Bull. Soc. (3) 9. 586.)

$\text{H}_4\text{Cr}_2(\text{SO}_4)_6$. As above.

$\text{H}_6\text{Cr}_2(\text{SO}_4)_8$. As above.

Ammonium chromosulphate,
 $(\text{NH}_4)_2\text{Cr}_2(\text{SO}_4)_4 + 5\text{H}_2\text{O}$.

Sol. in H_2O after a few minutes. (Recoura.)

Chromium potassium chromosulphate,
 $[\text{Cr}_2(\text{SO}_4)_3\text{CrO}_4]\text{K}_2, [\text{Cr}_2(\text{SO}_4)_3(\text{CrO}_4)_2]\text{K}_4$,
 and $[\text{Cr}_2(\text{SO}_4)_3(\text{CrO}_4)]\text{K}_4$.

Sol. in H_2O . (Recoura, Bull. Soc. 1897, (3) 17. 934.)

Potassium chromosulphate, $\text{K}_2\text{Cr}_2(\text{SO}_4)_4 + 4\text{H}_2\text{O}$.

Sol. in H_2O in a few minutes. (Recoura, Bull. Soc. (3) 9. 590.)

Sodium chromosulphate, $\text{Na}_2\text{Cr}_2(\text{SO}_4)_4 + 10\text{H}_2\text{O}$.

As K salt. (Recoura.)

Chromotelluric acid.

Ammonium chromotellurate,
 $2(\text{NH}_4)_2\text{O}, 4\text{CrO}_3, \text{TeO}_3$.

Sol. in H_2O . (Berg, C. R. 1911, 152. 1588.)

Potassium chromotellurate,
 $2\text{K}_2\text{O}, 4\text{CrO}_3, \text{TeO}_3$.

Sl. sol. in cold H_2O without decomp.

Sol. in boiling H_2O . (Berg, Bull. Soc. 1911, (4) 9. 583.)

Chromous acid, $\text{H}_2\text{Cr}_2\text{O}_4 = \text{Cr}_2\text{O}_3, \text{H}_2\text{O}$.

Chromic hydroxide shows slightly acid properties, and salts corresponding to the above acid are known.

Aluminum ferrous magnesium chromite
 (chrome iron ore), $(\text{Fe}, \text{Mg})\text{O}$,
 $(\text{Cr}_2, \text{Al}_2)\text{O}_3$.

Insol. in H_2O or acids, even a mixture of H_2SO_4 and HF . (Ebelmen.)

Barium chromite, BaCr_2O_4 .

Insol. in H_2O . (Gerber, Bull. Soc. (2) 27. 436.)

Barium tetrachromite, $\text{BaO}, 4\text{Cr}_2\text{O}_3$.

Undecomp. by steam at red heat; insol. in HCl , H_2SO_4 , HNO_3 ; sol. in fused $\text{KOH} + \text{KNO}_3$; decomp. in the air. (Dufau, C. R. 1896, 122. 1126.)

Cadmium chromite, CdCr_2O_4 .

Not attacked by acids. (Viard, C. R. 109. 142.)

Calcium chromite, CaCr_2O_4 .

Insol. in H_2O . (Gerber, Bull. Soc. (2) 27. 436.)

Insol. in HCl , HF , HNO_3 , H_2SO_4 ; sol. in gaseous HCl and HF at red heat; sol. in fused KOH , KNO_3 , KClO_3 , K_2CO_3 . (Dufau, C. R. 1895, 121. 690.)

$2\text{CaO}, \text{Cr}_2\text{O}_3$. Insol. in H_2O , KOH , $\text{NH}_4\text{OH} + \text{Aq}$; slowly decomp. by H_2O or $\text{M}_2\text{CO}_3 + \text{Aq}$; insol. in sugar solut (Pelouse, A. ch. (3) 33. 9.)

$4\text{CaO}, \text{Cr}_2\text{O}_3$. Attacked by H_2O . (J. san, C. R. 1894, 119. 188.)

Cobaltous chromite, CoCr_2O_4 .

(Elliot, Dissert, Göttingen, 1862.)

Cuprous chromite, $\text{Cu}_2\text{O}, \text{Cr}_2\text{O}_3$.

Insol. in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.4). (W. Z. phys. Ch. 1908, 62. 445.)

Cupric chromite basic, $5\text{CuO}, 4$
 (Wöhler, Z. phys. Ch. 1908, 62. 445.)

Cupric chromite, CuCr_2O_4 .

Not attacked by $\text{HNO}_3 + \text{Aq}$. (Per. ch. (3) 25. 283.)

Not attacked by conc. HCl .
 Insol. in dil. acids. (Wöhler, Z. Ch. 1908, 62. 446.)

$\text{CuO}, 3\text{Cr}_2\text{O}_3$. (Rosenfeld, B. 187 958.)

Glucinum chromite, GlCr_2O_4 .

Insol. in H_2O . (Mallard, C. R. 105.)

Iron (ferrous) chromite (chrome iron ore)

See Chromite, aluminum ferrous chromite.

Iron (ferroferic) chromite, $\text{FeO}, \text{Fe}_2\text{O}_3$.

Not attacked by $\text{HCl} + \text{Aq}$. (Ebelmen.)

Iron (ferrous) magnesium chromite.

Insol. in $\text{HCl} + \text{Aq}$. Scarcely attacked by H_2SO_4 .

Lead chromite, PbCr_2O_4 .

Ppt. Insol. in $\text{KOH} + \text{Aq}$. (C. R. 43. 927.)

Lithium chromite, $\text{Li}_2\text{Cr}_2\text{O}_4$.

Very sl. sol. in acids. (Weyberg, 1906, II. 1659.)

Magnesium chromite, $\text{MgO}, 2\text{Cr}_2\text{O}_3$.

Insol. in H_2O . (Nichols, Sill. Am. 47. 16.)

MgCr_2O_4 . Insol. in acids or alkalis except boiling H_2SO_4 . (Schweitzer, J. 1 259.)

Could not be obtained. (Viard, Bull. (3) 5. 934.)

Easily attacked by boiling H_2SO_4 .
 Less easily by HCl or $\text{HF} + \text{Aq}$ not attacked by boiling HNO_3 . (Dufau, 1896, 123. 886.)

$2\text{MgO}, \text{Cr}_2\text{O}_3$. Insol. in H_2O or (Nichols.)

$5\text{MgO}, 4\text{Cr}_2\text{O}_3$. Insol. in acids. (C. R. 112. 1003.)

$3\text{MgO}, 2\text{Cr}_2\text{O}_3$. As above. (V.)

chromite, MnCr_2O_4 .
insol. in acids. (Ebelmen, A. 44.)

ite, ZnCr_2O_4 .
acids and alkalies. (Viard, C. R.

(Chancel, C. R. 43. 927.)
 Cr_2O_3 . As above. (Viard, C. R.

Cr_2O_3 . As above. (V.)
 Cr_2O_3 . (Gröger, M. 1904, 25,

nic acid.

chromovanadate, $2(\text{NH}_4)_2\text{O}$,
 $\text{V}_2\text{O}_5 + 7\text{H}_2\text{O}$.
 H_2O . (Ditte, C. R. 102. 1105.)

amide, $\text{CrO}_2(\text{NH}_3)_2$.
 H_2O . (Ohly, C. N. 1899, 80. 134.)

trichloride, $(\text{CrO}_2)_3\text{Cl}_4$.
cent, sol. in H_2O with decomp.,
dry ether. (Pascal, C. R. 1909,

chloride (chlorochromic acid).
 H_2 .
by H_2O with evolution of much
in glacial acetic acid without de-
n.
 Cl_4 , C_6H_6 , (mol. wt. det.). (Oddo,
it. 1899, 29. (2) 318; Chem. Soc.
(2) 75.)

trichloride, $\text{Cr}_2\text{O}_3\text{Cl}_3$.
cent. Sol. in H_2O with gradual
tion. Sol. in conc. $\text{HCl} + \text{Aq}$.
Chem. Soc. (2) 8. 31.)
sol. in CS_2 .
alcohol and ether. (Rawson, C. N.
185.)

chlorides.

H_2O_2 .
chromium oxychlorides.

chloride nitrogen tetroxide,
 H_2O_7 , 2NO_2 .
 H_2O with decomp. (Thomas, C. R.
828.)

fluoride, CrO_2F_2 .
by H_2O with evolution of heat.
ass. ch. it. 16. 218.)

silicate, aluminum, Al_2O_3 , $\text{SiO}_2 +$

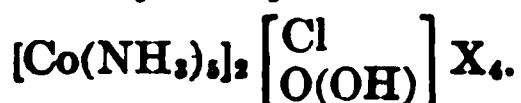
Cobalt, Co.

Not attacked by H_2O .
Sol. in dil. HCl , or H_2SO_4 , or $\text{HNO}_3 + \text{Aq}$.
Conc. hot H_2SO_4 , and HNO_3 decomp. with
evolution of SO_2 or NO gas.
Exists also in passive state. See Iron.
(Nickles, J. pr. 61. 186.)
Sol. in conc. $\text{KOH} + \text{Aq}$ when in finely
divided state. (Winkler, J. pr. 91. 211.)
Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ in presence of air.
(Hodgkinson and Bellairs, C. N. 1895, 71. 73.)

Cobalt ammonia compounds.

See—

Anhydrooxycobaltamine compounds,



Bromotetramine cobaltic compounds,
 $\text{BrCo}(\text{NH}_3)_4\text{X}_2$.

Bromopurpureocobaltic compounds,
 $\text{BrCo}(\text{NH}_3)_4\text{X}_2$.

Carbonatotetramine cobaltic compounds,
 $(\text{CO}_2)\text{Co}(\text{NH}_3)_4\text{X}$.

Chlorotetramine cobaltic compounds,
 $\text{ClCo}(\text{NH}_3)_4\text{X}_2$.

Chloropurpureocobaltic compounds,
 $\text{ClCo}(\text{NH}_3)_4\text{X}_2$.

Croceocobaltic compounds,
 $\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{X}$.

Decamine cobaltic sulphite,
 $\text{Co}_2(\text{NH}_3)_{10}(\text{SO}_3)_2$.

Diamine cobaltic nitrites,
 $\text{Co}(\text{NH}_3)_2(\text{NO}_2)_4\text{M}$.

Dichrocobaltic compounds, $\text{Co}(\text{NH}_3)_4\text{X}_2$.

Flavocobaltic compounds,
 $(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4\text{X}$.

Fuscocobaltic compounds,
 $(\text{OH})\text{Co}(\text{NH}_3)_4\text{X}_2$.

Iodotetramine cobaltic compounds,
 $\text{ICO}(\text{NH}_3)_4\text{X}_2$.

Luteocobaltic compounds, $\text{Co}(\text{NH}_3)_4\text{X}_2$.

Melanocobaltic compounds,
 $[\text{Co}(\text{NH}_3)_2\text{Cl}_2]_2, \text{NH}_4\text{Cl}$.

Nitratotetramine cobaltic compounds,
 $(\text{NO}_2)\text{Co}(\text{NH}_3)_4\text{X}_2$.

Nitratopurpureocobaltic compounds,
 $(\text{NO})_2\text{Co}(\text{NH}_3)_4\text{X}_2$.

Nitritocobaltic compounds,
 $(\text{NO}_2)\text{Co}(\text{NH}_3)_4\text{X}_2$.

Octamine cobaltic compounds,
 $\text{Co}_2(\text{NH}_3)_8\text{X}_6$.

(= Tetramine cobaltic compounds,
 $\text{Co}(\text{NH}_3)_4\text{X}_2$).

Oxycobaltamine compounds,
 $\text{Co}_2(\text{NH}_3)_{10}(\text{OOH})\text{X}_4$.

Praseocobaltic compounds, $\text{Co}(\text{NH}_3)_4\text{X}_2$.

Purpureocobaltic compounds,
 $\text{Co}(\text{NH}_3)_4\text{X}_2$.

Roseocobaltic compounds,
 $\text{Co}(\text{NH}_3)_4(\text{OH})_2\text{X}_2$.

Sulphatotetramine cobaltic compounds,
 $(\text{SO}_4)\text{Co}(\text{NH}_3)_4\text{X}$.

Sulphatopurpureocobaltic compounds,
 $(\text{SO}_4)\text{Co}(\text{NH}_3)_4\text{X}$.

"Tetramine cobaltic" compounds,
 $\text{Co}(\text{NH}_3)_4\text{X}_2$.

Xanthocobaltic compounds,
 $(\text{NO}_2)\text{Co}(\text{NH}_3)_4\text{X}_2$.

Cobalt arsenide, CoAs_2 .

As Co_2As_2 . (Ducelliez, C. R. 1908, 147. 425.)

CoAs_2 . As Co_2As_2 . (Ducelliez, C. R. 1908, 147. 425.)

Co_2As_2 . As Co_2As_2 . (Ducelliez, C. R. 1908, 147. 425.)

Co_2As_2 . Very sl. attacked by hot conc. HCl , less by H_2SO_4 . Easily sol. in HNO_3 and aqua regia. Sl. attacked by fused alkalis and alkali carbonates. (Ducelliez, C. R. 1908, 147. 425.)

CoAs_2 . Min. *Skutterudite*. Sol. in HNO_3 + Aq, with separation of As_2O_3 .

Cobalt arsenide sulphide, CoAs_2S_2 .

Min. *Cobaltite*. Sol. in HNO_3 + Aq, with separation of S and As_2O_3 .

Glaucodote. Completely sol. in HNO_3 + Aq.

Cobalt azoimide, basic, $\text{Co}(\text{OH})\text{N}_3$.

Insol. in H_2O .

Sol. in HN_3 + Aq. (Curtius, J. pr. 1898, (2) 58. 300.)

Cobalt potassium azoimide, KN_3 , $\text{Co}(\text{N}_3)_2$.

Sol. in H_2O ; Aq. solution decomp. on boiling. (Curtius, J. pr. 1898, (2) 58. 301.)

Cobalt boride, Co_2B .

Attacked by HNO_3 . (Jassonneix, C. R. 1907, 145. 240.)

CoB . Decomp. by moist air and by alkali nitrates, chlorates, hydroxides and carbonates; decomp. by steam at red heat and by acids. (Moissan, C. R. 1896, 122. 425.)

Not attacked by HCl ; rapidly attacked by HNO_3 . Not attacked by dil. but decomp. by conc. H_2SO_4 . Rapidly attacked by aqua regia. (Moissan, A. ch. 1896, (7) 9. 272.)

CoB_2 . (Jassonneix, C. R. 1907, 145. 241.)

Cobaltous bromide, CoBr_2 .

Deliquescent. Sol. in H_2O , alcohol, and ether.

Sat. CoBr_2 + Aq contains at:

59° 75° 97°
 86.7 66.8 68.1% CoBr_2 .

(Étard, A. ch. 1894, (7) 2. 542.)

Nearly insol. in AsBr_3 . (Walden, Z. anorg. 1902, 29. 374.)

Sol. in $\text{SO}_2\text{Cl}(\text{OH})$. (Walden.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

lg. CoBr_2 is sol. in 9.74g. methyl acetate at 18°. Sp. gr. 18°/4° of sat. solution = 1.013. (Naumann, B. 1909, 42. 3792.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Sol. in acetone. (Eidmann, C. C. 1893, II. 1014; Naumann, B. 1904, 37. 4328.)

Mol. weight determined in pyridine. (Werner, Z. anorg. 1897, 15. 24.)

+2, and $6\text{H}_2\text{O}$. (Hartley, Chem. Soc. (2) 12. 214.)

Cobaltous hydrazine bromide hydrazine, $2\text{CoBr}_2, 4\text{N}_2\text{H}_4\text{HBr}, \text{N}_2\text{H}_4(?)$.

Sol. in H_2O with decomp. (Ferratini, C. A. 1912. 1613.)

Cobaltous mercuric bromide, basic, $\text{CoBr}_2, \text{HgBr}_2, 6\text{CoO} + 20\text{H}_2\text{O}$.

(Mailhe, A. ch. 1902, (7) 27. 369.)

Cobalt stannic bromide.

See Bromostannate, cobalt.

Cobaltous bromide ammonia, $\text{CoBr}_2, 6\text{NH}_3$.

Sol. in H_2O with residue of cobalt hydrazide. (Rammelsberg, Pogg. 55. 245.)

Cobaltous bromide hydrazine, $\text{CoBr}_2, 2\text{N}_2\text{H}_4$.

Decomp. by boiling with H_2O . (Fransen, Z. anorg. 1908, 60. 270.)

Cobalt carbonyl, $\text{Co}(\text{CO})_4$.

Insol. in H_2O . More or less sol. in CS_2 , ether, alcohol and $\text{Ni}(\text{CO})_4$. Relatively stable with non-oxidizing acids. Quickly decomp. by oxidizing acids. (Mond, Hirtz and Cowap, C. N. 1908, 98. 165.)

Cobaltous chloride, CoCl_2 .

Deliquescent. Sol. in H_2O with evolution of heat. 100 pts. H_2O dissolve 43.3 pts. CoCl_2 at 0°. (Engel, A. ch. (6) 17. 355.)

100 pts. sat. CoCl_2 + Aq at t° contain pts. CoCl_2 .

t°	Pts. CoCl_2	t°	Pts. CoCl_2	t°	Pts. CoCl_2
-22	24.7	25	34.4	56	48.4
-4	28.0	34	37.5	78	48.8
+7	31.2	41	39.8	94	50.5
11	31.3	45	41.7	96	51.2
12	32.5	49	46.7	112	52.3

(Étard, C. R. 113. 699.)

Sp. gr. of CoCl_2 + Aq containing—

5 10 12 20 25% CoCl_2
 1.0496 1.0997 1.1579 1.2245 1.3002

Sat. solution, 1.3613.

(Franz, J. pr. (2) 5. 284.)

Sp. gr. of CoCl_2 + Aq containing in 1000 g. H_2O , g. $\text{CoCl}_2 + 6\text{H}_2\text{O}$ —

119 g. (= $\frac{1}{2}$ mol.) 238 357 476 594
 1.055 1.101 1.141 1.177 1.200

833 952 1071 1190
 1.238 1.264 1.287 1.309

Containing g. CoCl_2 (anhydrous)—
 65 g. (= $\frac{1}{2}$ mol.) 130 195 260 325 390
 1.058 1.112 1.164 1.213 1.260 1.304
 (Gerlach, Z. anal. 28. 466.)

Sp. gr. of $\text{CoCl}_2 + \text{Aq}$ at room temp. containing:

7.97 14.858 22.27% CoCl_2 .
 1.0807 1.1613 1.2645

(Wagner, W. Ann. 1883, 18. 267.)

Sp. gr. of $\text{CoCl}_2 + \text{Aq}$ at 20° containing M g mols. of salt per liter.

M 0.01 0.025 0.05 0.075
 Sp. gr. 1.001159 1.003052 1.006065 1.009190

M 0.10 0.25 0.50 0.75
 Sp. gr. 1.012386 1.03049 1.05492 1.09118

M 1.0 1.5 2.0
 Sp. gr. 1.11847 1.17502 1.23637
 (Jones and Pearce, Am. Ch. J. 1907, 38. 711.)

Sp. gr. of $\text{CoCl}_2 + \text{Aq}$ at 25°.

Concentration of $\text{CoCl}_2 + \text{Aq}$.	Sp. gr.
1—normal	1.0571
$\frac{1}{2}$ —	1.0286
$\frac{1}{4}$ —	1.0144
$\frac{1}{8}$ —	1.0058

(Wagner, Z. phys. Ch. 1890, 5. 37.)

Solubility in $\text{HCl} + \text{Aq}$ at 0°. $\frac{\text{CoCl}_2}{2} = \frac{1}{2}$ mols.

CoCl_2 in mgs. in 10 ccm. of solution.
 $\text{HCl} = \text{mols. HCl in mgs. in ditto. H}_2\text{O}$
 $= \text{g. H}_2\text{O.}$

$\frac{\text{CoCl}_2}{2}$	HCl	$\text{CoCl}_2 + \text{HCl}$	Sp. gr.	H_2O
62.4	0	62.4	1.343	9.36
58.525	3.7	62.2	1.328	9.34
50.8	11.45	62.25	1.299	9.27
37.25	25.2	62.45	1.248	9.13
12.85	55.0	67.85	1.167	...
4.75	74.75	79.50	1.150	8.46
12.0	104.5	116.5	1.229	7.5
25.0	139.0	164.0	1.323	...

(Engel, A. ch. (6) 17. 355.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Sol. in alcohol.

Sat. solution in alcohol (0.792 sp. gr.) contains 23.66 % CoCl_2 and has sp. gr. = 1.0107. (Winkler, J. pr. 91. 209.)

Very sol. in ether.

100 pts. absolute ether dissolve only 0.021 g. CoCl_2 . (Böttker, Z. phys. Ch. 1897, 22. 511.)

100 g. formic acid (95%) dissolve 6.2 g. CoCl_2 at 20.2°. (Aschan, Ch. Z. 1913, 37. 1117.)

1g. CoCl_2 is sol. in 271g. methyl acetate at 18°. Sp. gr. 18°/4° of sat. solution = 0.938. (Naumann, B. 1909, 42. 3791.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

100 pts. acetone dissolve 8.62 pts. anhydrous CoCl_2 . (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

0.08 pts. sol. in 100 pts. ethyl acetate at 14°.
 0.26 " " " 100 " " " 79°.
 9.11 " " " 100 " acetone " 0°.
 9.28 " " " 100 " " " 22.5°.

(Laszczynski, B. 1894, 27, 2286.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

1 g. CoCl_2 is sol. in 36.4 g. acetone at 18°. Sp. gr. of sat. solution 18°/4° = 0.825. (Naumann, B. 1904, 37. 4334.)

100 g. acetonitrile dissolve 4.08 g. CoCl_2 at 18°. (Naumann and Schier, B. 1914, 97. 249.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

Solubility in pyridine at t°.

t°	G. CoCl_2 sol. in 100 g. pyridine	Solid phase
—50.3	0.4200	$\text{CoCl}_2, 6\text{C}_5\text{H}_5\text{N}$
—45.0	0.4204	
—30.0	0.4224	
—19.6	0.4227	
—10.0	0.4329	
0	0.4326	
+23.0	0.572	$\text{CoCl}_2, 4\text{C}_5\text{H}_5\text{N}$
25.0	0.578	
34.6	0.755	
37.6	0.760	
44.6	0.959	
47.2	1.029	
51.0	1.122	$\text{CoCl}_2, 2\text{C}_5\text{H}_5\text{N}$
55.0	1.206	
60.0	1.342	
64.2	1.483	
68.0	1.597	
74.8	2.079	
78.2	2.330	CoCl_2
79.8	2.488	
88.0	3.397	
96.5	7.817	
98.8	8.862	
106.0	14.340	
110.0	16.500	

(Pearce and Moore, Am. Ch. J. 1913, 50. 226.)

Mol. weight determined in piperidine, and pyridine. (Werner Z. anorg. 1897, 15. 18 and 23.)

Sol. in urethane. (Castoro, Z. anorg. 1899, 20. 61.)

+ H_2O .

+2H₂O. Very deliquescent. (Bersch, J. B. 1867, 291.)
17.16 pts. sol. in 100 pts. acetone at 0°.
17.06 " " 100 " " 25°.

(Laszczynski, B. 1894, 27. 2287.)

+4H₂O. Deliquescent. (Bersch.)

+6H₂O. Not deliquescent. Easily sol. in H₂O.

Solubility of CoCl₂·6H₂O in ethyl alcohol + Aq at 11.5° under addition of increasing amounts of CoCl₂.

P = Percent of alcohol by volume.

G = Grams of CoCl₂ added.

C_c = Grams of CoCl₂ in 5 cc. of the solution.

C_w = Grams of water in 5 cc. of the solution, calculated from

(1) the water content of the alcohol.

(2) the water of crystallization which had gone into solution.

(3) the water held mechanically in CoCl₂·6H₂O.

P	G	C _w	C _c
91.3	0.0	1.325	1.168
98.3	0.0	1.134	1.214
98.3	0.0	1.068	1.181
99.3	0.0	1.045	1.199
"	0.194	0.899	1.204
"	0.400	0.829	1.325
"	0.612	0.764	1.459
"	0.813	0.688	1.568
"	1.022	0.634	1.713
"	1.240	0.553	1.831
"	1.446	0.483	1.943
"	0.650	0.500	2.186

(Bödtker, Z. phys. Ch. 1897, 22. 508.)

Easily soluble in absolute ethyl alcohol. 100 pts. absolute alcohol dissolve at room temperature 56.20 pts. CoCl₂. Water precipitates CoCl₂·6H₂O from a solution of CoCl₂ in absolute alcohol. (Bödtker.)

100 pts. absolute ether dissolve 0.291g. CoCl₂·6H₂O. (Bödtker, Z. phys. Ch. 1897, 22. 511.)

Anhydrous ethylene glycol dissolves 10.6% CoCl₂·6H₂O at 16.4°. (de Coninck, Chem. Soc. 1904, 86, (2) 741.)

Cobaltous hydrazine chloride,
CoCl₂·2N₂H₄·HCl + 2½H₂O.

Sol. in H₂O. (Ferratini, C. A. 1912. 1613.)

Cobaltous iodine chloride, CoCl₂·2ICl₃ + 8H₂O.

Hygroscopic. Decomp. by H₂O. CCl₄ separates ICl₃. (Weinland and Schlegelmilch, Z. anorg. 1902, 30. 137.)

Cobalt lithium chloride, CoCl₂·LiCl + 3H₂O.

Very deliquescent. Sol. in H₂O with decomp. Sol. in LiCl + Aq without decomp. Sol. in alcohol without decomp. (Chassevant, A. ch. (6) 30. 27.)

Cobaltous mercuric chloride basic, CoCl₂·HgCl₂·6CoO + 20H₂O.

(Mailhe, A. ch. 1902, (7) 27. 369.)

Cobaltous mercuric chloride, CoCl₂·HgCl₂.
Very deliquescent. (v. Bonsdorff.)

Cobaltous thallic chloride, 2TlCl₃·CoCl₂ + 8H₂O.

Hygroscopic; can be cryst. from H₂O. (Gewecke, A. 1909, 366. 222.)

Cobaltous tin (stannic) chloride, CoCl₂·SnCl₄ + 6H₂O.

See Chlorostannate, cobaltous.

Cobaltous chloride ammonia, CoCl₂·2NH₃.

Decomp. by H₂O. (F. Rose.)

CoCl₂·4NH₃. Decomp. by H₂O. (H. Rose.)

CoCl₂·6NH₃. Decomp. by H₂O. Sol. in dil. NH₄OH + Aq with ease, but difficultly in conc. NH₄OH + Aq. Insol. in absolute alcohol. (Fremy.)

Cobaltous chloride hydrazine, CoCl₂·2N₂H₄.

Insol. in cold H₂O.

Slowly decomp. by cold, rapidly by hot H₂O.

Easily sol. in dil. acids and NH₄OH + Aq. (Franzen, Z. anorg. 1908, 60. 270.)

Cobaltous chloride hydroxylamine,
CoCl₂·2NH₂OH.

Decomp. in the air; sol. in H₂O. (Feldt, B. 1894, 27. 403.)

Cobaltic chloride hydroxylamine,
CoCl₃·6NH₂OH.

Insol. in alcohol.

Sol. in acidified H₂O without decomp.; sol. in conc. H₂SO₄ without decomp. (Feldt, B. 1894, 27. 404.)

Cobaltous fluoride, CoF₂.

Sl. sol. in H₂O; insol. in alcohol and ether; slowly attacked by cold HCl, H₂SO₄, or HNO₃ + Aq. (Poulenc, C. R. 114. 1429.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 29. 827.)

+2H₂O. Sol. in a little H₂O without decomp. Decomp. into oxyfluoride by boiling with much H₂O. Sol. in HF + Aq. (Berzelius.)

+4H₂O. Two modifications.

Solubility of α mod. at 7° = 2.2328 %
" " β " " " = 2.3203 %.

(Costachescu, Ann. Sci. Univ. Jassy, 1912, 7, 1, 10.)

Cobaltic fluoride, CoF₃.

Sol. in conc. H₂SO₄. (Barbieri, Chem. Soc. 1905, 88, (2) 393.)

as hydrogen fluoride, CoF_2 , $5\text{HF} + \text{H}_2\text{O}$.

r sol. in H_2O and dil. acids.

1 $\text{NH}_4\text{OH} + \text{Aq}$ with decomp. (Böhm, Z. 1905, 43. 330.)

columbium fluoride.

luocolumbate, cobalt.

us iron (ferric) fluoride,

F_2 , $\text{FeF}_3 + 7\text{H}_2\text{O}$.

1 dil. $\text{HF} + \text{Aq}$. (Weinland, Z. anorg. 2. 269.)

us manganic fluoride, 2CoF_2 , Mn_2F_8 , $10\text{H}_2\text{O}$.

stensen, J. pr. (2) 34. 41.)

molybdenyl fluoride.

fluoxymolybdate, cobalt.

us potassium fluoride, CoF_2 , KF .

ol. in H_2O ; less in ethyl or methyl; insol. in amyl alcohol or benzene. p. by hot H_2SO_4 . (Poulenc, C. R. 114.

O. Sl. sol. in H_2O . (Wagner, B. 19.

, 2KF .

us sodium fluoride, CoF_2 , $\text{NaF} + \text{H}_2\text{O}$. in H_2O . (Wagner, B. 19. 896.)

us stannic fluoride.

fluostannate, cobaltous.

vanadium fluoride.

fluovanadate, cobalt.

us hydroxide, CoO_2H_2 .

l. in H_2O . Sol. in acids. Insol. in -Aq. Sol. in ammonium sulphate, e, nitrate, or succinate + Aq. (Brett.) in warm acetic acid; insol. in NH_4OH and cold $\text{NH}_4\text{Cl} + \text{Aq}$, but sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (de Schulten, C. R. 109.

l. in H_2O and dil. $\text{KOH} + \text{Aq}$; some-ol. in conc. $\text{KOH} + \text{Aq}$; easily sol. in salts + Aq. (Fresenius.)

ly sol. in $\text{KCN} + \text{Aq}$. (Rodgers, 1834.) in conc. $\text{K}_2\text{CO}_3 + \text{Aq}$. (Gmelin.)

pptd. by $\text{KOH} + \text{Aq}$ in presence of O_2 or NH_4 citrate. (Spiller.)

in large amt. in boiling $\text{NH}_4\text{SCN} + \text{Aq}$. mann, Z. anorg. 1908, 58. 269.)

l. in methyl, or amyl amine + Aq. l.)

y non-volatile organic substances pre-s pptn.

ic hydroxide, $3\text{Co}_2\text{O}_3$, $2\text{H}_2\text{O}$.

la, Phil. Mag. (4) 35. 257.)

), $2\text{H}_2\text{O}$. Decomp. by $\text{HCl} + \text{Aq}$;

gives brown solutions with cold HNO_3 or $\text{H}_2\text{SO}_4 + \text{Aq}$, which soon decomp. (Wernicke, Pogg. 141. 120.)

$\text{Co}_2\text{O}_3\text{H}_2 = \text{Co}_2\text{O}_3 \cdot 3\text{H}_2\text{O}$. Sol. in warm HCl , HNO_3 , and H_2SO_4 , with decomp. (Proust.)

Sol. in cold H_3PO_4 , H_2SO_4 , HNO_3 , or $\text{HCl} + \text{Aq}$, but decomp. on standing or warming. (Winkelblech.)

Sol. in racemic, tartaric, oxalic, or citric acid as cobaltous salt.

Sol. in conc. acetic acid without immediate decomp. (Remele). Solution is not decomp. by boiling. Sol. in warm sat. $(\text{NH}_4)_2\text{C}_2\text{O}_4 + \text{Aq}$ with decomp.

Not attacked by cold or hot $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$.

Sol. when freshly pptd. in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Geuther, A. 128. 157.)

Cobaltocobaltic hydroxide, Co_2O_4 , $3\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in oxalic acid; solution decomp. by heat. Sol. in $\text{HCl} + \text{Aq}$ with evolution of Cl . (Gibbs and Genth, Sill. Am. J. (2) 23. 257.)

$\text{Co}_2\text{O}_4 \cdot 7\text{H}_2\text{O}$. Sol. in weak acids, especially $\text{HC}_2\text{H}_3\text{O}_2$ without decomp. (Fremy.)

Co_2O_7 , $6\text{H}_2\text{O}$. Min. *Heterogenite*. Sol. in dil. $\text{HCl} + \text{Aq}$ with evolution of Cl .

Cobaltous iodide, CoI_2 .

Deliquescent, and very sol. in H_2O .

100 pts. sat. $\text{CoI}_2 + \text{Aq}$ at t° contain
pts. CoI_2 .

t°	Pts. CoI_2	t°	Pts. CoI_2	t°	Pts. CoI_2
-22	52.4	14	61.6	60	79.2
- 8	56.7	25	66.4	82	80.7
- 2	58.7	34	73.0	111	80.9
+ 9	61.4	46	79.0	156	83.1

(Étard, C. R. 113. 699.)

Sol. in $\text{SO}_2(\text{OCH}_3)_2$. (Walden, Z. anorg. 1902, 29. 388.)

Sol. in SOCl_2 . (Walden, Z. anorg. 1900, 25. 216.)

Sol. in POCl_3 . (Walden, Z. anorg. 1900, 25. 212.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, 25. 217.)

Nearly insol. in AsBr_3 . (Walden, Z. anorg. 1902, 29. 374.)

Sol. in AsCl_3 . (Walden, Z. anorg. 1900, 25. 214.)

Easily sol. in alcohol.

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.) (Naumann, B. 1904, 37. 4328.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

Sol. in methyl acetate. (Naumann, B. 1909, 42, 3790.)

+ $2\text{H}_2\text{O}$.

+ $4\text{H}_2\text{O}$. Very deliquescent. (Étard.)

+6H₂O. (Hartley, Chem. Soc. (2) 12. 214.)

+9H₂O. Very hygroscopic. (Bolschakoff, C. C. 1898, II. 660.)

Cobaltous lead iodide, 2 CoI₂, PbI₂+3H₂O.

Decomp. by H₂O. (Mosnier, A. ch. 1897, (7) 12. 412.)

Cobaltous mercuric iodide, CoI₂, HgI₂+6H₂O.

Partially decomp. by H₂O.

Sol. in alcohol and acetone. (Dobroserdoff, C. C. 1901, II. 332.)

CoI₂, 2HgI₂+6H₂O. Decomp. by H₂O; sol. in alcohol and acetone. (Dobroserdoff, C. C. 1901, II. 332.)

Cobaltous iodide ammonia, CoI₂, 4NH₃.

Decomp. by H₂O. Sol. in NH₄OH+Aq. (Rammelsberg, Pogg. 48. 155.)

CoI₂, 6NH₃. Insol. in NH₄OH+Aq. (Rammelsberg.)

Cobaltous iodide hydrazine, CoI₂, 2N₂H₄.

Sl. sol. in H₂O. Easily sol. in acids. (Franzen, Z. anorg. 1911, 70. 147.)

Cobaltic octamine compounds.

See Octamine cobaltic compounds.

Cobaltous oxide, CoO.

Insol. in H₂O. Easily sol. in dil. or conc. HCl or HNO₃+Aq. Slowly sol. in cold, but easily in hot dil. H₂SO₄+Aq, acetic, or tartaric acid +Aq. Insol. in NH₄OH+Aq. Sol. in hot NH₄Cl +Aq, KOH, or NaOH+Aq. (Rose.)

Insol. in NH₄Cl or NH₄NO₃+Aq. (Brett, 1834.)

Insol. in K₂CO₃+Aq. Sol. in boiling Ce and Ni nitrates +Aq, with pptn. of the oxides. (Persoz.)

Easily sol. in dil. acids, even tartaric, acetic, and oxalic acids. Not attacked by NH₄OH+Aq. Sol. in 13% NH₄Cl+Aq with evolution of NH₃; also in NH₄SCN+Aq. Sol. in warm conc. NaOH, and KOH+Aq. (Zimmerman, A. 232. 324.)

Solubility in (calcium succate+sugar)+Aq. 1 l. solution containing 418.6 g. sugar and 34.3 g. CaO dissolves 1.56 g. CoO; 1 l. solution containing 296.5 g. sugar and 24.2 g. CaO dissolves 0.29 g. CoO. (Bodenbender, J. B. 1865. 600.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

See also Cobaltous hydroxide.

Cobaltic oxide, Co₂O₃.

Decomp. by most acids, even in the cold, with formation of cobaltous salts. Sol. in acetic acid without immediate decomp.

See also Cobaltic hydroxide.

Cobaltocobaltic oxide, Co₂O₄=CoO, Co₂O₃.

Insol. in boiling conc. HCl, HNO₃, or aqua regia. Sol. by long standing with H₂SO₄. (Gibbs and Genth, Sill. Am. J. (2) 23. 257.)

See also Cobaltocobaltic hydroxide.

Co₂O₄=2CoO, Co₂O₃.

Co₂O₇=4CoO, Co₂O₃. Not attacked by boiling dil. HNO₃ or H₂SO₄+Aq. (Beetz.)

Co₂O₄=6CoO, Co₂O₃+20H₂O. Sol. in dil. acids, with residue of Co₂O₃, which dissolves on warming. (Gentele, J. pr. 69. 131.)

+8H₂O. As above. (Gentele.)

Cobaltous oxychloride, CoCl₂, 3CoO+3½H₂O.

Ppt. Very sl. sol. in H₂O. (Habermann, M. 5. 432.)

Cobaltous oxychloride hydroxylamine, CoOCl, 2NH₂OH.

Insol. in H₂O; unstable; insol. in alcohol. (Feldt, B. 1894, 27. 404.)

Cobaltous oxyfluoride, CoO, CoF₂+H₂O.

Ppt. (Berzelius, Pogg. 1. 26.)

Cobaltous oxyiodide, CoO, CoI₂.

Insol. in H₂O. (Rammelsberg.)

Cobaltous oxysulphide, CoO, CoS.

Cold HCl+Aq dissolves out CoO; hot HCl+Aq decomp. with evolution of H₂S. (Arfvedson, Pogg. 1. 64.)

Cobalt phosphide, Co₂P.

Sol. in conc. HNO₃. Slowly attacked by HCl and H₂SO₄. (Maronneau, C. R. 1900, 130. 658.)

Sol. in HNO₃, aqua regia, and in fused alkalis. (Granger, Bull. Soc. 1896, (3) 15. 1089.)

Co₂P₃. Insol. in HNO₃ and aqua regia; stable in the air even when heated. (Granger, Bull. Soc. 1896, (3) 15. 1087.)

Co₂P₂. Insol. in conc. HCl+Aq. Sol. in HNO₃+Aq. (Rose, Pogg. 24. 332.)

Cobalt subselenide, Co₂Se.

Sol. in bromine water.

Only sl. attacked by boiling fuming HCl. (Fonze-Diacon, C. R. 1900, 131. 704.)

Cobalt monoselenide, CoSe.

(Little, A. 112. 211.)

Cobalt diselenide CoSe₂.

Sol. in Br₂+Aq.

Only sl. attacked by boiling fuming HCl. (Fonze-Diacon, C. R. 1900, 131. 705.)

arsenide, Co_2Se_3 .

$\text{Br}_2 + \text{Aq.}$

sl. attacked by boiling fuming HCl .
Diacon, C. R. 1900, 131. 704.)

elenide, Co_2Se_4 .

$\text{Br}_2 + \text{Aq.}$

sl. attacked by boiling fuming HCl .
Diacon, C. R. 190, 131. 704.)

silicide, Co_2Si .

sl. in HF and aqua regia. Insol. in cold
Decomp. by steam at red heat. Sol.
in dil. alkali carbonates. (Vigouroux,
1895, 121. 687.)

Insol. in HNO_3 and H_2SO_4 . Sol.
in aqua regia and HCl , and in fused KOH .
(C. R. 1901, 132. 557.)

Not attacked by dil. or conc. HNO_3 , or
 H_2SO_4 . Sol. in aqua regia and in conc.

Not attacked by dil. alkali hydroxides +
not attacked by fused alkali. (Lebau, Bull.
1911, (3), 25. 540.)

Sl. sol. in hot conc. HCl and hot
conc. $\text{KOH} + \text{Aq.}$ Sol. in HF ; insol. in
conc. H_2SO_4 . (Lebeau, C. R. 1902, 135.

disulphide, CoS .

Amorphous. Easily sol. in acids, even
in H_2O , but only slowly in the latter case.
(Lebau, C. R. 65. 75.)

Not attacked by cold dil. $\text{HCl} + \text{Aq.}$ (Ebel-
sch, (3) 25. 94.)

Seymourite.

1 l. H_2O dissolve $41.62 + 10^{-6}$
of CoS at 18° . (Weigel, Z. phys. Ch.
1894, 294.)

Insol. in conc. mineral acids; very sl. sol. in
dil. acids; scarcely sol. in acetic acid.
(Lebeau, C. R. 1902, 135.)

When still moist in $\text{SO}_2 + \text{Aq.}$ (Ber-

Not sol. in HNO_3 , but only very sl. sol.
in $\text{HCl} + \text{Aq.}$ Not pptd. from very dil.
solutions by H_2S .

Insol. in H_2O , alkalies, and alkali carbon-
ates + Aq. (Fresenius.)

Insol. in NH_4Cl , and $\text{NH}_4\text{NO}_3 + \text{Aq.}$

Not pptd. by $(\text{NH}_4)_2\text{S} + \text{Aq.}$ shows a
yellow colour in presence of 200,000 pts. H_2O .

Ammonium cobaltic acid, etc. does not hinder the
precipitation of $(\text{NH}_4)_2\text{S} + \text{Aq.}$ (Rose.)

Insol. in potassium thiocarbonate + Aq.
(Ladit, Z. anal. 26. 15.)

Insol. in Na_2S_x or $\text{K}_2\text{S}_x + \text{Aq.}$ (de Koninck,
Bull. chim. 1891. 202.)

disulphide, Co_2S_3 .

Decomposes by $\text{HCl} + \text{Aq.}$; sol. in
conc. $\text{HCl} + \text{Aq.}$ with decomposition.

Not attacked by $\text{HCl} + \text{Aq.}$; and slowly even
in aqua regia. (Schneider, J. pr. (2) 9. 209.)

Min. Cobalt pyrite.

$+x\text{H}_2\text{O}$. Insol. in $\text{KCN} + \text{Aq.}$ (Fleck,
J. pr. 97. 303.) More sol. in $\text{HCl} + \text{Aq.}$ than
 CoS_2 . (Dingler, Berz, J. B. 10. 139.)

Cobaltocobaltic sulphide, Co_2S_4 .

Min. *Linnæite*. Sol. in warm $\text{HNO}_3 + \text{Aq.}$,
with residue of S .

Cobalt disulphide, CoS_2 .

Not attacked by alkalies or acids except
 HNO_3 and aqua regia. (Setterberg, Pogg. 7.
40.)

Cobalt sulphide, Co_3S_2 .

Easily sol. in hot HCl with evolution of
 H_2S (and H_2 ?). (Proust.)

Cobalt potassium sulphide, $\text{K}_2\text{Co}_{11}\text{S}_{10}$.

Slowly sol. in cold HCl and aqua regia.
Quickly sol. in warm aqua regia.

Sol. in HF and H_2SO_4 only on warming.
Insol. in $(\text{NH}_4)_2\text{S}$, organic acids, alkalies,
12% $\text{HCl} + \text{Aq.}$ and $\text{KCN} + \text{Aq.}$ (Milbauer,
Z. anorg. 1904, 42. 447.)

Cobalt telluride, CoTe .

(Fabre, C. R. 105. 673.)

Cobalt decamine sulphurous acid.

See Decamine cobaltisulphurous acid.

Cobaltic acid.

**Potassium cobaltate, $\text{K}_2\text{Co}_2\text{O}_7 + 2\text{H}_2\text{O}$, or
 $3\text{H}_2\text{O}$.**

Insol. in H_2O (Pebal, A. 100. 262), but de-
composes by long boiling. Sol. in conc. acids.

K_2O , $x\text{CoO}_2$. Sol. in H_2O . (Winkler, J.
pr. 91. 351.)

Does not exist. (Donath, W. A. B. 102, 2b.
71.)

Cobalticyanhydric acid, $\text{H}_3\text{Co}(\text{CN})_6 + \frac{1}{2}\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O and only
sl. decomp. on boiling.

Sol. in $\text{HCl} + \text{Aq.}$ without decomp. even on
boiling. Sl. sol. in conc., more sol. in dil.
 $\text{HNO}_3 + \text{Aq.}$ Not decomp. by boiling conc.
 $\text{HNO}_3 + \text{Aq.}$ or aqua regia. Insol. in conc.,
sl. sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ Sol. in alcohol.
Insol. in ether. (Zwenger, A. 162. 157.)

**Ammonium cobalticyanide, $(\text{NH}_4)_3\text{Co}(\text{CN})_6$
 $+ \frac{1}{2}\text{H}_2\text{O}$.**

Very sol. in H_2O ; sl. sol. in alcohol.

**Ammonium barium cobalticyanide,
 $\text{NH}_4\text{BaCo}(\text{CN})_6 + \text{H}_2\text{O}$.**

Sol. in H_2O . (Weselsky.)

**Ammonium calcium cobalticyanide,
 $\text{NH}_4\text{CaCo}(\text{CN})_6 + 10\text{H}_2\text{O}$.**

Sol. in H_2O .

Ammonium lead cobalticyanide,
 $\text{NH}_4\text{PbCo}(\text{CN})_6 + 3\text{H}_2\text{O}.$

Sol. in 8.31 pts. H_2O at 18° , and sl. sol. in 93% alcohol. (Schuler.)

Ammonium mercuric cobalticyanide,
 $(\text{NH}_4)_6\text{Co}_2\text{Hg}(\text{CN})_{14} + \text{H}_2\text{O}.$

Sol. in H_2O with decomp.

Insol. in alcohol. (Soenderop, Dissert. 1899.)

Ammonium sodium cobalticyanide,
 $\text{NH}_4\text{Na}_2\text{Co}(\text{CN})_6.$

Only sl. sol. in H_2O . (Weselsky, B. 2. 598.)

Ammonium strontium cobalticyanide,
 $\text{NH}_4\text{SrCo}(\text{CN})_6 + 9\text{H}_2\text{O}.$

Sol. in H_2O . (W.)

Barium cobalticyanide, basic, $\text{Ba}_3[\text{Co}(\text{CN})_6]_2,$
 $\text{BaO}_2\text{H}_2.$

Not very stable. Cannot be recryst. without partial decomp. (W.)

Barium cobalticyanide, $\text{Ba}_3[\text{Co}(\text{CN})_6]_2 +$
 $10\text{H}_2\text{O}.$

Sl. efflorescent. Very sol. in H_2O . Insol. in alcohol.

Barium cobalticyanide chloride,
 $\text{Ba}_3[\text{Co}(\text{CN})_6]_2, \text{BaCl}_2 + 16\text{H}_2\text{O}.$

Sol. in H_2O without decomp. (W.)

Barium lithium cobalticyanide, $\text{BaLiCo}(\text{CN})_6,$
 $+15\text{H}_2\text{O}.$

The most. sol. of the double cobalticyanides. (Weselsky.)

Barium potassium cobalticyanide,
 $\text{BaKCo}(\text{CN})_6 + 11\text{H}_2\text{O}.$

Sol. in H_2O . (W.)

Bismuth cobalticyanide $\text{BiCo}(\text{CN})_6.$

Ppt. (Mathews, J. Am. Chem. Soc. 1900, 22. 275.)

$+5\text{H}_2\text{O}.$ Moderately stable with dil. min. acids; more stable with conc. acids than Cd or Zn comp.

Decomp. by NH_3 and alkalis. (Fischer and Cuntze, Ch. Z. 1902, 26. 872.)

Cadmium cobalticyanide, $\text{Cd}_3[\text{Co}(\text{CN})_6]_2 +$
 $7\frac{1}{2}\text{H}_2\text{O}.$

Attacked by strong min. acids only when hot. Behaves as Zn salt, toward conc. H_2SO_4 and dil. and conc. HCl .

Insol. in $\text{K}_2\text{Co}(\text{CN})_6 + \text{Aq}$. Sol. in NH_4OH and $\text{NH}_4\text{Cl} + \text{Aq}$. (Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

Cadmium potassium cobalticyanide,
 $\text{KCdCo}(\text{CN})_6.$

Not attacked by H_2O . (Fischer and Cuntze, Ch. Z. 1902. 26, 873.)

Cadmium sodium cobalticyanide,
 $\text{NaCdCo}(\text{CN})_6.$

(Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

Cadmium cobalticyanide ammonia,
 $\text{Cd}_3[\text{Co}(\text{CN})_6]_2, 4\text{NH}_3 + 2\text{H}_2\text{O}.$

(Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

$\text{Cd}_3[\text{Co}(\text{CN})_6]_2, 5\text{NH}_3 + 3\text{H}_2\text{O}.$
 and Cuntze, Ch. Z. 1902, 26. 873.)

$\text{Cd}_3[\text{Co}(\text{CN})_6]_2, 7\text{NH}_3.$ (Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

$\text{Cd}_3[\text{Co}(\text{CN})_6]_2, 9\text{NH}_3 + 2\text{H}_2\text{O}.$
 and Cuntze, Ch. Z. 1902, 26. 873.)

Calcium potassium cobalticyanide,
 $\text{CaKCo}(\text{CN})_6 + 9\text{H}_2\text{O}.$

Sol. in H_2O . (W.)

Cobaltous cobalticyanide, $\text{Co}_3[\text{Co}(\text{CN})_6]_2,$
 $14\text{H}_2\text{O}.$

Insol. in H_2O and acids. Sl. sol. in $+ \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$.

Cupric cobalticyanide, $\text{Cu}_3[\text{Co}(\text{CN})_6]_2,$
 $7\text{H}_2\text{O}.$

Insol. in H_2O and acids. Sol. in $\text{NH}_3 + \text{Aq}$.

Cupric cobalticyanide ammonia,
 $\text{Cu}_3[\text{Co}(\text{CN})_6]_2, 4\text{NH}_3 + 7\text{H}_2\text{O}.$

Sol. in H_2O . (Zwenger.)

Lead cobalticyanide, basic, $\text{Pb}_3[\text{Co}(\text{CN})_6]_2,$
 $3\text{PbO}_2\text{H}_2 + 11\text{H}_2\text{O}.$

Insol. in H_2O or alcohol; somewhat hot $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$. (Schuler.)

Lead cobalticyanide, $\text{Pb}_3[\text{Co}(\text{CN})_6]_2,$
 $7\text{H}_2\text{O}.$

Very sol. in H_2O . Insol. in alcohol. (Zwenger.)

$+7\text{H}_2\text{O}.$ Sol. in 1.77 pts. H_2O at 19° . Insol. in absolute H_2O . Sl. sol. in 93% alcohol. (Schuler, V 79. 302.)

Lead potassium cobalticyanide, $\text{PbKCo}(\text{CN})_6,$
 $+3\text{H}_2\text{O}.$

Sol. in 6.74 pts. H_2O at 18° and much more easily in hot H_2O . Insol. in absolute H_2O . Sl. sol. in 93% alcohol. (Schuler.)

Lead cobalticyanide nitrate, $\text{Pb}_3[\text{Co}(\text{CN})_6]_2,$
 $\text{Pb}(\text{NO}_3)_2 + 12\text{H}_2\text{O}.$

Sol. in 16.91 pts. H_2O at 18° , 10 at 19° , and much less hot H_2O .

Nearly insol. in 93% alcohol. (Schuler.)

Mercurous cobalticyanide, $\text{Hg}_2\text{Co}(\text{CN})_6.$

Ppt. Decomp. by HCl . Not attacked by cold, but by hot conc. H_2SO_4 . Not attacked by HNO_3 , acetic or oxalic acid. Dec. by alkalis $+ \text{Aq}$. (Miller and Mathews, Chem. Soc. 1900, 22. 64.)

Mercuric cobalticyanide, $\text{Hg}_2[\text{Co}(\text{CN})_6]_2$.

Sl. sol. in H_2O , decomp. by boiling.
Insol. in alcohol and ether. Not attacked by HCl . (Soenderop, Dissert, 1899.)

Mercuric potassium cobalticyanide, $\text{K}_4\text{HgCo}_2(\text{CN})_{14}$.

Sol. in H_2O with decomp. Insol. in alcohol. Sl. sol. in ether. (Soenderop, Dissert, 1899.)

$\text{K}_4\text{Hg}_2\text{Co}_4(\text{CN})_{14}$. (Soenderop, Dissert. 1899.)

Mercuric sodium cobalticyanide, $\text{Na}_4\text{Hg}_2\text{Co}_4(\text{CN})_{14} + 4\text{H}_2\text{O}$.

Extremely deliquescent. (Soenderop, Dissert, 1899.)

Nickel cobalticyanide, $\text{Ni}_2[\text{Co}(\text{CN})_6]_2 + 12\text{H}_2\text{O}$.

Insol. in H_2O and acids. Not attacked by boiling $\text{HCl} + \text{Aq}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$.

Nickel cobalticyanide ammonia, $\text{Ni}_2[\text{Co}(\text{CN})_6]_2, 4\text{NH}_3 + 7\text{H}_2\text{O}$.

Insol. in H_2O .

Potassium cobalticyanide, $\text{K}_3\text{Co}(\text{CN})_6$.

Easily sol. in H_2O . Insol. in alcohol.

Potassium strontium cobalticyanide, $\text{KSrCo}(\text{CN})_6 + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Weselsky.)

Potassium thallium cobalticyanide, $\text{K}_3\text{Tl}_2[\text{Co}(\text{CN})_6]_2$.

More sol. in H_2O than corresponding K salt. (Fischer and Benzian, Ch. Z. 1902, 26. 49.)

Potassium zinc cobalticyanide, $\text{KZnCo}(\text{CN})_6 + 3\text{H}_2\text{O}$.

(Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

Potassium cobalticyanide mercuric chloride, $2\text{K}_3\text{Co}(\text{CN})_6, 3\text{HgCl}_2$.

(Soenderop, Dissert. 1899.)

Potassium cobalticyanide mercuric iodide, $4\text{K}_3\text{Co}(\text{CN})_6, \text{HgI}_2$.

Sol. in H_2O with subsequent decomp.
Sol. in alcohol and ether with decomp. (Soenderop, Dissert, 1899.)

Silver cobalticyanide, $\text{Ag}_3\text{Co}(\text{CN})_6$.

Insol. in H_2O and acids. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Silver cobalticyanide ammonia, $\text{Ag}_3\text{Co}(\text{CN})_6, \text{NH}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O . (Zwenger.)

Sodium cobalticyanide, $\text{Na}_3\text{Co}(\text{CN})_6 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O ; insol. in alcohol.

Sodium zinc cobalticyanide, $\text{NaZnCo}(\text{CN})_6 + \text{H}_2\text{O}$.

(Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

Strontium cobalticyanide, $\text{Sr}_2[\text{Co}(\text{CN})_6]_2 + 10\text{H}_2\text{O}$.

Very sol. in H_2O . (Weselsky.)

Thallium cobalticyanide, $\text{Tl}_3\text{Co}(\text{CN})_6$.

100 pts. H_2O dissolve 3.6 pts. at 0° , 5.86 pts. at 9.5° , 10.04 pts. at 19.5° . (Fronmüller, B. 11. 91.)

Yttrium cobalticyanide, $\text{YCo}(\text{CN})_6 + 2\text{H}_2\text{O}$.

Nearly insol. in H_2O . (Cleve.)

Zinc cobalticyanide, $\text{Zn}_2[\text{Co}(\text{CN})_6]_2 + 12\text{H}_2\text{O}$.

Sol. in $\text{HCl} + \text{Aq}$ and salt is pptd. by dilution with H_2O . Decomp. by H_2SO_4 . Insol. in $\text{K}_4\text{Co}(\text{CN})_6 + \text{Aq}$. Sol. in NH_4OH and $\text{NH}_4\text{Cl} + \text{Aq}$. (Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

Zinc cobalticyanide ammonia, $\text{Zn}_2[\text{Co}(\text{CN})_6]_2, 5\text{NH}_3$.

Decomp. by H_2O and acids. (Fischer and Cuntze, Ch. Z. 1902, 26. 873.)

$\text{Zn}_2[\text{Co}(\text{CN})_6]_2, 6\text{NH}_3$. (Fischer and Cuntze.)

$+ 3\text{H}_2\text{O}$. (Fischer and Cuntze.)

$\text{Zn}_2[\text{Co}(\text{CN})_6]_2, 10\text{NH}_3 + 9\text{H}_2\text{O}$. Decomp. by H_2O . (Fischer and Cuntze.)

Cobaltimolybdic acid.

Ammonium barium cobaltous cobaltimolybdate, $\frac{1}{2}(\text{NH}_4)_2\text{O}, 1\frac{1}{2}\text{BaO}, \text{CoO}, \text{CoO}_2, 10\text{MoO}_3 + 18\frac{1}{2}\text{H}_2\text{O}$.

Difficultly sol. in H_2O . (Friedheim and Keller, B. 1906, 39. 4306.)

Ammonium cobaltous cobaltimolybdate, $2(\text{NH}_4)_2\text{O}, \text{CoO}, \text{CoO}_2, 10\text{MoO}_3 + 12\text{H}_2\text{O}$.

Much more sol. in H_2O than $3(\text{NH}_4)_2\text{O}, \text{CoO}, \text{CoO}_2, 12\text{MoO}_3 + 20\text{H}_2\text{O}$. Sp. gr. of cold sat. solution = 1.096. (Friedheim and Keller.)

$3(\text{NH}_4)_2\text{O}, \text{CoO}, \text{CoO}_2, 12\text{MoO}_3 + 20\text{H}_2\text{O}$.

100 cc. cold sat. aqueous solution contain 3 g. of the salt. Sp. gr. of the solution = 1.0234.

Sol. in conc. HCl .

Decomp. by conc. H_2SO_4 , by $\text{KOH} + \text{Aq}$ and by $\text{NaOH} + \text{Aq}$. (Friedheim and Keller.)

Barium cobaltous cobaltimolybdate, $3\text{BaO}, \text{CoO}, \text{CoO}_2, 9\text{MoO}_3 + 25\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Friedheim and Keller.)

Cobaltous potassium cobaltimolybdate,
 $\text{CoO}, 3\text{K}_2\text{O}, \text{CoO}_2, 10\text{MoO}_3 + 10\text{H}_2\text{O}$.
 (Kurnakoff, Ch. Z. 1890, 14. 113.)

+11H₂O. Sol. in conc. HCl. Decomp. by KOH+Aq and by NaOH+Aq. (Friedheim and Keller.)

$3\text{K}_2\text{O}, \text{CoO}, \text{CoO}_2, 12\text{MoO}_3 + 15\text{H}_2\text{O}$. Sl. sol. in H₂O. Sol. in conc. HCl. Decomp. by KOH+Aq and by NaOH+Aq. (Friedheim and Keller.)

+20H₂O. (Kurnakoff, Ch. Z. 1890, 14. 113.)

Potassium cobaltimolybdate,
 $3\text{K}_2\text{O}, \text{CoO}_2, 9\text{MoO}_3 + 6\frac{1}{2}\text{H}_2\text{O}$.

Ppt. (Hall, J. Am. Chem. Soc. 1907, 29. 703.)

Cobaltinitrocyanhydric acid.

Potassium cobaltinitrocyanide,
 $\text{K}_4\text{Co}_2(\text{CN})_8\text{NO}_2 + 3\text{H}_2\text{O}$.

Very sol. in H₂O but quickly decomp. Insol. in alcohol. (Rosenheim and Koppel, Z. anorg. 1898, 17. 68.)

Silver cobaltinitrocyanide,
 $\text{Co}_2\text{Ag}_3\text{NO}_2(\text{CN})_{10} + 6\text{H}_2\text{O}$, and +21H₂O.
 (Rosenheim and Koppel.)

Sodium cobaltinitrocyanide,
 $\text{Na}_4\text{Co}_2(\text{NO}_2)(\text{CN})_{10} + 11\text{H}_2\text{O}$.

Very deliquescent. Sol. in H₂O. (Rosenheim and Koppel.)

Cobaltisulphurous acid, H₂Co₂(SO₃)₆.

Not obtained in a solid state. (Berglund, Acta Lund. 1872.)

Cobaltisulphites.

The cobaltisulphites are insol. or at least very sl. sol. in H₂O. (Berglund, Acta Lund. 1872. 23.)

Ammonium cobaltous cobaltisulphite,
 $(\text{NH}_4)_2\text{SO}_3, 2\text{CoSO}_3, \text{Co}_2(\text{SO}_3)_3 + 14\text{H}_2\text{O} =$
 $(\text{NH}_4)_4\text{Co}_2\text{Co}_2(\text{SO}_3)_6 + 14\text{H}_2\text{O}$.

Scarcely sol. in H₂O, but decomp. thereby. Easily sol. in acids, when finely divided; also in H₂SO₄+Aq. (Berglund.)

$2(\text{NH}_4)_2\text{SO}_3, \text{CoSO}_3, \text{Co}_2(\text{SO}_3)_3 + 8\text{H}_2\text{O} =$
 $(\text{NH}_4)_4\text{CoCo}_2(\text{SO}_3)_6 + 8\text{H}_2\text{O}$. As above. (Berglund.)

Barium cobaltisulphite, 3BaSO₃, Co₂(SO₃)₃ + 12H₂O = Ba₃Co₂(SO₃)₆ + 12H₂O.

Ppt. Insol. in H₂O. Not attacked by cold acids even H₂SO₄, but is decomp. by boiling therewith. (Berglund, Acta Lund. 1872.)

Bismuth cobaltisulphite, Bi₂Co₂(SO₃)₆.

Insol. in H₂O, dil. HNO₃, or HCl+Aq. (Berglund, Acta Lund. 1872. 31.)

Calcium cobaltisulphite, Ca₃Co₂(SO₃)₆.
 Ppt. Insol. in H₂O or HCl+Aq. (Berglund, Acta Lund. 1872. 30.)

Cobaltous cobaltisulphite, Co₂Co₂(SO₃)₆.
 $3\text{CoSO}_3, \text{Co}_2(\text{SO}_3)_3$.
 Ppt. (Berglund, B. 7. 470.)

Cobaltous potassium cobaltisulphite,
 $\text{CoK}_4\text{Co}_2(\text{SO}_3)_6$.
 Insol. in H₂O. (Berglund.)

Silver cobaltisulphite, Co₂(SO₃)₃, 3.
 Properties as the following comp. (Berglund.)

Silver cobaltous cobaltisulphite, Co₂(SO₃)₃, 2Ag₂SO₃ + 9H₂O.
 Insol. in H₂O. Insol. in HNO₃+Aq. comp. by HCl or H₂S+Aq. (Berglund.)

Sodium cobaltous cobaltisulphite.

Decomp. by H₂O, so that it has not been obtained pure. (Berglund, Acta Lund. 29.)

Cobaltoctamine sulphurous acid.

See Octamine cobaltisulphurous acid.

Cobaltocobalticyanhydric acid,
 $\text{H}_2\text{Co}_2(\text{CN})_{11}$.

Unstable. (Jackson and Comey, A. J. 1897. 19, 277.)

Barium cobaltocobalticyanide,
 $\text{BaHCo}_2(\text{CN})_{11} + 1\frac{1}{2}\text{H}_2\text{O}$.

Somewhat sol. in H₂O when pure. The crude salt is insol. even in hot H₂O. (Jackson and Comey.)

Cupric cobaltocobalticyanide, Cu₂Co₂(CN)₁₁ + 4H₂O.

Ppt. (Jackson and Comey.)

Potassium hydrogen cobaltocobalticyanide,
 $\text{K}_2\text{HCo}_2(\text{CN})_{11} + 2\text{H}_2\text{O}$.

Sl. sol. in cold, easily sol. in hot H₂O. Insol. in alcohol. (Jackson and Comey.)
 $\text{KH}_2\text{Co}_2(\text{CN})_{11} + \text{H}_2\text{O}$. Insol. in hot H₂O when impure.

The pure salt is slowly sol. in cold H₂O. More sol. in warm H₂O. (Jackson and Comey.)

Silver cobaltocobalticyanide, Ag₂Co₂(CN)₁₁ + H₂O.

Ppt. (Jackson and Comey, B. 16. 1021.)

Zinc cobaltocobalticyanide, ZnHCo₂(CN)₁₁ + 3H₂O.

Ppt. (Jackson and Comey.)

ocyanhydric acid, $H_4Co(CN)_4$.

unstable. Sol. in H_2O . Insol. in

potassium cobaltocyanide, $K_3Co(CN)_6$.

us, Z. anorg. 1895, 9. 17.)

um cobaltocyanide, $K_4Co(CN)_6$.

mp. on air. Very deliquescent, and
 H_2O . Insol. in alcohol and ether.
nps, Zeit. Ch. 1888. 952.)

ous acid.

cobaltite, $BaCoO_3$.

. in H_2O or dil. HCl , H_2O_2 + Aq. Sol.
+ Aq. (Rousseau, C. R. 109. 64.)
 H_2O_2 . As above. (Rousseau.)

**us potassium cobaltite, $3CoO_2$, CoO ,
 O .**

dly hydrolysed by H_2O .
in conc. HCl . (Bellucci, Chem. Soc.
2, (2) 354.)

sium cobaltite, $MgCoO_3$.

. in H_2O , NH_4OH , or $(NH_4)_2CO_3$ + Aq.
sol. in NH_4Cl + Aq, from which it is
by KOH + Aq. (Berzelius, Pogg. 33.

in HF , HCl , HNO_3 + H_2SO_4 ; decomp.
tially dissolved by NH_4OH + Aq; quite
when heated. (Dufau, C. R. 1896,
0.)

um cobaltite.

rding to Bellucci and Dominici the
nds formerly described are more or
comp. by hydrolysis. (C. C. 1907.
)

cobaltite.

n $NaOH$ + Aq, but pptd. by diluting
tion.

**bic acid (Niobic acid), $3Cb_2O_5$,
 $2O$, or $3Cb_2O_5$, $7H_2O$.**

y sol. in HF ; very sl. sol. in HCl + Aq,
sol. in H_2O after being treated with
Aq. Sol. in conc. H_2SO_4 . Sol. in
Aq. Insol. in $NaOH$ + Aq, but be-
sol. in H_2O by being treated with
+ Aq. Sol. in boiling Na_2CO_3 + Aq.
Pogg. 113. 109.)

. in liquid NH_3 . (Gore, Am. Ch. J.
. 830.)

, $4H_2O$.

, $7H_2O$. (Santesson, Bull. Soc. (2)

um columbate, Al_2O_3 , $3Cb_2O_5$ + $12H_2O$.

(E. F. Smith, J. Am. Chem. Soc.
. 1652.)

Barium columbate, $7BaO$, $6Cb_2O_5$ + $18H_2O$.

Ppt. (Bedford, J. Am. Chem. Soc. 1905,
27. 1218.)

Cadmium columbate, CdO , Cb_2O_5 .

Sol. in boiling conc. H_2SO_4 ; insol. in most
acids; decomp. by $HKSO_4$ at red heat. (Lars-
son, Z. anorg. 1896, 12. 199.)

+ $3\frac{1}{2}H_2O$. Ppt. (E. F. Smith, J. Am.
Chem. Soc. 1908, 30. 1652.)

Cæsium columbate, $4Cs_2O$, $3Cb_2O_5$ + $14H_2O$.

Very sol. in H_2O . (E. F. Smith, J. Am.
Chem. Soc. 1908, 30. 1654.)

$7Cs_2O$, $6Cb_2O_5$ + $30H_2O$. Ppt. (E. F.
Smith, J. Am. Chem. Soc. 1908, 30. 1655.)

Calcium columbate, $2CaO$, Cb_2O_5 .

Insol. in H_2O . (Joly, C. R. 81. 266.)

CaO , Cb_2O_5 . Sol. in boiling conc. H_2SO_4 ;
insol. in most acids; decomp. by $HKSO_4$ at
red heat. (Larsson, Z. anorg. 1896, 12. 198.)

Cobalt columbate, CoO , Cb_2O_5 .

Sol. in conc. boiling H_2SO_4 ; insol. in most
acids; decomp. by $HKSO_4$ at red heat. (Lars-
son.)

Copper columbate, CuO , Cb_2O_5 .

Sol. in boiling conc. H_2SO_4 ; insol. in most
acids; decomp. by $HKSO_4$ at red heat. (Lars-
son.)

+ $3\frac{1}{2}H_2O$. Ppt. (E. F. Smith, J. Am.
Chem. Soc. 1908, 30. 1652.)

Iron (ferrous) columbate, $Fe(CbO_3)_2$.

Min. *Columbite*. Insol. in acids.

Iron (ferrous) columbate tantalate,

$xFe(TaO_3)_2$, $yFe(CbO_3)_2$.

Min. *Tantalite*. Not attacked by acids.

$Fe(CbO_3)_2$, $4Fe(TaO_3)_2$. Min. *Tapiolite*.

Lithium columbate, $7Li_2O$, $6Cb_2O_5$ + $26H_2O$.

Ppt. (E. F. Smith, J. Am. Chem. Soc.
1908, 30. 1655.)

Magnesium columbate, MgO , Cb_2O_5 .

Sol. in boiling conc. H_2SO_4 ; insol. in most
acids; decomp. by $KHSO_4$ at red heat. (Lars-
son, Z. anorg. 1896, 12. 196.)

+ $4H_2O$. Precipitate. (Rammelsberg.)

+ $7H_2O$. Ppt. (E. F. Smith, J. Am.
Chem. Soc. 1908, 30. 1651.)

$4MgO$, Cb_2O_5 . Insol. in H_2O . (Joly, C. R.
81. 266.)

$3MgO$, Cb_2O_5 . As above.

Manganous columbate.

Insol. in H_2O . (Joly, C. R. 81. 266.)

$3MnO$, $5Cb_2O_5$. Sol. in boiling conc. H_2SO_4 ;
insol. in most acids; decomp. by $HKSO_4$ at
red heat. (Larsson, Z. anorg. 1896, 12. 201.)

Potassium columbate, KCbO_3 .

Sol. in H_2O . (Joly, in Fremy's Encyc. Ch.)
 $\text{K}_2\text{Cb}_4\text{O}_7 + 5\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O . (Santesson.)

$\text{K}_2\text{Cb}_5\text{O}_{10} + 5\text{H}_2\text{O}$. Nearly insol. in H_2O .
 $\text{K}_4\text{Cb}_2\text{O}_7 + 11\text{H}_2\text{O}$. Insol. in H_2O . (Santesson, Bull. Soc. (2) 24. 53.)

$\text{K}_4\text{Cb}_3\text{O}_{12} + 11\text{H}_2\text{O}$. (Santesson.)
 $\text{K}_5\text{Cb}_4\text{O}_{13} + 13\text{H}_2\text{O}$. Sol. in H_2O .
 $\text{K}_5\text{Cb}_5\text{O}_{15} + 16\text{H}_2\text{O}$. Efflorescent. Sol. in H_2O . (Marignac, A. ch. (4) 8. 20.)

Very sol. in H_2O . (E. F. Smith, J. Am. Chem. Soc. 1908, 80. 1652.)

$\text{K}_{11}\text{Cb}_{12}\text{O}_{37} + 27\text{H}_2\text{O}$. Sol. in H_2O . Insol. in alcohol. (E. F. Smith.)

$\text{K}_{16}\text{Cb}_{14}\text{O}_{43} + 32\text{H}_2\text{O}$. Sol. in H_2O .

Potassium sodium columbate, $3\text{K}_2\text{O}, \text{Na}_2\text{O}, 3\text{Cb}_2\text{O}_5 + 9\text{H}_2\text{O}$.

Very slightly sol. in H_2O . Insol. in alkalies. (Marignac.)

Rubidium columbate, $3\text{Rb}_2\text{O}, 4\text{Cb}_2\text{O}_5 + 9\frac{1}{2}\text{H}_2\text{O}$.

(E. F. Smith, J. Am. Chem. Soc. 1908, 30. 1655.)

$4\text{Rb}_2\text{O}, 3\text{Cb}_2\text{O}_5 + 14\text{H}_2\text{O}$. Very sol. in H_2O . (E. F. Smith.)

Silver columbate, $\text{Ag}_2\text{O}, \text{Cb}_2\text{O}_5 + 2\text{H}_2\text{O}$.

Ppt. (E. F. Smith.)

$7\text{Ag}_2\text{O}, 6\text{Cb}_2\text{O}_5 + 5\text{H}_2\text{O}$. Insol. in H_2O . (Bedford, J. Am. Chem. Soc. 1905, 27. 1218.)

Sodium columbate, $\text{NaCbO}_3 + 3\frac{1}{2}\text{H}_2\text{O}$.

Completely sol. in H_2O . (Rose.)

Ppt. (E. F. Smith, J. Am. Chem. Soc. 1908, 30. 1651.)

$+ 2\frac{1}{2}\text{H}_2\text{O}$. Sl. sol. in cold H_2O . Insol. in $\text{NaOH} + \text{Aq}$. (Santesson.)

$2\text{Na}_2\text{O}, 3\text{Cb}_2\text{O}_5 + 9\text{H}_2\text{O}$. Insol. in H_2O or $\text{NaOH} + \text{Aq}$. (Santesson.)

$8\text{Na}_2\text{O}, 7\text{Cb}_2\text{O}_5$. 1 pt. is sol. in 195–200 pts. H_2O at 14–20°; in ether 75–80 pts. or in 103 pts. boiling water. (Rose.)

$7\text{Na}_2\text{O}, 6\text{Cb}_2\text{O}_5 + 32\text{H}_2\text{O}$. Very stable. Sol. in H_2O . (Bedford, J. Am. Chem. Soc. 1905, 27. 1217.)

Thorium columbate, $5\text{Th}_2\text{O}_3, 16\text{Cb}_2\text{O}_5$.

Sol. in boiling conc. H_2SO_4 ; insol. in most acids; decomp. by HKSO_4 at red heat. (Larson, Z. anorg. 1896, 12. 202.)

Yttrium columbate, $\text{Y}_2\text{O}_3, \text{Cb}_2\text{O}_5$.

Insol. in H_2O . (Joly, C. R. 81. 1261.)

Sol. in boiling conc. H_2SO_4 ; insol. in most acids; decomp. by HKSO_4 at red heat. (Larson.)

Zinc columbate, $\text{ZnO}, \text{Cb}_2\text{O}_5$.

Sol. in boiling conc. H_2SO_4 ; insol. in most acids; decomp. by HKSO_4 at red heat. (Larson.)

$7\text{ZnO}, 6\text{Cb}_2\text{O}_5 + 25\text{H}_2\text{O}$. Insol. in H_2O . (Bedford, J. Am. Chem. Soc. 1905, 27. 1218.)

Zirconium columbate, $\text{ZrO}_2, 5\text{Cb}_2\text{O}_5$.

Sol. in boiling conc. H_2SO_4 ; insol. in most acids; decomp. by HKSO_4 at red heat. (Larson.)

Percolumbic acid.

See Percolumbic acid.

Columbium (Niobium), Cb .

Scarcely attacked by HCl , HNO_3 , or aqua regia. Conc. H_2SO_4 dissolves easily on warming.

Sol. in fused oxidizing agents; sol. in hot conc. H_2SO_4 and in HF ; also in $\text{HF} + \text{HNO}_3$; insol. in other acids. (Moissan, C. R. 1901, 133. 24.)

Columbium pentabromide, CbBr_5 .

(Rose, Pogg. 104. 422.)

Columbium carbide nitride, $3\text{CbC}, 2\text{CbN}$.

(Joly, Bull. Soc. (2) 25. 508.)

Columbium trichloride, CbCl_3 .

Not deliquescent; not attacked by H_2O , but easily oxidised by $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Roscoe, C. N. 37. 25.)

Columbium pentachloride, CbCl_5 .

Decomp. by H_2O with separation of a hydrate of Cb_2O_5 . Sol. in cold $\text{HCl} + \text{Aq}$, forming a solution which soon gelatinises, and separates out Cb_2O_5 by heat or dilution; with hot $\text{HCl} + \text{Aq}$, forms a cloudy solution which does not gelatinise. Sol. in H_2SO_4 to form a clear liquid which gelatinises on heating. Sol. in $\text{KOH} + \text{Aq}$. Sol. in alcohol with slight residue. (Rose, Pogg. 104. 432.)

Columbium pentafluoride, CbF_5 .

Very hygroscopic; sol. in H_2O without separation of columbic acid. (Ruff, B. 1900, 42. 492.)

Columbium fluoride with MF.

See Fluocolumbate, M.

Columbium hydride, $\text{CbH}(\text{?})$.

Insol. in HCl , HNO_3 , and dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, even on boiling. Sol. in boiling conc. H_2SO_4 , and in fused KHSO_4 . Sol. in cold $\text{HF} + \text{Aq}$ if not too dilute. Also attacked by $\text{KOH} + \text{Aq}$. (Marignac, N. Arch. Phys. Nat. 31. 89.)

Not attacked by boiling H_2O , or boiling HCl .

Oxidized by hot H_2SO_4 . Insol. in boiling HNO_3 . (Muthmann, A. 1907, 355. 90.)

Columbium hydroxide $\text{Cb}_2\text{O}_5, x\text{H}_2\text{O}$.

See Columbic acid.

nitride.

cked by boiling nitric acid or aqua ol. in a cold mixture of HNO_3 and , Pogg. 111. 426.)

Not attacked by boiling H_2O or l. in conc. HNO_3 , and H_2SO_4 .

by fused KOH . Not attacked by h $\text{KOH} + \text{Aq.}$ (Muthmann, A. 34.)

dioxide, Cb_2O_2 .

n still moist in boiling dil. $\text{HCl} +$ in hot HNO_3 ; less sol. in aqua in $\text{HCl} + \text{Aq.}$ Sol. in conc. H_2SO_4 heating. (Rose.)

H_2O , KOH , or conc. acids, even ng. (Delafontaine.)

trioxide, Cb_2O_3 .

acids except HF . (Smith, Z. 7. 28.)

tetroxide, Cb_2O_4 .

acked by cold or hot H_2O , HCl , SO_4 , or aqua regia. Slightly at-boiling $\text{KOH} + \text{Aq.}$ (Delafon-

pentoxide, Cb_2O_5 .

mitted insol. in hot conc. H_2SO_4 . is not been ignited it forms a clear ith H_2SO_4 , which can be diluted rming any precipitate. (Rose, 549.)

ised KHSO_4 , which can be diluted without causing pptn. Insol. in

oxybromide, CbOBr_2 .

by H_2O into Cb_2O_3 and HBr . Sol. SO_4 and conc. $\text{HCl} + \text{Aq.}$ (Rose, 442.)

Cb_2O_4 , $\text{HBr}(\text{?})$. Easily sublimed. anorg. 1894, 7. 97.)

oxybromide rubidium bromide, r_2 , 2RbBr .

in moist air. Decomp. by H_2O . B. 1903, 39. 3059.)

oxychloride, CbOCl_2 .

H_2O from air without deliquescing posed. Decomp. with H_2O with f heat. Insol. in hot or cold $\text{HCl} +$ by long contact with H_2SO_4 to a uid, which clears up on warming, eparates out Cb_2O_3 . Sol. in cold and hot $\text{K}_2\text{CO}_3 + \text{Aq.}$ (Rose.)cohol, from which it is precipitated (Blomstrand.)

Cb_2O_4 , HCl . Sublimate. (Smith, 894, 7. 97.)

oxychloride rubidium chloride, l_2 , 2RbCl .

by H_2O . (Weinland, B. 1906,

Columbium oxyfluoride, CbOF_2 .

(Joly, C. R. 81. 1266.)

Columbium oxyfluoride with MF.

See Fluoxycolumbate, and Fluoxyhypocolumbate, M.

Columbium oxysulphide, Cb_2OS_2 .

Insol. in boiling $\text{HCl} + \text{Aq.}$ Slowly decomp. into Cb_2O_3 by boiling with HNO_3 or aqua regia. Insol. in boiling dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ Converted into columbic sulphate, sol. in H_2O , by boiling conc. H_2SO_4 . Sl. sol. in hot HF . Insol. in boiling $\text{K}_2\text{S} + \text{Aq.}$ (Rose, Pogg. 111. 193.)

Copper, Cu.

Copper is not attacked by distilled H_2O , or by NH_4NO_3 , KNO_3 , or $(\text{NH}_4)_2\text{SO}_4 + \text{Aq.}$, or by a mixture of those salts in solution. (Muir, cited by Carnelly, Chem. Soc. 30. 1.)

Distilled H_2O has slight action on Cu. 100 ccm. H_2O dissolved from 2 sq. dcm. Cu from 0.035 mg. Cu in one hour up to 0.280 mg. in 72 hours. 100 ccm. H_2O dissolved 0.44 mg. from 6 sq. dcm. in 48 hours. Presence of solder diminishes solubility about one-half. At 90–100° the amount dissolved is about one-half that at ord. temp. (Carnelley, Chem. Soc. 30. 1.)

100 ccm. distilled H_2O dissolved only 1 mg. Cu from 11.8 sq. cm. during a week, while air free from CO_2 was conducted through the solution. When the air contained CO_2 , 3 mg. were dissolved. (Wagner, Dingl. 221. 259.)

100 l. sea water dissolved 12.96 g. Cu from 1 sq. m. (Calvert and Johnson, C. N. 11. 171.)

Solubility in H_2SO_4 .

Not attacked by dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Vogel, Schw. J. 32. 301.)

Action of H_2SO_4 at ordinary temp. is very slight even after a long time. (Barruel, J. Pharm. 20. 13 [1834].)

H_2SO_4 has no action below 130°. (Calvert and Johnson, Chem. Soc. 19. 438.)

H_2SO_4 acts slightly even at 20°.

16.3 g. H_2SO_4 (1.843 sp. gr.) dissolved the following amts. from 3 g. Cu, having a surface of 65 sq. cm. at the given temp.

Temp.	Time	% Cu dissolved
19°	14 days	About 6
60	120 min.	2.5
80	30 "	1.5
100	30 "	3.1
124	30 "	22.7
130	30 "	32.6
137	30 "	35 0
150	30 "	69.2
170	10 "	51.92
195	2 "	53.5
220	½ "	70.57
270	few seconds	nearly 100

With dilute acid the action was much less violent, as is seen in the following table—

Tem.	Time	Acid	Sp. gr.	% Cu dissolved
100°	30 min.	H ₂ SO ₄	1.843	2.380
100	30 "	2H ₂ SO ₄ , H ₂ O	1.8295	0.585
100	30 "	H ₂ SO ₄ , H ₂ O	1.780	0
100	30 "	H ₂ SO ₄ , 2H ₂ O	1.620	0
130	30 "	H ₂ SO ₄	1.843	32.6
130	30 "	H ₂ SO ₄ , H ₂ O	1.780	1.18
130	30 "	H ₂ SO ₄ , 2H ₂ O	1.620	0
165	15 "	H ₂ SO ₄	1.843	70
165	30 "	H ₂ SO ₄ , H ₂ O	1.780	16.5
165	30 "	H ₂ SO ₄ , 2H ₂ O	1.620	2.7

(Pickering, Chem. Soc. 33. 112.)

Cu is very sl. attacked by cold HCl + Aq of 1.12 sp. gr., but somewhat more on warming. Even less sol. in dil. HCl + Aq. (Löwe, Z. anal. 4. 361.)

Sol. in warm conc. HI + Aq. (Rose.)

Slowly attacked by H₂SO₃ + Aq. (Causse, Bull. Soc. (2) 45. 3.)

More or less sol. in all dil. mineral acids and also in organic acids, as acetic, tartaric, etc., when supply of air is afforded; but absolutely insol. in the latter acids when air is wholly excluded. The importance of this fact in the use of Cu cooking utensils is manifest.

Easily attacked by ord. HNO₃ + Aq.

With very conc. HNO₃ + Aq (sp. gr. 1.52) it becomes passive, as in the case of Fe.

Pure dil. HNO₃ + Aq of 1.07 sp. gr. or less does not attack Cu at 20°, but if NO₂ or KNO₃ is added the action begins at once. If HNO₃ + Aq is more conc. the Cu is attacked. HNO₃ + Aq of 1.108 sp. gr. begins to act at -2°, and of 1.217 sp. gr. at -10°.

HNO₃ + Aq of 1.512 sp. gr. attacks Cu violently at 20°, but action soon ceases on account of formation of a crust of Cu(NO₃)₂, insol. in pure HNO₃. (Millon, A. ch. (3) 6. 95.)

Easily sol. in 2N-HClO₄ + Aq at 50°. (Hendrixson, J. Am. Chem. Soc. 1904, 26. 756.)

Not appreciably sol. in anhydrous HF. (Poulenc, A. ch. 1894, (7) 2. 12.)

When in contact with the air, Cu is soon oxidised by acids, alkalies (especially NH₄OH + Aq), and many fatty bodies.

Sol. in (NH₄)₂CO₃ + Aq. (Traube, B. 10. 1887.)

Slowly sol. in NH₄OH + Aq. (Schonbein, B. A. B. 1856. 580.)

Sol. in KI + Aq when warm and conc. (Rose.)

When finely divided, Cu is easily sol. in hot FeCl₃ + Aq.

Action of dilute solutions of salts on solubility of Cu in H₂O.

100 ccm. solution of the following salts dissolve the amts. of Cu given below, from surface of 1 sq. dm. in 48 hours.

Salts	G. salt dissolved in 100 ccm. H ₂ O	Mg. Cu dissolved
H ₂ O	...	0.11
KNO ₃	0.01 0.05 5.00	0.07 0.13 0.16
NaNO ₃	0.05 5.00	0.18 0.19
CaSO ₄	0.05	0.11
K ₂ SO ₄	0.05 5.00	0.12 0.28
MgSO ₄	0.05 5.00	0.16 0.34
Na ₂ CO ₃	0.01 0.05 5.00	0.05 0.11 2.80
K ₂ CO ₃	0.05 5.00	0.14 2.35
NaCl	0.01 0.05 5.00	0.05 0.18 7.50
KCl	5.00	8.17
(NH ₄) ₂ SO ₄	0.05 5.00	0.66 28.50
NH ₄ NO ₃	0.01 0.05 5.00	0.17 0.66 60.00
NH ₄ Cl	0.05 5.00	0.92 158.75

At 100° the action of KNO₃, K₂SO₄, and NH₄NO₃ is diminished, while that of (NH₄)₂SO₄, Na₂CO₃, and NaCl is increased.

Tables are also given for mixtures of the above salts. (Carnelley, Chem. Soc. 80. 1. Solubility of Cu in dilute salt solutions.)

11.8 sq. cm. Cu were used, and the action continued one week, while air with or without CO₂ was passed through the solution continually.

ccm. solution of the following salts dis-
the given amts. Cu.

	G. salt dis- solved in 100 ccm. H ₂ O	Mg. Cu dis- solved with- out CO ₂	Mg. Cu dissolved with CO ₂
1	0.50	4	115
1	0.50	4	115
l ₂	0.83	5	112
Cl	1.00	904	138
l ₄	1.00	0	4
l ₃	1.00	0	3
O ₂	1.00	0	...
H	0.923	0	...
H ₂	sat.	0	...

(Wagner, Dingl. 221. 260.)

illed H₂O dissolved no Cu from 420 sq.
150 hours at ord. temp.
NO₂+Aq with less than 0.4 g. per litre
l the same result.
O₂+Aq or (NH₄)₂SO₄+Aq contain-
to 0.2 g. per litre dissolved no Cu.
containing carbonates+nitrates, car-
s+sulphates, or chlorides+nitrates
ssolved no Cu.
NO₂+Aq containing 0.4 g. per litre
ed 3 mg. per litre after 150 hours
t.
n a surface of 2100 sq. m. of Cu, H₂O
d with CO₂ at ord. pressure, and con-
g the following salts in solution, dis-
the given amts. Cu. in 120 hours.

Salt	G. salt dissolved in 1 l. H ₂ O	Mg. Cu dissolved
I ₂ O	1.0
CO ₂	0.2	0.2
aCl ₂	0.2	1.80
l ₂ NO ₂	0.02	1.40
l ₄ NO ₂	0.04	1.40
CO ₂ + +NO ₂	0.1 0.02	1.00
CO ₂ + +NO ₂	0.2 0.04	0.1
+NO ₂ + aCl ₂	0.2 0.2	3.6

n a surface of 2100 sq. m., H₂O charged
O₂ at pressure of 6 atmos. dissolved 0.6
48 hours.
when charged with CO₂ at 6 atmos. and
ning:
mg. NH₄NO₂ per litre, dissolved 0.8 mg.
ours.
mg. NH₄NO₂ per litre, dissolved 1.4
48 hours.
ng. K₂CO₃ per litre, dissolved 1.2 mg
hours. (Muir, Proc. Soc. Manchester,
.)
in KCN+Aq. (Goyder, C. N. 1894,
2.)
lution of (NH₄)₂S₂O₃ containing 0.829 g.

in 110 cc. dissolves 0.2050-0.2279 g. Cu.
(Turrentine, J. phys. Chem. 1907, 11. 625.)
Sl. attacked by liquid NH₃. (Franklin,
Am. Ch. J. 1898, 20. 827.)

Amts. Cu dissolved by action of various oils
on 8 sq. in. Cu by 10 days' exposure and
subsequent 67 days—

	Amt. Cu dis- solved in 10 days	Amt. Cu dissolved in subsequent 67 days
Linseed oil .	0.3000 grain	0.2435 grain
Olive oil . .	0.2200 "	0.0200 "
Colza oil . .	0.0170 "	0.1230 "
Almond oil .	0.1030 "	0.1170 "
Seal oil . .	0.0485 "	0.0315 "
Sperm oil .	0.0030 "	0.0575 "
Castor oil .	0.0065 "	0.0035 "
Neatsfoot oil	0.1100 "
Sesame oil .	0.1700 "	0.0015 "
Paraffine oil .	0.0015 "

(Watson, C. N. 36. 200.)

Qualitative results of the action of various
oils on Cu are also given by Thompson.
(C. N. 34. 176, 200, 219.)
½ ccm. oleic acid dissolves 0.0157 g. Cu
in 6 days. (Gates, J. phys. Chem. 1911,
15. 143.)
Sol. in an alkaline solution of gelatine (3.54
%) copper gauze dissolved in 48 hours.
(Lidoff, C. C. 1899, II. 471.)

Cuprous acetylide, Cu₂C₂.

Decomp. by heating with H₂O or KCN +
Aq. Decomp. by HNO₃. (Keiser, Am. Ch.
J. 1892, 14. 289.)
Not decomp. by H₂SO₄, NH₄OH, KOH +
Aq or acetic acid, even on warming. The
dry salt is sol. in very dil. HCl+Aq without
evolution of gas. Sol. in conc. KCN+Aq.
(Böttger, A. 1859, 109. 356.)

Cupric acetylide, CuC₂.

Easily sol. in HCl. Turns brown in the air,
and becomes insol. in acids. (Phillips, Z.
anorg. 1894, 6. 241.)
3Cu₂C₂O+2H₂O. Solubility as that of
Cu₂C₁₇H₄O₃. (Söderbaum, B. 1897, 30. 764.)
Cu₂C₁₇H₄O₃. Insol. in H₂O.
When dry is violently decomp. by conc.
H₂SO₄ or not too dil. HNO₃. Rapidly de-
comp. by warming with dil. acids, especially
HCl.
Insol. in NH₄OH+Aq in absence of air,
partially sol. in presence of air.
Insol. in organic solvents. (Söderbaum,
B. 1897, 30. 762.)

Cuprous acetylide iodide, Cu₂C₂I, CuI.

Ppt. (Berthelot and Delépine, A. ch.
1900, (7) 19. 54.)

Cupric arsenide, Cu_3As_2 .(Reinsch, J. pr. **24**. 244.) Cu_3As_2 . (Gehlen.) Cu_3As_2 . Ppt. Decomp. by acids. (Kane, Pogg. **44**. 471.) Cu_3As . Min. *Domeykite*. Insol. in HCl + Aq; sol. in HNO_3 . Cu_3As . Min. *Algodonite*. Cu_3As . Min. *Darwinite*.**Cuprous azoimide, CuN_3 .**Insol. in H_2O . (Curtius.)Sensitive to sunlight. (Wöhler, B. 1913, **46**. 2053.)**Cupric azoimide, basic, CuO , CuN_3 .**Insol. in H_2O . (Wöhler, B. 1913, **46**. 2055.)**Cupric azoimide, CuN_3 .**

Very explosive.

Very sl. sol. in H_2O . Decomp. by boiling with H_2O . (Curtius, J. pr. 1898, (2) **58**. 296.)**Copper azoimide ammonia, CuN_3 , 2NH_3 .**Ppt. Insol. in H_2O . Easily sol. in dil. acids. (Dennis, J. Am. Chem. Soc. 1907, **29**, 19.)**Copper boride, Cu_3B_2 .**

(Marsden, J. B., 1880. 330.)

Cuprous bromide, Cu_2Br_2 .1 l. H_2O dissolves at 18° – 20° :

0.4320 millimols bromine.

0.3157 " cupric copper.

0.1061 " cuprous copper.

(Bodländer, Z. anorg. 1902, **31**. 460.)Sol. in HBr , HCl without decomp., or HNO_3 + Aq with decomp., also in NH_4OH + Aq. Insol. in boiling conc. H_2SO_4 or $\text{HC}_2\text{H}_3\text{O}_2$ + Aq. (Berthémot, A. ch. **44**. 385.)Sol. in H_2SO_3 + Aq. (Lean and Whatmough, Chem. Soc. 1898, **73**. 151.)Sol. in NaCl , and $\text{Na}_2\text{S}_2\text{O}_3$ + Aq. (Renault, C. R. **59**. 319.)Solubility of Cu_2Br_2 in KBr + Aq.

All values recorded in millimols per litre.

KBr	Total copper	Cupric copper	Cuprous copper
25	0.119	0.012	0.107
40	0.200	0.013	0.187
60	0.310	0.025	0.285
80	0.423	0.012	0.411
100	0.5836	...	0.5836
120	0.6934	...	0.6934
500	8.719	...	8.719

(Bodländer and Storbeck, Z. anorg. 1902, **31**. 462.)Difficultly sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)Sol. in ethyl acetate. (Naumann, B. 1904, **37**. 314.)Insol. in acetone. (Naumann, B. 1904, **37**. 4329; Eidmann, C. C. 1897, II. 1014.)100 g. acetonitrile dissolve 3.86 g. Cu_2Br_2 at 18° . (Naumann and Schier, B. 1914, **47**. 249.)Sol. in pyridine. (Naumann, B. 1904, **37**. 4609.)Mol. weight determined in pyridine, methyl and ethyl sulphides. (Werner, Z. anorg. 1897, **15**. 19, 26, and 28.)**Cupric bromide, CuBr_2 .**Deliquescent. Very sol. in H_2O . Insol. in benzene. (Franchimont, B. **16**. 387.)Very sl. attacked by cold or even hot H_2SO_4 . (Viard, C. R. 1902, **135**. 169.)Moderately sol. in liquid NH_3 . (Horn, Am. Ch. J. 1908, **39**. 219.)100 g. 95% formic acid dissolve 0.16 g. CuBr_2 at 21° . (Aschan, Ch. Z. 1913, **37**. 1117.)100 g. acetonitrile dissolve 24.43 g. CuBr_2 at 18° . (Naumann and Schier, B. 1914, **47**. 249.)Sol. in benzonitrile. (Naumann, B. 1914, **47**. 1369.)Sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)Sol. in acetone. (Naumann, B. 1904, **37**. 4328.)

Sol. in acetone with a brown color. (Eidmann, C. C. 1899, II. 1014.)

+ $2\text{H}_2\text{O}$ (?). (Berthémot, A. ch. 1830, **44**. 385.)+ $4\text{H}_2\text{O}$. Very sol. in H_2O . (Sabatier, Bull. Soc. 1894, (3) **11**. 677.)**Cupric hydrogen bromide, CuBr_2 , HBr + $2\text{H}_2\text{O}$.**Decomp. by H_2O . (Sabatier, Bull. Soc. 1894, (3) **11**. 681.)+ $10\text{H}_2\text{O}$. (Weinland and Knöll, Z. anorg. 1905, **44**. 116.)**Cupric lithium bromide, CuBr_2 , 2LiBr + $6\text{H}_2\text{O}$.**Very hygroscopic. (Sementschenko, Z. anorg. 1899, **19**. 336.)Very hygroscopic; decomp. by H_2O . (Kurnakoff, C. C. 1899, I. 16.)**Cupric potassium bromide, CuBr_2 , KBr .**Decomp. by H_2O . (Sabatier, Bull. Soc. 1894, (3) **11**. 683.)**Cuprous bromide ammonia, Cu_2Br_2 , 2NH_3 .**

Stable when dry.

Easily sol. in HNO_3 and NH_4OH + Aq. Other mineral acids and acetic acid separate Cu_2Br_2 . (Richards, Z. anorg. 1898, **17**. 245.) Cu_2Br_2 , 6NH_3 . (Lloyd.) Cu_2Br_2 , 3NH_3 . (Lloyd, J. phys. Chem. 1908, **12**. 399.)

bromide ammonia, $\text{CuBr}_2 \cdot 2\text{NH}_3$.
in $\text{NH}_4\text{Br} + \text{Aq}$ without decomp.
ds, B. 23. 3790.)
 Br_2 , 10NH_3 . Decomp. by H_2O .
ds, Am. Ch. J. 15. 651.)
 Br_2 , 3NH_3 . Completely sol. in a little
it is decomp. by dilution. Insol. in
(Rammelsberg, Pogg. 55. 246.)
 Br_2 , $4\text{NH}_3 + \text{H}_2\text{O}$. 100 pts. H_2O dis-
.03 pts. CuBr_2 , 4NH_3 at 25° . (Pud-
Dissert.)
 Br_2 , 5NH_3 . As above. (Rammels-
 Br_2 , 6NH_3 . Sol. in small amts. of H_2O ,
comp. on dilution. (Richards.)

bromide nitric oxide, CuBr_2 , NO .
np. by H_2O . (Manchot, B. 1914,
.)

chloride, Cu_2Cl_2 .
g. Cu_2Cl_2 dissolve in 100 g. H_2O at
.55 g. at 26.5° . (Kremann and Noss,
, 33. 1206.)

ty of Cu_2Cl_2 in H_2O in an atmosphere
of hydrogen.
ility is recorded in mg-atoms per l.

Total Cu	CuCl_2 by analysis	Cl	Cu_2Cl_2 calc.	Cu_2Cl_2 by analysis
2.752	2.124	5.672	0.628	...
2.919	2.254	5.525	0.665	0.420
2.971	2.294	5.464	0.677	0.474
2.861	2.245	5.464	0.616	0.499

dländer, Z. anorg. 1902, 31. 12.)

ty of Cu_2Cl_2 in H_2O in an atmosphere
of CO_2 .
ility is recorded in mg-atoms per l.

Total Cu	CuCl_2 by analysis	Cl	Cu_2Cl_2 by analysis	Cu_2Cl_2 calc.
2.818	...	5.235	0.525	...
2.805	2.243	5.430	0.516	0.562
2.880	2.258	5.312	0.391	0.662
2.805	2.138	5.390	0.336	0.667

(Bodländer, l. c.)

conc. $\text{HCl} + \text{Aq}$; insol. in dil. HNO_3 ,
 $\text{H}_2\text{O}_2 + \text{Aq}$. Not attacked by cold conc.
and only sl. on warming. (Rosen-
12. 954.) Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; sol.
 NaCl , KCl , FeCl_3 , ZnCl_2 , MnCl_2 , etc.
mol. $\text{Na}_2\text{S}_2\text{O}_3$ in aqueous solution dis-
mol. Cu_2Cl_2 . (Winkler, J. pr. 88.
sl. in KI , I_2 , KCN , or $(\text{NH}_4)_2\text{SO}_4 +$
nault, C. R. 59. 558.)

Solubility in $\text{HCl} + \text{Aq}$ at 17° . $\frac{\text{Cu}_2\text{Cl}_2}{2} =$
 $\frac{1}{2}$ mols. CuCl_2 in mgs. in 10 ccm. solu-
tion. $\text{HCl} =$ mols. HCl in ditto.

$\frac{\text{Cu}_2\text{Cl}_2}{2}$	HCl	Sp. gr.
0.475	8.975	1.050
1.4	15.7	...
1.575	18.2	...
4.5	34.5	1.080
8.25	47.8	1.135
11.5	57.0	...

(Chatelier, calc. by Engel, A. ch. (6) 17. 377.)

Solubility of Cu_2Cl_2 in $\text{HCl} + \text{Aq}$ at 0° .

$\frac{\text{Cu}_2\text{Cl}_2}{2}$	HCl	Sp. gr.
1.5	17.5	1.049
2.9	26.0	1.065
8.25	44.75	1.132
15.5	68.5	1.261
33.0	104.0	1.345

(Engel, l. c.)

Freshly pptd. Cu_2Cl_2 is sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$.
(Lean and Whatmough, Chem. Soc. 1898,
73. 150.)

Sl. sol. in normal $\text{NH}_4\text{OH} + \text{Aq}$ only by
shaking several hours, a 0.02 normal solution
of cuprous copper being obtained. (Gaus,
Z. anorg. 1900, 25. 258.)

Insol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Siewert, Gm. K.
5. 1, 893.)

Sol. in alkyl triphosphites. (Arbusoff,
C. C. 1906, II. 750.)

Solubility in FeCl_2 , $4\text{H}_2\text{O} + \text{Aq}$ at 21.5° .

In 100 g. H_2O		Solid phase
g. FeCl_2	g. Cu_2Cl_2	
...	1.535	Cu_2Cl_2
6.015	1.33	"
11.62	1.81	"
16.30	3.11	"
26.305	7.125	"
29.35	8.06	"
33.125	9.565	"
43.75	12.44	"
54.00	17.04	"
66.40	21.60	"
73.20	23.20	$\text{Cu}_2\text{Cl}_2 + \text{FeCl}_2 \cdot 4\text{H}_2\text{O}$
71.895	21.655	$\text{FeCl}_2 \cdot 4\text{H}_2\text{O}$
69.34	11.895	"
65.10	"

(Kremann and Noss, M. 1912, 33. 1208.)

Solubility of Cu_2Cl_2 in $\text{KCl} + \text{Aq}$ at t° . Determined in an atmosphere of CO_2 .

t°	g. mol. KCl per l.	g. atoms Cu per l.
18.3	0.05	0.002411
16	0.1	0.004702
16	0.2	0.009458
19.2	1.0	0.0970
16.4	2.0	0.3840

(Bodländer and Storbeck, Z. anorg. 1902, 31. 17.)

Solubility of Cu_2Cl_2 in $\text{KCl} + \text{Aq}$ at t° . Determined in an atmosphere of CO_2 . All values recorded in millimols per litre.

t°	KCl	Cupric copper	Total copper	Cuprous copper calc.	Cl
20°	0	2.222	2.851	0.629	5.436
19	1	1.901	2.385	0.484	5.287
19	2	1.571	2.150	0.589	5.614
19	2.5	1.421	1.955	0.534	6.015
19	3	1.523	1.983	0.460	6.247
16	5	1.008	1.522	0.514	7.525
18	10	0.475	1.236	0.761	11.735
20	15	0.322	1.344	1.022	16.437
19	20	0.324	1.446	1.122	21.356
19	30	0.1308	1.761	1.630	31.911
18	50	0.1038	2.411	2.302	...
16	100	0	4.702	4.702	...
16	200	0	9.485	9.485	...
19	1000	0	97.0	97.0	...
16	2000	0	384.0	384.0	...

(Bodländer and Storbeck, Z. anorg. 1902, 31. 24.)

Solutions of 0.05–0.4 normal KCl dissolve Cu_2Cl_2 with the formation of KCuCl_2 ; those of higher concentration with the formation of K_2CuCl_3 . (Bodländer and Storbeck, Z. anorg. 1902, 31. 41.)

Solubility of $\text{Cu}_2\text{Cl}_2 + \text{KCl}$ in H_2O at 22° .

G. in 1 g. of solution		Solid Phase
Cu_2Cl_2	KCl	
0.00115	0.0387	Cu_2Cl_2
0.00405	0.0656	"
0.00861	0.0824	"
0.0137	0.0984	"
0.0219	0.1133	"
0.0390	0.1406	"
0.0484	0.1530	"
0.0675	0.1639	"
0.0719	0.1747	"
0.0863	0.1839	"
0.1043	0.2027	"
0.1084	0.2018	"
0.1021	0.2031	"
0.1204	0.2095	"
0.1332	0.2164	"

Solubility of $\text{Cu}_2\text{Cl}_2 + \text{KCl}$ in H_2O at 22° Continued

G. in 1 g. of solution		Solid Phase
Cu_2Cl_2	KCl	
0.1621	0.2330	Cu_2Cl_2
0.1723	0.2384	"
0.1907	0.2374	"
0.2148	0.2516	"
0.2145	0.2506	$\text{Cu}_2\text{Cl}_2 + \text{Cu}_2\text{Cl}_2 \cdot 4\text{K}$
0.2149	0.2549	$\text{Cu}_2\text{Cl}_2 \cdot 4\text{KCl}$
0.1548	0.2387	"
0.1473	0.2363	"
0.1399	0.2357	"
0.1439	0.2389	"
0.1451	0.2363	"
0.1155	0.2320	"
0.1139	0.2350	"
0.0953	0.2359	"
0.0735	0.2349	"
0.0555	0.2389	"
0.0453	0.2404	"
0.0366	0.2433	"
0.0314	0.2503	"
0.0285	0.2499	"
0.0265	0.2523	"
0.0220	0.2628	"
0.0193	0.2687	"
0.0176	0.2698	"
0.0193	0.2703	$\text{Cu}_2\text{Cl}_2 \cdot 4\text{KCl} + \text{KCl}$
0.0160	0.2706	KCl
0.0124	0.2668	"
0.0058	0.2632	"
0.0000	0.2568	"

(Brönsted, Z. phys. Ch. 1912, 80. 208)

Solubility in $\text{NaCl} + \text{Aq}$.
Sat. $\text{NaCl} + \text{Aq}$ dissolves 16.9 % Cu_2Cl_2 90° ; 11.9 % at 40° ; and 8.9 % at 11° .
15 % $\text{NaCl} + \text{Aq}$ dissolves 10.3 % Cu_2Cl_2 90° ; 6.0 % at 40° ; and 3.6 % at 14° .
5 % $\text{NaCl} + \text{Aq}$ dissolves 2.6 % Cu_2Cl_2 90° , and 1.1 % at 40° . (Hunt, Sill. Ar (2) 49. 154.)

Solubility in $\text{NaCl} + \text{Aq}$ at 26.5° .

In 100 g. H_2O		Solid phase
Cu_2Cl_2	NaCl	
1.55	...	Cu_2Cl_2
3.15	10.80	"
7.30	20.70	"
40.60	27.00	"
49.10	36.48	"
57.21	44.14	$\text{Cu}_2\text{Cl}_2 + \text{NaCl}$
41.40	55.95	NaCl
18.70	50.90	"

(Kremann and Noss, M. 1912, 33. 121)

solubility of Cu_2Cl_2 in $\text{CuSO}_4 + \text{Aq}$ at t° .
Values recorded in millimols per l.

	Conc. of CuSO_4	Cupric copper	Total copper	Cuprous copper calc.	Cl
		2.258	2.880	0.622	5.312
	0.49375	2.746	3.125	0.379	4.805
	0.9875	3.145	3.602	0.457	4.908
	1.4812	3.315	3.915	0.600	4.530
	1.975	4.131	4.553	0.422	4.687
	2.4687	4.349	4.786	0.437	4.287
	2.9625	4.625	5.193	0.509	4.256
	4.9375	6.546	7.276	0.730	4.329

dländer and Storbeck, Z. anorg. 1902, 31. 22.)

insol. in SbCl_3 . (Klemensiewicz, Bull. Acad. Crac. 1908, 6, 485.)

sl. sol. in liquid NH_3 . (Franklin, Am. J. 1898, 20, 827.)

insol. in alcohol.

sl. sol. in ether. (Gehlen.)

sol. in quinoline. (Beckmann and Gabel, anorg. 1906, 51. 236.)

sol. in pyridine. (Schroeder, Dissert. 01.)

insol. in phosgene. (Eidmann, Dissert. 09.)

insol. in acetone. (Naumann, B. 1904, 1. 4329.)

insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

insol. in ethyl acetate. (Alexander, Dissert. 1899.) (Naumann, B. 1904, 37. 3601.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

100 g. acetonitrile dissolve 13.33 g. Cu_2Cl_2 at 18° . (Naumann and Schier, B. 1914, 47. 49.)

sol. in benzonitrile. (Naumann, B. 1914, 7. 1369.)

sol. in hot benzonitrile and other aromatic nitriles. (Werner, Z. anorg. 1897, 15.)

Mol. weight determined in pyridine methyl and ethyl sulphides. (Werner, Z. anorg. 1897, 15. 19, 25 and 28.)

Min. Nantokite. Sol. in HCl , HNO_3 , or $\text{H}_2\text{OH} + \text{Aq}$.

Cupric chloride, CuCl_2 .

Deliquescent. 100 pts. H_2O dissolve 70.6 pts. CuCl_2 at 0° ; 100 pts. $\text{CuCl}_2 + \text{Aq}$ contain 1.4 pts. CuCl_2 . (Engel, A. ch. (6) 17. 350.)

100 pts. H_2O dissolve 76.2 pts. CuCl_2 at 11° , or 100 pts. $\text{CuCl}_2 + \text{Aq}$ sat. at 16.1° contain 43.25 pts. CuCl_2 . (Rudorff, B. 6. 484.)

100 pts. $\text{CuCl}_2 + \text{Aq}$ sat. at 17° contain 106 pts. CuCl_2 ; at 31.5° , contain 44.7 pts. CuCl_2 .

Coefficient of solubility = $41.4 + 105t$. (Reicher and Deventer, Z. phys. 1. 5. 560.)

Sat. $\text{CuCl}_2 + \text{Aq}$ contains at:

-20°	-5°	$+12^\circ$	17°	32°
37	38.8	39.3	41.7	43.2% CuCl_2
39	55	68	73	91
44.0	46.5	47.9	48.6	51.0% CuCl_2

(Étard, A. ch. 1894, (7) 2. 536.)

Sp. gr. of $\text{CuCl}_2 + \text{Aq}$ at 17.5° .

% CuCl_2	Sp. gr.	% CuCl_2	Sp. gr.
5	1.0455	25	1.2918
10	1.0920	30	1.3618
15	1.1565	35	1.4447
20	1.2223	40	1.5284

(Franz, J. pr. (2) 5. 274.)

Sp. gr. of $\text{CuCl}_2 + \text{Aq}$ at 22.9° , containing in 1000 g. H_2O , g. $\text{CuCl}_2 + 2\text{H}_2\text{O}$.

85.5 (= $\frac{1}{2}$ mol.) 171 255.5 g. $\text{CuCl}_2 + 2\text{H}_2\text{O}$,
1.057 1.108 1.154

342 427.5 513 g. $\text{CuCl}_2 + 2\text{H}_2\text{O}$,
1.197 1.238 1.275

598.5 684 g. $\text{CuCl}_2 + 2\text{H}_2\text{O}$,
1.309 1.341

769.5 855 g. $\text{CuCl}_2 + 2\text{H}_2\text{O}$,
1.371 1.399

940.5 1.026 g. $\text{CuCl}_2 + 2\text{H}_2\text{O}$.
1.425 1.449

Containing CuCl_2 (anhydrous).

67.5 (= $\frac{1}{2}$ mol.) 135 202.5 270 g. CuCl_2 ,
1.059 1.114 1.165 1.213

337.5 405 472.5 g. CuCl_2 .
1.257 1.299 1.30

540 607.5 675 g. CuCl_2 .
1.379 1.416 1.453

(Gerlach, Z. anal. 28. 468.)

Sp. gr. of $\text{CuCl}_2 + \text{Aq}$ at 0° . S = pts. CuCl_2 in 100 pts. solution; S_1 = mols. CuCl_2 in 100 mols. of solution.

S	S_1	Sp. gr.
39.4170	8.00	1.4797
35.3839	6.82	1.4173
30.9255	5.65	1.3529
26.1129	4.51	1.2881
20.6697	3.36	1.2204
14.5820	2.23	1.1494
8.0732	1.16	1.0796

(Charpy, A. ch. (6) 29. 25.)

Tables for 7° , 30.5° , 49.2° , and 65° are also given by Charpy.

Sp. gr. of $\text{CuCl}_2 + \text{Aq}$ at room temp., containing:

12.006 21.349 33.027% CuCl_2 .
1.1037 1.2154 1.3312

(Wagner, W. Ann. 1883, 18. 273.)

Sp. gr. at 20° of $\text{CuCl}_2 + \text{Aq}$ containing M. mols. CuCl_2 per liter.

M.	0.01	0.05	0.075
Sp. gr.	1.001208	1.00637	1.009264
M.	0.10	0.20	0.50
Sp. gr.	1.012614	1.030991	1.051479
M.	0.75		1.0
Sp. gr.	1.090912		1.120249
M.	1.5		2.0
Sp. gr.	1.177618		1.234551

(Jones and Pearce, Am. Ch. J. 1907, 33, 717.)

Sp. gr. of $\text{CuCl}_2 + \text{Aq}$ at 25°.

Concentration of $\text{CuCl}_2 + \text{Aq}$.	Sp. gr.
1—normal	1.0624
$\frac{1}{2}$ —	1.0313
$\frac{1}{4}$ —	1.0158
$\frac{1}{8}$ —	1.0077

(Wagner, Z. phys. Ch. 1890, 5, 38.)

Much less sol. in $\text{HCl} + \text{Aq}$ than in H_2O . 1 l. $\text{HCl} + \text{Aq}$ containing 45 pts. HCl to 100 pts. H_2O dissolves only 290 g. CuCl_2 at 12°, whereas 1 l. H_2O at 12° dissolves 630 g. CuCl_2 . (Ditte, C. R. 1881, 92, 353.)

Solubility in $\text{HCl} + \text{Aq}$ at 0°. $\frac{\text{CuCl}_2}{2}$ =

$\frac{1}{2}$ mols. in milligrammes in 10 cc. solution. HCl = mols. HCl in ditto; H_2O = g. H_2O .

$\frac{\text{CuCl}_2}{2}$	HCl	Sum of equiv.	Sp. gr.	H_2O
91.75	■	91.75	1.490	8.73
86.8	4.5	91.3	1.475	8.74
83.2	7.8	91	1.458	
79.35	10.5	89.85	1.435	8.64
68.4	20.25	88.65	1.389	8.56
50.0	37.5	87.5	1.319	8.47
22.8	70.25	93.05	1.231	8.21
23.5	102.5	126	1.288	7.56
26.7	128	154.7	1.323	6.77

(Engel, A. ch. (6) 17, 351.)

Not decomp. by cold H_2SO_4 .

Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. Very sol. in conc. $\text{NaCl} + \text{Aq}$ (Boussingault.)

Solubility of CuCl_2 in $\text{NH}_4\text{Cl} + \text{Aq}$ at 30°.

$\frac{\text{CuCl}_2}{2}$	$\frac{\text{NH}_4\text{Cl}}{2}$	Solid phase
29.5	0	NH_4Cl
28.6	1.9	$\text{NH}_4\text{Cl} + \text{CuCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O}$
12.1	15.6	$\text{CuCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O}$
2.03	43.2	$\text{CuCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O} + \text{CuCl}_2, 2\text{H}_2\text{O}$
0	43.95	$\text{CuCl}_2, 2\text{H}_2\text{O}$

(Schreinmakers, Z. phys. Ch. 1909, 66, 688.)

See also $\text{NH}_4\text{Cl} + \text{CuCl}_2$ under ammonium chloride.

Solubility of $\text{CuCl}_2 + \text{HgCl}_2$ in H_2O at 37°

% HgCl_2	% CuCl_2	Solid Phase
0	44.47	$\text{CuCl}_2, 2\text{H}_2\text{O}$
21.03	33.50	"
37.30	26.07	"
44.47	23.31	"
50.47	21.50	$\text{CuCl}_2 + \text{HgCl}_2$
52.44	19.40	HgCl_2
52.54	18.46	"
52.81	18.06	"
51.03	14.73	"
49.50	5.94	"
23.87	2.64	"
8.51	8.51	"

(Schreinmakers and Thonius, Proc. K. Ak. Wet. 1912, 15, 472.)

Solubility of $\text{CuCl}_2 + \text{KCl}$ in H_2O at t°

t°	Cl per g. of solution		Mols. per 100 mols. H_2O		
	As CuCl_2	As KCl	CuCl_2	KCl	
39.4	0.120	0.107	5.50	9.93	$\text{CuCl}_2, 2\text{KCl}, 2\text{H}_2\text{O}$
49.9	0.129	0.115	6.39	11.4	"
50.4	0.142	0.125	7.71	13.6	"
79.1	0.168	0.142	11.1	18.8	"
90.5	0.189	0.154	14.9	24.4	"
93.7	0.194	0.155	16.2	26.0	$\text{CuCl}_2, \text{KCl} + \text{D}$
98.8	0.197	0.162	17.5	28.7	"
0	0.214	0.021	9.84	1.94	$\text{CuCl}_2, 2\text{KCl}, 2\text{H}_2\text{O}$
					$\text{CuCl}_2, 3\text{H}_2\text{O}$
39.6	0.232	0.049	12.9	5.44	"
50.1	0.233	0.059	13.7	6.90	"
52.9	0.241	0.062	14.8	7.63	"
60.2	0.246	0.066	15.8	8.49	$\text{CuCl}_2, \text{KCl} + \text{Cu}$
					$2\text{H}_2\text{O}$
72.6	0.255	0.063	16.8	8.35	"
94.2			14.9	11.6	$\text{CuCl}_2, 2\text{KCl}, 2\text{H}_2\text{O}$
					$\text{CuCl}_2, \text{KCl}$
72.5			14.8	1.50	$\text{CuCl}_2, \text{KCl}$

(Meyerhoffer, Z. phys. Ch. 1890, 5, K)

100 g. H_2O dissolve 72.6 g. $\text{CuCl}_2 + 16 \text{ NaCl}$. (Rudorff, B. 6, 684.)

Solubility of $\text{CuCl}_2 + \text{NaCl}$ in H_2O at 3°

% NaCl	% CuCl_2	Solid Phase
0	43.95	$\text{CuCl}_2, 2\text{H}_2\text{O}$
3.10	41.14	"
4.28	41.06	"
6.41	39.40	"
10.25	36.86	$\text{CuCl}_2 + \text{NaCl}$
12.02	32.38	"
12.25	32.40	"
13.54	28.64	"
15.40	23.72	"
18.44	16.98	"
20.61	11.03	"
26.47	0	"

(Schreinmakers and de Baat, Z. phys. 1909, 65, 586.)

Solubility of CuCl₂ in CuSO₄+Aq at 30°.

Composition of the solution		Solid phase.
Pt. Cl ₂	% by wt. CuSO ₄	
8	20.32	CuSO ₄ , 5H ₂ O
8	13.62	
8	8.93	
7	4.77	
8	3.21	CuSO ₄ , 5H ₂ O + CuCl ₂ , 2H ₂ O
7	2.89	
7	2.90	CuCl ₂ , 2H ₂ O
5	1.14	
5	0	

reinemakers, Z. phys. Ch. 1909, 69. 561.)

sol. in liquid NH₃. (Gore, Am. Ch. J. 8, 20. 827.)

l. sol. in liquid HF. (Franklin, Z. anorg. 5, 46. 2.)

sol. in alcohol and ether.

sol. in 1 pt. strong alcohol.

100 pts. absolute methyl alcohol dissolve 68 CuCl₂ at 15.5°; 100 pts. absolute ethyl alcohol dissolve 53 pts. CuCl₂ at 15.5°. (de laun, Z. phys. Ch. 10. 783.)

See also under CuCl₂+2H₂O.

Solubility of CuCl₂+KCl in absolute alcohol at 25°.

% CuCl ₂	% KCl	Solid Phase
.27	0.28	KCl + KCl, CuCl ₂
.51	0.28	
.15	KCl, CuCl ₂
.25	
.16	KCl, CuCl ₂ + CuCl ₂ , C ₂ H ₅ OH
.17	0.21	
.45	0.21	"
.29	0.21	
.97	0.00	CuCl ₂ , C ₂ H ₅ OH

Foot and Walden, J. Am. Chem. Soc. 1911, 33. 1032.)

Solubility of CuCl₂+KCl in acetone at 25°.

% CuCl ₂	% KCl	Solid Phase
.34	0.38	KCl + KCl, CuCl ₂
.18	
.50	"
.16	
.19	0.27	KCl, CuCl ₂ + CuCl ₂ , C ₂ H ₆ O
.12	

Foot and Walden, J. Am. Chem. Soc. 1911, 33. 1032.)

100 g. of sat. solution of CuCl₂ in ethyl alcohol contains 33.97 g. CuCl₂. (Foot and Walden, J. Am. Chem. Soc. 1911, 33. 1032.)

Easily sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)
Insol. in benzene.

Solubility in organic solvents.

Solvent	t°	Sat. solution contains % CuCl ₂
Methyl alcohol	22	36.8
	40	37.5
	50	37.1
	60	37.5
Abs. ethyl alcohol	0	32.0
	19	35.7
	20	35.9
	38	38.5
	50	41.7
Propyl alcohol	-15	26.8
	19	30.9
	37	30.7
	57	30.3
	62	30.5
Allyl alcohol	-20	23.4
	-4	23.6
	27	22.9
	32	23.3
N-butyl alcohol	0	15.2
	23	15.8
	37	15.7
	55	16.1
	84	16.2
Ethyl formate	92	16.7
	-20	10.2
	+24	9.4
	37	7.4
Ethyl acetate	50	7.2
	+20	3.0
	40	2.5
Acetone	72	1.3
	-20	18.4
Isopropyl alcohol	+8	18.8
	32	11.0
	70	28.3
	84	28.7

(Étard, A. ch. 1894, (7) 2. 565.)

See also under CuCl₂+2H₂O.

1 g. CuCl₂ is sol. in 181 g. methyl acetate at 18°. Sp. gr. 18°/4° of the sat. solution = 0.939. (Naumann, B. 1909, 42. 3793.)

1 g. CuCl₂ is sol. in 249 g. ethyl acetate at 18°. Sp. gr. of sat. solution 18°/4° = 0.9055. (Naumann, B. 1904, 37. 3603.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

1 pt. sol. in 249.3 pts. ethyl acetate at 10°. (Alexander, Dissert. 1899.)
Solubility in acetone.
34.7 g. acetone dissolve 1 g. CuCl₂ at 18°. Sp. gr. of sat. solution 18°/4° = 0.8154. (Naumann, B. 1904, 37. 4329.)
1.40 pts. are sol. in 100 pts. acetone at 56°. (Laszczynski, B. 1894, 27. 2287.)
Sol. in acetone and methylal. 1 gram. dissolves in 34.08 grams of acetone at 18°. (Eidmann, C. C. 1899, II, 1014.)
100 pts. absolute ether dissolve 0.043 g. CuCl₂. (Bödtker, Z. phys. Ch. 1897, 22. 511.)
Mol. weight determined in pyridine and methyl sulphide. (Werner, Z. anorg. 1897, 15. 20 and 25.)
100 g. sat. solution in acetonitrile contains 1.57 g. CuCl₂. (Naumann and Schier, B. 1914, 47. 249.)
Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)
Sol. in boiling dipropylamine. (Werner, Z. anorg. 1897, 15. 34.)
Sol. in urethane. (Ley, Z. phys. Ch. 1897, 22. 81); (Castoro Z. anorg. 1899, 20, 61.)
+H₂O. (Ditte, A. ch. (5) 22. 551.)
Sol. in H₂O with slight decomp. (Sabatier, Bull. Soc. 1895, (3) 13. 601.)
+2H₂O. Deliquescent. 100 g. H₂O dissolve 121.4 g. CuCl₂+2H₂O at 16.1°. (Rüddorff.)
Aq. sol. at 35° contains 9.689 Mol. % CuCl₂.
" 15° " 8.934 " % "
(Schreinemakers, C. C. 1911, II. 349.)

CuCl₂+2H₂O+Aq. sat. at 30° contains 43.95% CuCl₂. (Meerburg, C. C. 1904, II. 1362.)
43.95 g. anhydrous CuCl₂ are dissolved in 100 g. CuCl₂+Aq at 30°. (Schreinemakers, Arch. neer. Sc. 1910 [2], 15. 117.)
44.47% by weight anhydrous CuCl₂ is dissolved in H₂O at 35°. (Schreinemakers and Thonus, Verh. k. Akad. Wet. Amst. 1912, 21. 333.)

Solubility of CuCl₂+2H₂O in ethyl alcohol +Aq at 11.5°.

Percent of ethyl alcohol by volume	Grams CuCl ₂ dissolved in 5 cc.
99.3	1.175
98.3	1.116
96.3	1.097
95.3	1.070
94.3	1.059

Anhydrous salt dissolves readily in absolute ethyl alcohol; CuCl₂+2H₂O is precipitated by H₂O.

Solubility of CuCl₂+2H₂O in ethyl alcohol Aq at 11.5° C. under addition of creasing amounts of CuCl₂.
P = Percent of ethyl alcohol by volume
G = Grams of CuCl₂ added.
C₀ = Grams of CuCl₂ in 5 cc. of the solution
C_w = Grams of water in 5 cc. of the solution calculated from
(1) the water content of the alcohol.
(2) the water of crystallization which gone into solution.
(3) the water held mechanically in C +2H₂O.

P	G	C _w	C ₀
89.3	0.000	0.794	1.13
90.3	"	0.744	1.12
91.3	"	0.695	1.10
92.3	"	0.648	1.09
94.3	"	0.561	1.09
95.3	"	0.517	1.09
96.3	"	0.478	1.11
97.3	"	0.440	1.14
98.3	"	0.396	1.19
99.3	"	0.369	1.20
"	0.223	0.330	1.29
"	0.444	0.290	1.39
"	0.665	0.270	1.50
"	0.887	0.247	1.63
"	1.106	0.223	1.77
"	1.324	0.205	1.92
"	1.540	0.191	2.08
"	1.739	0.179	2.23
"	1.957	0.164	2.40

(Bödtker, Z. phys. Ch. 1897, 22. 506-5)

100 pts. absolute ether dissolve 0.06 CuCl₂+2H₂O. (Bödtker, Z. phys. Ch. 1 22. 511.)
0.11 pts. are sol. in 100 pts. ether at
0.11 " " " " 100 " " "
8.86 " " " " 100 " acetone "
8.92 " " " " 100 " " " 1
(Laszczynski, B. 1894, 27. 2286 and 2

Solubility in organic solvents at t°.

Solvent	t°	P Percent ethyl alcohol by volume
Pure methyl alcohol	20°	"
"	18.9	"
Ethyl alcohol (95°)	20.3	1
"	19.6	1
Pure acetone	22.1	4
"	20.0	4
90 pts. ethyl alcohol (98°) +10 pts. H ₂ O	21.8	"
"	23.0	"

Solubility in organic solvents at t°.—Cont.

Solvent	t°	Pts. of solvent required to dissolve 1 pt. CuCl ₂ + 2H ₂ O at t°
1 pta. abs. alcohol + 20 pta. H ₂ O	28.1	6.0
"	20.7	6.2
1 pta. acetone + 20 pta. H ₂ O	23.1	5.3
"	21.8	5.6
1 pta. acetone + 20 pta. methyl alcohol	23.1	12.0
"	24.0	11.6
0 pta. methyl alcohol + 90 pta. ethyl alcohol (98°)	24.2	5.4
"	25.0	5.1
10 pta. methyl alcohol + 80 pta. ord. ether	24.1	15.1
"	22.4	15.7
80 pta. abs. alcohol + 20 pta. ord. ether	24.1	8.8
"	25.0	8.5
Comm. methyl alcohol	23.9	5.4
"	23.0	5.6
5 pta. pyridine + 15 pta. H ₂ O	24.4	63.4
"	23.6	63.7
6 pta. pyridine + 40 pta. H ₂ O	27.3	26.7
"	28.0	26.2
15 pta. α picoline + 25 pta. H ₂ O	26.1	51.6
"	25.1	52.3
70 pta. α picoline + 30 pta. H ₂ O	26.1	47.3

(de Coninck, C. R. 1900, 131. 59.)

Solubility in organic solvents.

Sol. in propyl alcohol, hot glycol, hot glycerine, hot paraldehyde, hot crys. acetic acid, pure acetone, 30% methylamine + Aq, pure pyridine, pure α-picoline, acetonitrile; sl. sol. in isobutyl and amyl alcohols, crystallisable formic acid, ethyl acetate; insol. in cold glycol, cold glycerine, cold paraldehyde, benzaldehyde, cold crystallisable acetic acid, ord. ether, abs. ether, hot and cold CS₂, cold aniline, cold orthotoluidine, methylene chloride, ethyl iodide, propyl iodide, ethylene bromide, benzene, toluene, xylene, ligroin, nitrobenzene, cold piperidine and essence of termenthine. (de Coninck, C. R. 1900, 131. 59.)

+ 5H₂O.

Aq. sol. at 35° contains 2.921 Mol. % CuCl₂.
15° " 1.11

(Schreinemakers, C. C. 1911, II. 349.)

Cuprous hydrogen chloride, CuCl, HCl.

Sol. in H₂O. (Neumann, M. 1894, 15. 493.)Cupric hydrogen chloride, CuCl₂, HCl + 3H₂ODecomp. by H₂O. Sol. in HCl + Aq below 0° (Engel, C. R. 106. 273.)CuCl₂, 2HCl. Deliquescent. Very sol. in H₂O. (Alexander, Dissert. 1899.)+ 5H₂O. Properties as above. (Sabbatier, C. R. 106. 1724.)CuCl₂, 3HCl. Sol. in H₂O. (Neumann M. 1894, 15. 493.)Cupric gold (auric) chloride, CuCl₂, 2AuCl₃ + 6H₂O.10% is sol. in H₂O at 18°. (Mylius, Z. anorg. 1911, 70. 210.)Cupric lithium chloride, CuCl₂, LiCl + 2½H₂O.Decomp. on air. Decomp. by dissolving in H₂O. Sol. in conc. LiCl + Aq without decomp. Decomp. by alcohol. (Chassevant, A. ch. (6) 30. 33.)+ 2H₂O. (Meyerhoffer, W. A. B. 100, 2b. 621.)

Cupric mercuric chloride.

Easily sol. in H₂O. (v. Bonadorff.)Cupric mercuric potassium chloride, CuCl₂, 3HgCl₂, 6KCl + 2H₂O.Deliquescent in moist air. Sol. in boiling H₂O without decomp., and recrystallises if cooled slowly. Insol. in absolute alcohol. (v. Bonadorff, Pogg. 33. 81.)Cuprous nitrosyl chloride, Cu₂Cl₂, 2NOCl.Very deliquescent and sol. in H₂O with immediate decomp. (Sudborough, Chem. Soc. 59. 658.)Cuprous potassium chloride, Cu₂Cl₂, 4KCl.Sol. in H₂O. (Mitscherlich, A. ch. 73. 384.)For solubility data, see Cu₂Cl₂ + KCl under cuprous chloride.

Cupric potassium chloride.

CuCl₂, KCl. (Meyerhoffer, Z. phys. Ch. 3. 336.)Sol. in H₂O; only sl. sol. in conc. HCl + Aq. (Gröger, Z. anorg. 1899, 19. 330.)CuCl₂, 2KCl + 2H₂O. Sol. in H₂O and alcohol. (Berzelius, Pogg. 13. 458.)

The composition of the hydrates formed by this salt at different dilutions is calculated from determinations of the lowering of the fr. pt. produced by the salt and of the conductivity and sp. gr. of its aqueous solutions. (Jones, Am. Ch. J. 1905, 34. 322.)

For solubility data, see CuCl₂ + KCl under cupric chloride.Cupric rubidium chloride, CuCl₂, 2RbCl.Easily sol. in H₂O and HCl + Aq. (Godefroy, B. 8. 9.)+ 2H₂O. Sol. in H₂O. (Wyrouboff, J. B. 1887. 538.)

Cuprous sodium chloride.

Very sol. in H₂O.

Cupric sodium chloride.

Easily sol. in conc. NaCl + Aq. Sol. in alcohol of 0.837 sp. gr.

No double salt exists. (Schreinemakers and de Baat, Z. phys. Ch. 1909, **65**. 586.)

Cupric thallic chloride, $\text{CuCl}_2, 2\text{TlCl}_2$.

Sol. in H_2O . (Willm, A. ch. (4) **5**. 55.)
+ $6\text{H}_2\text{O}$. Can be cryst. from H_2O . (Gewecke, A. 1909, **366**. 225.)

Cuprous chloride ammonia, $\text{Cu}_2\text{Cl}_2, \text{NH}_3$.
(Lloyd, J. phys. Chem. 1908, **12**. 399.)

$\text{Cu}_2\text{Cl}_2, 2\text{NH}_3$. Decomp. by H_2O or acids, not by alcohol. (Ritthausen, J. pr. **59**. 369.)

$\text{Cu}_2\text{Cl}_2, 3\text{NH}_3$. (Lloyd, J. phys. Chem. 1908, **12**. 399.)

$\text{Cu}_2\text{Cl}_2, 6\text{NH}_3$. (Lloyd, J. phys. Chem. 1908, **12**. 399.)

Cupric chloride ammonia, $\text{CuCl}_2, 2\text{NH}_3$.

Decomp. by H_2O . (Kane, A. ch. **72**. 273.)
 $\text{CuCl}_2, 4\text{NH}_3$. Sol. in H_2O . (Bouzat, C. R. 1902, **135**. 294.)

+ H_2O (*Cuprammonium chloride*). Sol. in H_2O and hot $\text{NH}_4\text{OH} + \text{Aq}$.

+ $2\text{H}_2\text{O}$. Sol. in small amt. of H_2O . $\text{Cu}(\text{OH})_2$ is pptd. by dilution. (Bouzat, A. ch. 1903, (7) **29**. 350.)

$\text{CuCl}_2, 5\text{NH}_3$. (Bouzat, A. ch. 1903, (7) **29**. 350.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O . On dilution $\text{Cu}(\text{OH})_2$ is pptd. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; solubility decreases as NH_3 concentration increases. (Bouzat, A. ch. 1903, (7) **29**. 350.)

$\text{CuCl}_2, 6\text{NH}_3$. Completely sol. in H_2O . (Rose, Pogg. **20**. 55.)

Sol. in H_2O but decomp. by great dilution with pptn. of $\text{Cu}(\text{OH})_2$.

Insol. in liquid NH_3 . (Bouzat, A. ch. 1903, (7) **29**. 350.)

Cuprocupric chloride ammonia, $\text{Cu}_2\text{Cl}_2, \text{CuCl}_2, 4\text{NH}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O or alcohol. Abundantly sol. in $\text{NH}_4\text{Cl} + \text{Aq}$, but with partial decomposition. (Ritthausen.)

Cupric chloride ammonia platinous chloride, $\text{CuCl}_2, 4\text{NH}_3, \text{PtCl}_2$.

See Platodiamine cupric chloride.

Cuprous chloride carbon monoxide, $\text{Cu}_2\text{Cl}_2, 2\text{CO} + 4\text{H}_2\text{O}$.

Very sol. in HCl (sp. gr. 1.19) with evolution of CO. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Manchot and Friend, A. 1908, **359**. 110.)

$2\text{Cu}_2\text{Cl}_2, \text{CO} + 2\text{H}_2\text{O}$. Insol. in H_2O . (Berthelot, A. ch. 1856, (3) **46**. 488.)

$4\text{Cu}_2\text{Cl}_2, 3\text{CO} + 7\text{H}_2\text{O}$. Insol. in H_2O , but decomp. therewith very quickly. Sol. in $\text{Cu}_2\text{Cl}_2 + \text{HCl}$.

Cupric chloride hydrazine, $\text{CuCl}_2, 2\text{N}_2\text{H}_4$.

Easily decomp. (Hofmann and Marburg, A. 1899, **305**. 222.)

Cuprous chloride mercuric sulphide, $\text{Cu}_2\text{Cl}_2, 2\text{HgS}$.

Insol. in H_2O ; sol. in conc. hot $\text{HCl} + \text{Aq}$; not decomp. by boiling dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, but decomp. by conc. H_2SO_4 . (Heumann, B. **1**. 1390.)

Cuprous fluoride, Cu_2F_2 .

Insol. in H_2O or HF. Sol. in conc. $\text{HCl} + \text{Aq}$, from which it is precipitated by H_2O . Insol. in alcohol. (Berzelius, Pogg. **1**. 28.)

Decomp. by H_2O into sol. CuF_2 . Sol. in boiling $\text{HCl} + \text{Aq}$ and in $\text{HNO}_3 + \text{Aq}$. Only sl. attacked by warm H_2SO_4 . (Poulenc, C. R. **116**. 1447.)

Cupric fluoride, CuF_2 .

Easily takes up H_2O to form $\text{CuF}_2 + 2\text{H}_2\text{O}$. Sol. in HCl , HNO_3 , or $\text{HF} + \text{Aq}$. (Poulenc, C. R. **116**. 1448.)

Solubility at 25° in $\text{HF} + \text{Aq}$.

Normality of $\text{HF} + \text{Aq}$.	g. atoms Cu in 1000 c. c. of solution
0.12	0.0307
0.28	0.1164
0.57	0.2494
1.08	0.388
2.28	0.463

Solubility is decreased by presence of KF. (Jaeger, Z. anorg. 1901, **27**. 29.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 827.)

Insol. in methyl acetate (Naumann, B. 1909, **42**. 3790); ethyl acetate. (Naumann, B. 1910, **43**. 314.)

Insol. in acetone. (Eidmann, C. C. **1898**, II. 1014; Naumann, B. 1904, **37**. 4329.)

+ $2\text{H}_2\text{O}$. Sl. sol. in cold, decomp. by hot H_2O . (Berzelius.)

Cupric hydrogen fluoride, $\text{CuF}_2, 5\text{HF} + 5\text{H}_2\text{O}$.

Deliquescent.

Easily sol. in H_2O and dil. acids.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ with decomp. (Böhm, Z. anorg. 1905, **43**. 329.)

Cupric potassium fluoride, $\text{CuF}_2, 2\text{KF}$.

Easily sol. in H_2O .

CuF_2, KF . Very sl. sol. in H_2O ; sl. sol. in dil. acids. (Helmholt, Z. anorg. **3**. 115.)

Cupric rubidium fluoride, CuF_2, RbF .

As the K salt. (Helmholt.)

Cupric silicon fluoride.

See Fluosilicate, cupric.

cupric fluoride.
cupric stannate, copper.
cupric tantalum fluoride.
cupric tantalate, copper.
cupric antimony fluoride.
cupric titanate, copper.
cupric tungstic fluoride.
cupric tungstate, copper.
cupric cerium fluoride.
cupric zirconate, copper.
cuprous fluoride ammonia, CuF_2 , 4NH_3 +
sol. rapidly in the air.
sol. in H_2O .
sol. by boiling with H_2O with evolu-
 H_2 .
sol. in dil. acids. (Böhm, Z. anorg.
333.)

cuprous hydride, CuH .
insol. in H_2O . Sol. in HCl + Aq. (Wurtz,
102.)
sol. in warm conc. HCl with decomp.
Am. Ch. J. 1895, 17. 187.)

cuprous hydride, CuH_2 .
sol. in HCl with decomp. (Bartlett, Am.
95, 17. 187.)

cuprous hydrosulphide, 7CuS , H_2S .
sol. in H_2S . (Linder and Picton, Chem. Soc. 1892, 61.

H_2S . (Linder and Picton.)
 H_2S . (Linder and Picton.)

cuprous hydroxide, Cu_2O , $x\text{H}_2\text{O}$.
insol. in acids as cupric salt. Insol. in NaOH ,
+ Aq.
sol. in NH_4OH , and $(\text{NH}_4)_2\text{CO}_3$ + Aq; sol.
in O_2 + Aq.

cupric hydroxide, CuOH , $3\text{Cu}(\text{OH})_2$,
 I_2O .
insol. in acids. (Francke, Dissert. 1907.)

cuprous hydroxide, 3CuO , H_2O .
insol. in H_2O or dil. alkalies. Easily sol.
in NH_4Cl + Aq. (Rose.)
more difficultly sol. than CuO_2H_2 in
aq. (Chodnew, J. pr. 28. 220.)
composition is 6CuO , H_2O .
is Cupric oxide.

I_2 . Insol. in H_2O , but decomp. into
 I_2O by being boiled therewith.
easily sol. in acids.
sol. in NH_4OH , and NH_4 salts + Aq.

Solubility in NH_4OH + Aq at 25° .

NH_3 conc. m.	g. Cu per l.	equiv. CuO_2H_2 per l.
2.63	3.05	0.096
2.00	2.12	0.067
1.32	1.08	0.034
2.540	6.26	0.197
1.965	6.28	0.166
1.280	4.13	0.129
0.973	3.36	0.106
0.870	3.08	0.097
0.540	2.36	0.074
0.391	2.04	0.064
3.176	8.06	0.253
2.070	5.72	0.180
1.272	4.75	0.149
0.451	2.54	0.080
0.320	2.13	0.067

The non-agreement of the results is due to the presence of different modifications of CuO_2H_2 .

(Bonsdorff, Z. anorg. 1904, 41. 182.)

Solubility in NH_4OH + Aq at 18° .

NH_3 mols per l.	Cu g. atoms per l.
0.20	0.00054
0.50	0.0033
1.0	0.0109
1.5	0.0204
2.0	0.0314
2.5	0.0442
3.0	0.0548
4.0	0.0784
5.0	0.1041
6.0	0.1254
8.0	0.1599
9.96	0.1787

(Dawson, Z. phys. Ch. 1909, 69. 111.)

Sol. in cold NaOH , or KOH + Aq (Proust); but CuO is pptd. on boiling (Berthollet); is not pptd. (Chodnew, J. pr. 28. 220.)

Insol. in NaOH or KOH + Aq unless they contain organic matter (Berzelius). This is contradicted by Volcker (A. 59. 34).

Entirely sol. in conc. KOH + Aq, but solution is decomp. by heating. (Freymy, A. ch. (3) 12. 510.)

Sol. in NaOH + Aq (70% NaOH). (Löw, Z. anal. 9. 463.)

The solubility in NaOH of CuO_2H_2 , prepared either from CuSO_4 or $\text{Cu}(\text{NO}_3)_2$, decreases with decrease in concentration of the base. The solubility of CuO_2H_2 in NaOH is only very slightly affected by the addition of sodium or potassium carbonate. (Fischer, Z. anorg. 1904, 40. 41.)

Solubility of crystalline CuO_2H_2 in ammoniacal $\text{Ba}(\text{OH})_2$ and NaOH solutions at 18° .

Solvent contains per litre	Cu concentration g atoms per l.
1 mol. NH_3 + 0 mol. $\text{Ba}(\text{OH})_2$	0.01090
" + 0.0025 "	0.00907
" + 0.005 "	0.00801
" + 0.01 "	0.00633
" + 0.02 "	0.00526
2 mols. NH_3 + 0 mol. $\text{Ba}(\text{OH})_2$	0.0314
" + 0.01 "	0.0277
4 mols. NH_3 + 0 mol. $\text{Ba}(\text{OH})_2$	0.0784
" + 0.01 "	0.0747
1 mol. NH_3 + 0 mol. NaOH	0.0109
" + 0.01 "	0.00766
" + 0.02 "	0.00655
" + 0.03 "	0.00531
" + 0.05 "	0.00456
" + 0.10 "	0.00410

(Dawson, Chem. Soc. 1909, 95. 377.)

Solubility of crystalline CuO_2H_2 in ammoniacal salt solutions at 18° .

Solvent contains per litre	Conc. of dissolved Cu g atoms per l.
1 mol. NH_3	0.0109
0.05 mol. NH_3 + 0.01 mol. $(\text{NH}_4)_2\text{SO}_4$	0.00129
" + 0.025 "	0.00511
0.1 mol. NH_3 + 0.01 mol. $(\text{NH}_4)_2\text{SO}_4$	0.00326
" + 0.025 "	0.0108
" + 0.05 "	0.0233
0.2 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	0.00054
" + 0.01 "	0.00649
" + 0.025 "	0.0175
" + 0.05 "	0.0384
" + 0.10 "	0.0690
0.3 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	0.0033
" + 0.01 "	0.0127
" + 0.025 "	0.0284
" + 0.05 "	0.0536
" + 0.10 "	0.1013
" + 0.20 "	0.1844
1.0 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	0.0109
" + 0.01 "	0.0210
" + 0.025 "	0.0386
" + 0.05 "	0.0660
" + 0.10 "	0.1185
" + 0.20 "	0.2275
" + 0.40 "	0.4135
2 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	0.0314
" + 0.01 "	0.0462
" + 0.025 "	0.0605
" + 0.05 "	0.0886
" + 0.10 "	0.1468
" + 0.20 "	0.2591
" + 0.40 "	0.4718
4 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	0.0548
" + 0.01 "	0.0672
" + 0.025 "	0.0847
" + 0.05 "	0.1156

Solubility of crystalline CuO_2H_2 in ammoniacal salt solutions at 18° —Contd.

Solvent contains per litre	
3 mol. NH_3 + 0.10 mol. $(\text{NH}_4)_2\text{SO}_4$	
" + 0.20 "	
" + 0.40 "	
4 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	
" + 0.01 "	
" + 0.025 "	
" + 0.05 "	
" + 0.10 "	
" + 0.20 "	
" + 0.40 "	
5 mol. NH_3 + 0.00 mol. $(\text{NH}_4)_2\text{SO}_4$	
" + 0.01 "	
" + 0.025 "	
" + 0.05 "	
" + 0.10 "	
" + 0.20 "	
" + 0.40 "	
1 mol. NH_3 + 0.00 mol. Na_2SO_4	
" + 0.025 "	
" + 0.10 "	
" + 0.20 "	
" + 0.40 "	
4 mol. NH_3 + 0.10 mol. Na_2SO_4	
" + 0.20 "	

(Dawson, Chem. Soc. 1909, 95.

Sl. sol. in alkali carbonates + Aq. KHCO_3 and NaHCO_3 . (Berselius. Sol. in cold $\text{Na}_2\text{S}_2\text{O}_3$ + Aq, but warming. (Field, Chem. Soc. (2) 1 Partially sol when freshly pptd + Aq. (Rodgers, 1834.)

Sol. in $(\text{NH}_4)_2\text{S}_2\text{O}_8$ + Aq. (Mores Ztg. 1901, 16. 383.)

Sol. in MSCN + Aq., more diff. in NH_4SCN + Aq. than ZnO_2H_2 . mann, Z. anorg. 1908, 58. 269.)

Very sol. in hydroxylamine. and Cohen, J. pr. 1905, (2), 72. 14.

Insol. in acetone. (Eidmann, C. H. 1014.)

Sol. in large amt. in $\text{NaC}_2\text{H}_3\text{O}_2$ + Aq. (1844.)

Not pptd. in presence of Na citrate.)

Insol. in cane sugar + Aq, unless an alkaline earth is present. (Peschier

Recently pptd. CuO_2H_2 is easily sol. in sugar with NaOH , KOH , or CaO less sol. in presence of SrO_2H_2 or (Becquerel)

Not pptd. by KOH + Aq. in solution containing tartaric acid, cane sugar, or other non-volatile organic substance

Sol. in Ca , Ba , Sr , K or Na such and ppts. of double succrates form solutions of the first three bases are heated

at two cases even at 100°.

a, Ba, or K succates + Aq.
when an excess of cane
t. (Peligot.)

amyl amine, easily sol. in
amine. (Wurts.)

q. (Pelouse.)

ence of aromatic oxyacids
the series. Thus in pres-
cid, pyrocatechin, gallic
l, etc., NaOH + Aq does
from Cu solutions, but
ted by benzoic acid, re-
re, etc. (Weith, B. 9.

of alkali salts of "spal-
albumen. (Kalle and

usion which has become
ternating magnetic field.
08, I. 593.)

abénovitch, C. R. 1899,

emann, Z. anorg. 1893,

ross, Gm.—K. 5. 1, 753.)

sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
2, (7) 27. 393.)

monia, CuO_2H_2 , 4NH_3 .
iacal solution of CuO_2H_2 .
th. 1909, 69. 110.)

NH_3 .

forming Cu_2N .

ed by H_2O .

I_2 solutions of NH_4NO_3 .
rem. Soc. 1912, 34. 1502.)

I_2 .

dil. acids.

electrical conductivity of
O dissolves about 8 mg.
ausch and Rose, Z. phys.

SO_4 . (Vitali, Gm.—K.

in conc $\text{HCl} + \text{Aq}$.

HNO_3 , or H_2SO_4 . Insol.

SO_4 , KBr , or $\text{NH}_4\text{Cl} +$

$\text{Na}_2\text{S}_2\text{O}_3$, KCN , or $\text{KI} +$

t. 59. 558.)

1 N/10 HCl . Practically

.. (Moser, Z. anal. 1904,

ments on solubility of

presence of acids and

ray and Mackay.

to be sol. in H_2O but

dissolves in presence of

on of CuI_2 and CuI . (J.

0, 32. 1207.)

Solubility of Cu_2I_2 in $\text{I}_2 + \text{Aq}$ at 20°.

g. per l.		Solid Phase
Cu	I	
0.285	0.585	Cu_2I_2
0.482	1.305	"
0.583	1.922	"
0.678	2.557	"
0.756	3.204	"
0.844	3.954	"
0.898	4.436	"
0.964	5.085	"
1.032	5.685	"
1.090	6.292	"
1.112	6.530	"
1.232	7.653	$\text{Cu}_2\text{I}_2 + \text{I}_2$
1.040	6.449	I_2
0.898	5.594	"
0.748	4.711	"
0.606	3.856	"
0.448	2.949	"
0.300	2.069	"
0.159	1.230	"
0.925*	5.461	$\text{Cu}_2\text{I}_2 + \text{I}_2$
1.658**	11.366	"

* at 0°. ** at 40°.

(Fedotieff, Z. anorg. 1911, 69. 26.)

Solubility in $\text{NH}_4\text{Br} + \text{Aq}$ at 20°.

$\text{NH}_4\text{Br} + \text{Aq}$.	g. Cu_2I_2 in 1 l. of the solution
2-N	1.9068
3-N	3.6540
4-N	6.0588

(Kohn and Klein, Z. anorg. 1912, 77. 254.)

Sol. in $\text{FeCl}_3 + \text{Aq}$. (Fleischer, C. N. 1869,
19. 206.)

Solubility in $\text{KBr} + \text{Aq}$ at t°.

t°	$\text{KBr} + \text{Aq}$.	g. Cu_2I_2 in 1 l. of the solution
19.5	2-N	1.4666
24.0	2-N	1.5576
19.5	3-N	3.4094
23.0	3-N	3.5949
22.0	4-N	7.1263
22.0	4-N	6.9768

The solutions undergo change in the course
of a few days, iodine being set free; the sol-
ubility of the cuprous iodide is not markedly
affected thereby.

(Kohn, Z. anorg. 1909, 63. 337.)

1 l. of a 0.2N solution of KI dissolves
0.000157 g. mol Cu_2I_2 . (Bodländer, Z.
anorg. 1902, 31. 475.)

Very sol. in liquid NH_3 . (Franklin, Am.
Ch. J. 1898, 20. 827.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Practically insol. in methylene iodide. (Retgers, Z. anorg. 1893, 3. 347.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in acetone (Naumann, B. 1904, 37. 4329); (Eidmann, C. C. 1899, II. 1014.)

100 g. acetonitrile dissolve 3.52 g. Cu_2I_2 at 18° . (Naumann and Schier, B. 1914, 47. 249.)

Min. *Marshite*. (Gm.—K. 5. 1, 945.)

Cupric iodide, CuI_2 .

Exists only in very dil. aqueous solution. (Traube, B. 17. 1064.)

Copper periodide, CuI_4 .

Sol. in H_2O . (Walker and Dover, Chem. Soc. 1905, 87. 1588.)

Copper ammonium iodide ammonia.

See Cupriammonium iodide ammonia.

Cuprous mercuric iodide, $\text{Cu}_2\text{I}_2, \text{HgI}_2$.

$\text{KI} + \text{Aq}$ dissolves out HgI_2 .

Cuprous mercuric iodide ammonia, $\text{CuI}_2, 2\text{HgI}_2, 4\text{NH}_3$.

Decomp. by H_2O or acids. Sol. in a mixture of acetic acid and alcohol.

$\text{CuI}_2, \text{HgI}_2, 4\text{NH}_3$. As above. (Jørgensen, J. pr. (2) 2. 347.)

Cupric nitrogen iodide, $\text{CuI}_2, \text{N}_2\text{H}_4\text{I}_2$.

Decomp. by H_2O ; or $\text{NH}_4\text{OH} + \text{Aq}$. (Guyard, C. R. 97. 526.)

Cupric thallic iodide ammonia, $\text{CuI}_2, 2\text{THI}_2, 4\text{NH}_3$.

Decomp. slowly by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ with decomp. Sol. in alcohol.

Cuprous iodide ammonia, $\text{Cu}_2\text{I}_2, \text{NH}_3$.

Ppt. (Anderline, Gazz. ch. it. 1912, 42. I, 321.)

$+4\text{H}_2\text{O}$. Insol. in H_2O . Very sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Silberrad, Chem. Soc. 1905, 87. 67.)

$\text{Cu}_2\text{I}_2, 3\text{NH}_3$. (Lloyd, J. phys. Chem. 1908, 12. 399.)

$\text{Cu}_2\text{I}_2, 4\text{NH}_3$. (Levol, J. Pharm. 4. 328.)

$+ \text{H}_2\text{O}$. (Saglier, C. R. 104. 1440.)

$\text{Cu}_2\text{I}_2, 6\text{NH}_3$. (Lloyd.)

Cupric iodide ammonia, $\text{CuI}_2, 4\text{NH}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ without decomp. Not attacked by cold

alcohol or ether. (Berthemon 15. 445.) (Pozzi-Escot, C. R. 1

$\text{CuI}_2, 6\text{NH}_3$. Sol. in liquid H_2O . Am. Ch. J. 1908, 39. 205.)

$3\text{CuI}_2, 10\text{NH}_3$. Decomp. by acids, Am. Ch. J. 1895, 17. 302.

Sol. in liq. NH_3 . (Horn, Am 39. 204.)

Cupriammonium iodide ammonia, $3\text{Cu}(\text{NH}_3)_2\text{I}_2, 4\text{NH}_3$.

Decomp. by air and by H_2O . Am. Ch. J. 1895, 17. 302.

Copper periodide ammonia, $2\text{C} + \text{H}_2\text{O}$.

Because of its insolubility recryst from any solvent. (Sill Soc. 1905, 87. 66.)

Copper tetraiodide ammonia, C (Jørgensen, J. pr. (2) 2. 353.)

Copper hexaiodide ammonia, C

Not decomp. in H_2O in c (Jørgensen.)

Copper mercuric iodide ammonia, $\text{CuHg}_2\text{I}_2, 5\text{NH}_3$.

$\text{CuHgI}_2, 2\text{NH}_3$. Ppt. Dec and by alcohol.

$\text{CuHg}_2\text{I}_2, 3\text{NH}_3$. Ppt. Dec washing with H_2O .

$\text{CuHg}_2\text{I}_2, 4\text{NH}_3$. Ppt. Dec Sl. attacked by abs. alcohol.

$\text{CuI}_2, \text{HgI}_2, 4\text{NH}_3$. Ppt. (A ch. it. 1912, 42, (1) 321; C. C.

Copper nitride, Cu_4N_2 .

Decomp. by dil. or conc. acid Easily decomp. by H_2O when dered. (Rossel, C. R. 1895, 12

Copper suboxide, Cu_2O .

Not attacked by H_2O . De $\text{H}_2\text{SO}_4 + \text{Aq}$ into Cu and CuSO_4 has similar action. Not attack $+ \text{Aq}$ or $\text{NH}_4\text{OH} + (\text{NH}_4)_2\text{CO}_3$, Pogg. 120. 1.)

Cu_2O . Not attacked by dil. acids, even aqua regia. Slowly Aq. (Bailey and Hopkins, Ch 57. 272.)

Is a solution of oxide in Zeit. angew. Ch. 1908, 21. 51.

Cuprous oxide, Cu_2O .

Insol. in H_2O . Decomp. by $\text{H}_3\text{PO}_4 + \text{Aq}$, or cold very dil. H a cupric salt and Cu. Conver Aq into cuprous chloride.

[Cu₂O in NH₄OH + Aq at 25°.

Total Cu G. mol. in 1000 g. of solution	Conc. of total NH ₃	
	G. in 1000 g. of solution	G. mol. in 1000 g. of solution
Preparation I		
0.00566	3.91	0.23
0.00791	12.07	0.71
0.01080	13.77	0.81
0.01095	16.15	0.95
0.01597	27.03	1.59
0.01645	32.64	1.92
0.01660	36.89	2.17
0.01924	45.73	2.69
0.02081	68.68	4.04
0.02340	74.12	4.36
0.02375	81.26	4.78
0.02565	98.52	5.56
0.02670	122.40	7.20

Preparation II		
0.00665	7.82	0.46
0.01050	8.16	0.48
0.01555	22.61	1.33
0.01650	28.39	1.67
0.02127	54.15	3.19
0.02366	72.08	4.24
0.02510	78.20	4.60
0.02603	102.05	6.00

l Thomas, Chem. Soc. 1911, 99. 1791.)

ng NH₄Cl + Aq. (Rose.)
cess of KOH + Aq. (Chodnew.)
nc. MgCl₂, and FeCl₂ + Aq.
69. 1357.)
l by liquid NH₃. (Gore, Am.
0. 827.)
ite. Sol. in HCl, HNO₃, and

CuO.
O. Easily sol. in acids. Sol.
l. Insol. in NH₄OH + Aq, but
ddition of a few drops of acid
+ Aq. Insol. in dil., but sol. in
iaOH, and KOH + Aq. (Low,
.)
ed at a low temp. is easily sol.
out when ignited is slowly sol.
s. acids, but moderately rapidly
ture of NH₄I + HCl. (Joannis,
12. 1161.)
1 N-HNO₃. 1 l. of the solution
2 g. atoms Cu at 25°. (Jaeger,
, 27, 33.)

Solubility of CuO in HF + Aq at 25°.

	Time	G. CuO in 10 cem. of the solution
0.25N-HF	1¼ hrs.	0.0431
	3¾ "	0.0619
	25¼ "	0.0812
	71½ "	0.0823
	170½ "	0.0907
N-HF (a)	5½ "	0.3018
	21½ "	0.2797
	52 "	0.2747
	201½ "	0.2339
	226½ "	0.2353
N-HF (b)	4¼ "	0.3220
	44¼ "	0.2930
	117¾ "	0.2431
	167¾ "	0.2219
2.02N-HF	1½ "	0.3646
	5 "	0.4533
	71½ "	0.3583
	156½ "	0.3311

(Deussen, Z. anorg. 1905, 44. 421.)

Solubility of CuO in HF at 25°.
Cu = g-atoms Cu in 1 l. of the solution.

HF normality	Cu
0.12	0.0307
0.28	0.1164
0.57	0.2494
1.08	0.388
2.28	0.463

(Jaeger, Z. anorg. 1901, 27. 29.)

Solubility of CuO in HF + KF at 25°.
Cu = g-atoms Cu in 1 l. of the solution.

HF normality	Cu
0.12	0.0356
0.28	0.06437
0.57	0.1442
1.11(1.08)	0.2451
2.17(2.28)	0.2517

(Jaeger, l. c.)

Sl. sol. in large excess of KOH + Aq. (de
Coninck, C. C. 1904, II, 65.)
Slowly sol. in boiling NH₄Cl + Aq. and
less easily in NH₄NO₃ + Aq. (Rose.)
Sol. in boiling H₂O solutions of Al₂, Gl,
U, Cr₂, Fe₂, or Bi nitrates and chlorides,
Hg(NO₃)₂, Hg₂(NO₃)₂, SbCl₃, SnCl₂, and
SnCl₄, with pptn. of oxides of the bases of
those salts. Unacted upon by boiling H₂O
solutions of Mn, Mg. Ni, Co, Zn, Ce, or

Fe nitrates or chlorides, AgNO_3 , $\text{Pb}(\text{NO}_3)_2$, $\text{Cd}(\text{NO}_3)_2$, and HgCl_2 . (Persoz.)

Pure CuO is very sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$ but the solution is greatly increased by the addition of NH_4 salts. (Muthmann, C. C. 1904, II, 410.)

Sol. in hot $(\text{NH}_4)_2\text{SO}_4$ or $(\text{NH}_4)_2\text{SO}_3 + \text{Aq}$. (Jumau, Electrochem. Ind. 1908, 6. 258.)

15% dissolves in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ in 24 hrs. (Schnabel, Z. B. H. Sal. 1880, 28. 282.)

Sl. attacked by liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Solubility in N-acetic acid. 1 l. of the solution contains 0.1677 g-atoms Cu at 25° . (Jaeger, Z. anorg. 1902, 27. 33.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Sl. sol. in benzamide. (Dessaigues, A. ch. 1852, (3), 34. 146.)

Insol. in piperidine. (Cahours, C. R. 1852, 34. 481.)

Sol. in acid amines as asparagin. (Piria, A. ch. 1848, (3), 22. 160.)

Sol. in amines alone or mixed with NH_3 . (Lance, Dissert. 1905.)

Slowly sol. in Ca or any other alkali succrate + Aq, but not in cane sugar + Aq. (Hunton.)

Solubility in (calcium succrate + sugar) + Aq.

1 l. solution containing 418.6 g. sugar and 34.3 g. CaO dissolves 10.26 g. CuO .

1 l. solution containing 296.5 g. sugar and 24.2 g. CaO dissolves 5.68 g. CuO .

1 l. solution containing 174.4 g. sugar and 14.1 g. CaO dissolves 3.47 g. CuO . (Bodenbender, J. B. 1865. 600.)

Polypeptides in aqueous solution dissolve CuO by short boiling. (Fischer, B. 1906, 39. 576.)

$+ \frac{1}{2}\text{H}_2\text{O} = 6\text{CuO} + \text{H}_2\text{O}$. Insol. in dil., but sol. in conc. KOH or $\text{NaOH} + \text{Aq}$.

Sol. in volatile oils.

See also Cupric hydroxide.

Min. *Melaconite*. Sol. in HCl , or $\text{HNO}_3 + \text{Aq}$.

Cuprocupric oxide, $\text{Cu}_3\text{O}_2 = 2\text{Cu}_2\text{O}$, CuO .

(Favre and Maumené.)

$\text{Cu}_3\text{O}_2 + \text{H}_2\text{O} = \text{Cu}_2\text{O}$, $\text{CuO} + \text{H}_2\text{O}$. When freshly pptd., sol. in $\text{HCl} + \text{Aq}$, but insol. after drying. (Siewert, J. B. 1866. 257.)

$\text{Cu}_3\text{O}_2 = \text{Cu}_2\text{O}$, 2CuO . (Siewert.)

All oxides of Cu except Cu_2O , Cu_2O , CuO , and Cu_2O_2 are mixtures. (Osborne, Sill. Am. J. (3) 32. 33; Debray, C. R. 99. 583.)

Copper dioxide, $\text{CuO}_2 + \text{H}_2\text{O}$.

Insol. in H_2O . Decomp. by acids with formation of cupric salt and H_2O_2 . (Weltzien, A. 140. 207.)

Cuprous oxide ammonia (cuprosammonium oxide).

Known only in solution. (Wagner, C. C. 1863. 239.)

Cupric oxide ammonia (cuprammonia-dioxide), 3CuO , $4\text{NH}_3 + 6\text{H}_2\text{O}$.

Insol. in H_2O . (Kane, A. ch. 72. 2)

CuO , $4\text{NH}_3 + 4\text{H}_2\text{O}$. Very deliq.

Decomp. in the air and by H_2O . (Kane and Sarzeau, A. ch. (3) 9. 438.)

Cuprous oxybromide, Cu_2Br_2 , $\text{CuO} +$

(Spring and Lucion, Bull. Ac. F. 24. 21.)

Cupric oxybromide, CuBr_2 , $3\text{CuO} + 3$

Insol. in H_2O . Easily sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Brun, C. R. 109. 66.)

Insol. in H_2O but decomp. by boiling. Sol. in conc. acetic acid, a conc. $\text{CuBr}_2 + \text{Aq}$. Insol. in dil. K (Richards, Proc. Am. Acad. 1890, 25.)

Cupric oxybromide ammonia, 2CuO , $2\text{NH}_3 + 3\text{H}_2\text{O}$.

(Kohlschütter and Pudschies, B. 1159.)

Cuprous oxychloride, Cu_2Cl_2 , $\text{CuO} +$

(Spring and Lucion, Bull. Ac. F. 24. 21.)

Cupric oxychloride, CuO , $\text{CuCl}_2 + \text{H}_2\text{O}$

Decomp. by H_2O . (Rousseau, C. 110. 1263.)

2CuO , CuCl_2 . Insol. in H_2O . Sol. + Aq, from which it is reprecipitated by dilution with H_2O .

+ H_2O . (Kane, A. ch. 72. 277.)

+ $4\text{H}_2\text{O}$. (Gladstone, Chem. Soc. 3CuO, $\text{CuCl}_2 + 2\text{H}_2\text{O}$. (Miller and

rick, Trans. Roy. Soc. Can. 1901, (2) 35.)

+ $3\text{H}_2\text{O}$. (Dupont and Janse Soc. 1893, (3), 9. 193.)

+ $3\frac{1}{2}\text{H}_2\text{O}$. Insol. in cold H_2O , sl. by boiling. (Reindel, J. pr. 106. 378.)

Insol. in boiling H_2O . (Haberman B. 90. 2. 268.)

+ $4\text{H}_2\text{O}$. Sol. in alkaline solution tartrate. (Gröger, Z. anorg. 1902, (Brunswick green). Insol. in H_2O

sol. in acids.

Min. *Atacamite*. Sol. in acids, and + Aq.

Sol. in cold sat. citric acid + Aq. B. 1880, 13. 732.)

4CuO , $\text{CuCl}_2 + 6\text{H}_2\text{O}$. (Kane, 5. 1, 919.)

+ $8\text{H}_2\text{O}$. Min. *Tallingite*. (Chun — K. 5. 1, 919.)

$5\text{Cu}(\text{OH})_2$, $\text{Cu}_2\text{Cl}_2\text{Cl}(\text{OH})$. Insol. Decomp. by hot H_2O . (Kühling, 34. 2852.)

7CuO , $2\text{CuCl}_2 + 9\text{H}_2\text{O}$. (Reindel 6CuO , $\text{CuCl}_2 + 9\text{H}_2\text{O}$. Insol. in H_2O , acetic acid. (Neumann, Repert, 37.)

8CuO , $\text{CuCl}_2 + 12\text{H}_2\text{O}$. Min. (König, Zeit. Kryst. 1891, 19. 601.)

inc oxychloride, ZnO , 2ZnCl_2 , 5CuO , H_2O .

é, C. R. 1888, 106. 855.)

oxychloride ammonia, 2CuO , CuCl_2 , I_2 , $+3\text{H}_2\text{O}$.

rain, Gm.—K. 5. 1, 932.)

oxyfluoride, CuO , CuF_2 , $+ \text{H}_2\text{O}$.

in H_2O . (Berzelius.) (Balbiano, it. 14. 74.)

oxyfluoride ammonia (cuprammonium fluoride), $\text{Cu}(\text{OH})\text{F}$, 2NH_3 .

ano, Gazz. ch. it. 14. 74.)

$\text{CuI}_2 + x\text{H}_2\text{O}$. (Tschiriwinski, Gm. l, 1584.)

oxyiodide, Cu_2I_2 , $\text{CuO} + \text{H}_2\text{O}$.

g and Lucion, Bull. Ac. Belg. (3)

oxyiodide, 2CuI_2 , $\text{CuO} + 4\text{H}_2\text{O}$.

decomp. by H_2O . (Carnegie, Watts' 257.)

oxysulphide, $2\text{Cu}_2\text{S}$, CuO .

in H_2O . (Maumené, A. ch. (3) 18.

CuO . Ppt. (Pelouze.)

CuO . Insol. in H_2O .

CuO . Insol. in H_2O .

comps. do not exist. (Pickering, oc. 33. 136.)

phosphide, Cu_3P_2 .

sol. in HNO_3 or aqua regia; insol. in q. (Rose, Pogg. 6. 209.)

in HNO_3 and $\text{Br}_2 + \text{Aq}$. Decomp. by H_2SO_4 . (Rubénovitch, C. R. 1899, 9.)

Sol. in $\text{HNO}_3 + \text{Aq}$. (Granger, 1898, (7), 14. 64.)

Used. Completely sol. in hot HNO_3 , HCl and $\text{HF} + \text{HNO}_3$. Slowly sol. in H_2SO_4 . Not attacked by hot or cold acetic acid. (Maronneau, C. R. 8. 939.)

Easily sol. in HNO_3 . Sol. in hot H_2SO_4 . Sol. in conc. $\text{HCl} + \text{Aq}$ before phosphide has been heated. (Rose, Pogg.

Easily sol. in HNO_3 , or $\text{HCl} + \text{Aq}$. $\text{NH}_4\text{OH} + \text{Aq}$. (Granger, Bull. Soc. l.)

Decomp. by HNO_3 ; not readily Cl_2 . Easily attacked by Cl_2 or Br_2 , + Granger, C. R. 1895, 120. 924.)

(Granger, C. N. 1898, 77. 229.)

sol. in HNO_3 and $\text{Br}_2 + \text{Aq}$. Decomp. by conc. H_2SO_4 . (Rubénovitch, C. R. 9. 338.)

Cupric zinc phosphide, $10\text{Cu}_3\text{P}_2$, $\text{Zn}_3\text{P}_2(?)$.

(Hvoslef, A. 100. 99.)

Copper phosphoselenide, CuSe , P_2Se_3 .

Insol. in H_2O or $\text{HCl} + \text{Aq}$; sol. in $\text{HNO}_3 + \text{Aq}$. Insol. in cold alkalies, but decomp. slowly when heated therewith. (Hahn, J. pr. 98. 436.)

2CuSe , P_2Se_3 . Attacked only by fuming HNO_3 . (Hahn.)

2CuSe , P_2Se_3 . Sol. only in $\text{HNO}_3 + \text{Aq}$. (Hahn.)

Copper phosphosulphide, $2\text{Cu}_2\text{S}$, P_2S_5 .

Cu_2S , P_2S_5 . (Berzelius.)

$2\text{Cu}_2\text{S}$, P_2S_5 . (Berzelius.)

CuS , P_2S_5 . Insol. in H_2O and dil. $\text{HCl} + \text{Aq}$. Sol. in conc. $\text{HCl} + \text{Aq}$, from which it is precipitated by H_2O . (Berzelius, A. 46. 252.)

8CuS , P_2S_5 . (Berzelius.)

Cu_4PS_5 . Sol. in conc. HNO_3 and in aqua regia. Insol. in HCl . Not attacked by hot H_2SO_4 or conc. $\text{NaOH} + \text{Aq}$. (Ferrand, A. ch. 1899, (7), 17. 407.)

Cuprous selenide, Cu_2Se .

Ppt. Sol. in HCl and in H_2SO_4 .

Decomp. by HNO_3 . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Fonzes-Diacon, C. R. 1900, 131. 1207.)

Sol. in $\text{KCN} + \text{Aq}$. (Heyn and Bauer, Metall. 1903, 3. 84.)

Min. Berzelianite.

Cupric selenide, CuSe .

(Little, A. 112. 211.)

Ppt. Sol. in HCl and in H_2SO_4 . Decomp. by HNO_3 . (Fonzes-Diacon, C. R. 1900, 131. 1207.)

Cuprocupric selenide, Cu_3Se_2 .

Min. Umangite. Sol. in HNO_3 . (Klockmann, Zeit. Kryst. 1891. 19, 270.)

Cuprous lead selenide, $3\text{Cu}_2\text{Se}$, PbSe .

Min. Zorgite. Sol. in cold conc. $\text{HNO}_3 + \text{Aq}$ with separation of Se.

Cupric lead selenide, CuSe , PbSe .

Sol. in cold conc. HNO_3 with separation of Se. (Karsten.)

CuSe , 2PbSe . As above.

CuSe , 4PbSe . As above.

Cuprous silver selenide, Cu_2Se , Ag_2Se .

Min. Eucainite. Sol. in hot HNO_3 with decomp. (Berzelius.)

Cuprous silicide, Cu_4Si .

Sol. in warm dil. or conc. HNO_3 . Only sl. sol. in HCl , H_2SO_4 and HF . Sol. in a mixture of HNO_3 and HF . Not attacked by solutions of alkalies. (Vigouroux, C. R. 1906, 142. 88.)

Cu_2Si . Sol. in aqua regia and fused sodium potassium carbonate. (de Chalmot, Am. Ch. J. 1896, 18, 95.)

Cu_2Si . Decomp. by water and moist air, and by acids and fused alkali. (Vigouroux, C. R. 1896, 122, 319.)

Cuprous sulphide, Cu_2S .

More sol. in H_2O than Ag_2S , but much less than PbS . (Bodländer, Z. phys. Ch. 1898, 27, 64.)

1 l. H_2O dissolves 3.1×10^{-6} moles Cu_2S at 18° . (Weigel, Z. phys. Ch. 1907, 58, 294.)

Very slowly decomp. by dil. H_2SO_4 in presence of oxygen. (Thompson, Electrochem, Ind. 1904, 2, 225.)

Decomp. by conc. H_2SO_4 . (Pickering, C. N. 1878, 37, 37.)

Cold $\text{HNO}_3 + \text{Aq}$ dissolves out Cu and leaves CuS ; hot HNO_3 dissolves with separation of S. Sl. sol. in boiling conc. $\text{HCl} + \text{Aq}$. Insol. in $(\text{NH}_4)_2\text{S} + \text{Aq}$.

5N- HCl dissolves Cu_2S very slightly (0.0038 g. Cu in $7\frac{1}{2}$ hrs.) but it is more sol. in presence of Cl, when 0.672 g. are dissolved in $7\frac{1}{2}$ hours. (Egli, Z. anorg. 1902, 30, 46.)

Sol. with exclusion of air in $\text{NH}_4\text{OH} + \text{Aq}$. (Malzac, Pat. 1904.)

Insol. in acetone. (Naumann, B. 1904, 37, 4329; Eidmann, C. C. 1899, II, 1014.)

Min. *Chalcocite*. Completely sol. in warm HNO_3 with separation of S.

Cupric sulphide, CuS .

Almost absolutely insol. in H_2O ; sol. in 950,000 pts. H_2O . When exposed to the air, dissolves in H_2O as CuSO_4 . Easily sol. in boiling HNO_3 with separation of S. Difficultly sol. in hot conc. $\text{HCl} + \text{Aq}$. Insol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ (1:6). (Hoffmann, A. 115, 286.)

Pptd. by H_2S or $(\text{NH}_4)_2\text{S} + \text{Aq}$ in presence of 100,000 pts. H_2O (Pfaff), 200,000 pts. H_2O (Lassaigne), 15,000 pts. H_2O and 7500 pts. HCl , but with 40,000 pts. H_2O and 20,000 pts. HCl no colour is visible (Reinsch).

1 l. H_2O dissolves 3.51×10^{-6} moles CuS at 18° . (Weigel, Z. phys. Ch. 1907, 58, 294.)

Insol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Guerout, C. R. 1872, 75, 1276.)

Decomp. by conc. H_2SO_4 . (Kliche, J. B. 1890, 593.)

Sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Berzelius.) Sol. in alkali bicarbonates + Aq .

Insol. in NH_4NO_3 or $\text{NH}_4\text{Cl} + \text{Aq}$. (Brett.)

Insol. in acidified conc. alkali chlorides + Aq . (Cushmann, Am. Ch. J. 1895, 17, 382.)

Sol. in $\text{FeCl}_3 + \text{Aq}$ with separation of S. (Cumenge and Wimmer, Dingl. 1883, 250, 123.)

Decomp. by boiling $\text{CuCl}_2 + \text{Aq}$ in presence of HCl or NaCl . (Raschig, Gm.—K. 5, 1, 819.)

Sol. in $\text{Fe}_2(\text{SO}_4)_3 + \text{Aq}$ in presence of large excess of air. (Thompson, Electrochem. Ind. 1904, 2, 228.)

Insol. in KOH , or $\text{K}_2\text{S} + \text{Aq}$, especial boiling; appreciably sol. in colourless even more readily in hot yellow $(\text{NH}_4)_2\text{S} + \text{Aq}$. (Becker, Sill. Am. J. (3) 33, 199.)

100 cc. sat. $\text{Na}_2\text{S} + \text{Aq}$ (sp. gr. = 1.1) dissolve 0.0032 g. CuS . (Holland, Chim. Anal. 1897, 2, 243.)

Sol. in K polysulphides (3-64%). (P. Bull. Soc. Belg. Chim. 1897, 103.)

Appreciably sol. in alkali polysulphide Aq . (Rössing, Z. anal. 1902, 41, 1.)

Sol. in considerable quantity in alkali arsenates, sulphantimonates, and stannates + Aq . Therefore when a ppt. of CuS and As_2S_3 , Sb_2S_3 , or SnS is treated with K_2S , a portion of the CuS is dissolved (Wöhler, A. 34, 236.)

Sol. in alkali sulphovandates, or tungstates + Aq . (Storch, B. 16, 2015.)

Sol. in alkali sulphomolybdates + (Debray, C. R. 96, 1616.)

Insol. in K thiocarbonate + Aq . (Rebladt, Z. anal. 26, 15.)

Sol. in $\text{KCN} + \text{Aq}$.

Insol. in liquid NH_3 . (Franklin, Am. J. 1898, 20, 827.)

Insol. in methyl acetate (Naumann, 1909, 42, 3790); ethyl acetate. (Naumann, B. 1910, 43, 314.)

Insol. in acetone. (Naumann, B. 1904, 37, 4329; Eidmann, C. C. 1899, II, 1014.)

Insol. in Na xanthogenate. (Ragg, Z. 1908, 32, 677.)

Solubility of CuS in sugar + Aq at t°
g. CuS per l. of solution.

t°	10% sugar	30% sugar	50% sugar
17.5	0.5672	0.8632	0.907
45	0.3659	0.7220	1.059
75	1.1345	1.2033	1.289

(Stolle, Z. Ver. Zuckerind. 1900, 50, 33)

Min. *Corellite*.

Colloidal. Aqueous solution is stable; it contains 5 g. CuS in a litre; when it contains 4 or 5 times that amount it is decomposed in an hour.

Solutions of salts of the following concentration cause a precipitate in the above solution. Salts of univalent elements—

Salts of univalent elements—

$\text{K}_3\text{Fe}(\text{CN})_6$.	.	.	1 : 62
$\text{K}_4\text{Fe}(\text{CN})_6$.	.	.	1 : 127
$\text{Na}_2\text{S}_2\text{O}_3$.	.	.	1 : 157
Na_2CO_3	.	.	.	1 : 200
Na_2HPO_4	.	.	.	1 : 252
Na_2SO_4	.	.	.	1 : 333
$\text{K}_2\text{Cr}_2\text{O}_7$.	.	.	1 : 2083
KI	.	.	.	1 : 80
KBr	.	.	.	1 : 133
KClO_3	.	.	.	1 : 166

univalent elements—Continued.

$C_2H_5O_2$	1 : 221
$H_4)_2C_2O_4$	1 : 255
Cl	1 : 400
HCO_2	1 : 2500
SO_4	1 : 117
CrO_4	1 : 133
$C_7H_5O_2$	1 : 166
S_2O_6	1 : 222
I	1 : 333
O_2	1 : 500

bivalent metals—

S_2O_6	1 : 2242
$NO_3)_2$	1 : 3483
SO_4	1 : 6830
$NO_3)_2$	1 : 2677
Cl_2	1 : 3921
$ClO_3)_2$	1 : 6988
SO_4	1 : 3442
SO_4	1 : 5518

trivalent metals—

monia alum	1 : 31,896
ome alum	1 : 58,889
$SO_4)_3$	1 : 90,909

cinic	1 : 100
lic	1 : 162
l	1 : 733
O_4	1 : 208
ic	1 : 20
tic	Not at all
taric	" "

and de Boeck, Bull. Soc. (2) 58. 165.)

polysulphide, Cu_2S_2 .

rhous. Ppt. Decomp. by boiling (Rössing, Z. anorg. 1900, 25. 413.)

Amorphous. Ppt. can be boiled without decomposition. (Rössing, Z. anorg. 1900, 25. 4, 11.)

Ppt.; insol. in alkali sulphides; by conc. HNO_3 . (Bodroux, C. R. 1900, 1398.)

not be obtained. (Rössing, Z. anorg. 1900, 25. 414.)

Ppt. Decomp. by H_2O . Sol. in H_2O and barium polysulphides + Aq. by colorless alkali sulphides + Aq. (Rössing, Z. anorg. 1900, 25. 407.)

iron (ferric) sulphide, Cu_2S , Fe_2S_3 .
 Prep. by conc. HCl + Aq. Sol. in boiling Aq of 1.2 sp. gr. (Schneider, J. pr. 1869.)

Chalcopyrite. Insol. in HCl + Aq. heated in a sealed tube with H_2S + Aq, 1 of it dissolves with difficulty and without deposition of S. (Senarmont, A. l. 168.)

cupric iron (ferric) sulphide, Cu_2S , CuS ,

Bornite. Sol. in HCl + Aq with a precipitate of S.

Cupric iron (ferric) sulphide, CuS , Fe_2S_3 .

Min. *Cubanite*.

Copper iron potassium sulphide, $K_2FeCu_2S_4$.

Sl. attacked by cold dil. HCl + Aq. Decomp. by warming. (Schneider, Pogg. 138. 318.)

Copper iron sodium sulphide, $Na_2FeCu_2S_4$.

Sl. attacked by cold dil., easily decomp. by hot HCl + Aq. (Schneider, Pogg. 138. 318.)

Cuprous lead sulphide, $9Cu_2S$, $2PbS$.

$3Cu_2S$, $2PbS$.

$2Cu_2S$, $2PbS$. Min. *Cuproplumbite*.

Copper phosphorus sulphide.

See Copper phosphosulphide.

Cupric platinum sulphide.

See Sulphoplatinate, cupric.

Cuprous potassium sulphide, $4Cu_2S$, K_2S .

(Ditte, C. R. 98. 1429.)

Cuprocupric potassium sulphide, $3Cu_2S$, $2CuS$, K_2S .

Not decomp. by very dil. HCl + Aq, but easily by conc. HCl + Aq on warming. (Schneider, Pogg. 138. 311.)

Copper potassium polysulphide, $KCuS_4$.

Sl. sol. in cold H_2O . Decomp. by hot H_2O . Decomp. by conc. and dil. HCl , H_2SO_4 and HNO_3 . Sl. sol. in alcohol. (Biltz and Herms, B. 1907, 40. 977.)

$2CuS_2$, K_2S . Decomp. by H_2O , NH_4OH , or NH_4SH + Aq. (Priwoznik, B. 5. 1291.)

$K_2Cu_2S_{10}$. Easily sol. in H_2O . 1 g. is sol. in less than 5 ccm. H_2O . Rapidly decomp. by dil. acids, slowly by conc. acids. Sl. sol. in alcohol. (Biltz and Herms, B. 1907, 40. 983.)

Cupric rubidium polysulphide, $RbCuS_4$.

As K salt. (Biltz and Herms, B. 1907, 40. 978.)

$Rb_2Cu_2S_{10}$. Easily sol. in H_2O . Decomp. by acids. Sl. sol. in alcohol. (Biltz and Herms, B. 1907, 40. 985.)

Cuprous silver sulphide, Cu_2S , Ag_2S .

Min. *Stromeyerite*. Sol. in HNO_3 + Aq with separation of S.

Cu_2S , $3Ag_2S$. Min. *Jalpaite*. As above.

Cuprous sodium sulphide, Na_2S , Cu_2S .

(Bodländer, Z. Elektrochem. 1905, 11. 181.)

Na_2S , $2Cu_2S$. (Bodländer, Z. Elektrochem. 1905, 11. 181.)

Cuprocupric sodium sulphide, Cu_2S , CuS , Na_2S .

Scarcely decomp. by cold dil. $\text{HCl} + \text{Aq}$; conc. $\text{HCl} + \text{Aq}$ decomp. easily on warming, without, however, dissolving all the Cu_2S . Completely decomp. by warm $\text{HNO}_3 + \text{Aq}$. (Schneider, Pogg. 188. 315.)

Copper zinc sulphide, CuS , 3ZnS .

Copper sulphophosphide.

See Copper phosphosulphide.

Cupric telluride, CuTe .

Cu_2Te_3 . Insol. in H_2O . (Parkmann, Sill. Am. J. (2) 3. 335.)

Cu_2Te . (Brauner, M. 1889. 423.)

Croceocobaltic bromide,

$\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{Br}$.

Very sl. sol. in cold, easily in hot H_2O . (Gibbs, Proc. Am. Acad. 10. 1.)

— **chloraurate**, $2\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{Cl}$, AuCl_3 .
Difficultly sol. in H_2O .

— **chloride**, $\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{Cl}$.

Very sl. sol. in cold easily in hot H_2O , but more sol. than the sulphate. (Gibbs.)

— **chloroplatinate**, $2\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{Cl}$, PtCl_4 .

Can be recrystallised without decomp. with difficulty. (Gibbs and Genth, Sill. Am. J. (2) 24. 91.)

— **chromate**, $[\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2]_2\text{CrO}_4$.

Sl. sol. in H_2O . (Gibbs.)

— **dichromate**, $[\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2]_2\text{Cr}_2\text{O}_7$.

Sl. sol. in H_2O . (Gibbs.)

— **periodide**, $\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{I}$, I_2 .

Difficultly sol. in cold H_2O and alcohol. Decomp. by hot H_2O . (Gibbs.)

— **nitrate**, $\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2\text{NO}_3$.

Sl. sol. in cold, easily sol. in hot H_2O or dil. acids. Much more sol. than the sulphate. (Gibbs.)

Sol. in about 400 pts. cold H_2O . (Jørgensen, Z. anorg. 5. 163.)

— **nitrite cobaltic nitrite**, $3\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2$, $\text{Co}(\text{NO}_2)_3$.

Somewhat sol. in H_2O . (Jørgensen, Z. anorg. 5. 178.)

— **nitrite diamine cobaltic nitrite**, $\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2$, $(\text{NO}_2)_2(\text{NH}_3)_2\text{Co}(\text{NO}_2)_3$.

Nearly insol. in cold, very sl. sol. in boiling H_2O . (Jørgensen.)

Croceocobaltic phosphomolybdate,
 $[\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2]_2\text{O}$, 24MoO_3 , P_2O_5 .

Sl. sol. in cold, easily in hot H_2O . (Gibbs, Am. Ch. J. 3. 317.)

— **sulphate**, $[\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2]_2\text{SO}_4$.

Very sl. sol. in cold or hot H_2O ; more easily in hot dil. $\text{H}_2\text{SO}_4 + \text{Aq}$.

Cuprammonium compounds.

See Copper compounds, ammonia.

Cuprotetrammonium tetraiodide.

See Cupric tetraiodide ammonia.

Cupric acid.

Known only in solution. (Krüger, Pogg. 62. 445.)

Calcium cuprate.

Decomp. by H_2O with evolution of oxygen. (Krüger and Crum, A. 55. 213.)

Cyanhydric acid, HCN .

Miscible with H_2O , alcohol, and ether with absorption of heat.

Sp. gr. of $\text{HCN} + \text{Aq}$.

% HCN	Sp. gr.	% HCN	Sp. gr.
1.60	0.9979	4.0	0.9940
1.68	0.9978	4.6	0.9930
1.77	0.9975	5.0	0.9923
2.0	0.9974	5.3	0.9914
2.1	0.9973	5.8	0.9900
2.3	0.9970	6.4	0.9890
2.5	0.9967	7.3	0.9870
2.7	0.9964	8.0	0.9840
3.0	0.9958	9.1	0.9815
3.2	0.9952	10.6	0.9768
3.6	0.9945	16.0	0.9570

(Ure, Quar. J. Sci. 13. 321.)

2HCN mixed with $3\text{H}_2\text{O}$ causes a diminution of temp. of 9.75° . (Bussy and Buignet, A. ch. (4) 3. 231.)

Miscible with volatile oils and other organic compounds.

Cyanhydric iodhydric acid, HI , HCN .

Easily sol. in H_2O or alcohol, with rapid decomp. Sl. sol. in ether. (Gal, A. 128. 33.)

Cyanides.

The alkali cyanides are easily sol. in H_2O ; those of the alkali-carths are less sol., while all others are insol. with the exception of $\text{Hg}(\text{CN})_2$. All cyanides are sol. in $\text{KCN} + \text{Aq}$.

Ammonium cyanide, NH_4CN .

Unstable; easily sol. in H_2O and alcohol.

m cobaltic mercuric cyanide.
calticyanide, ammonium mercuric.

m cuprous cyanide, NH_4CN , $(\text{CN})_2$.

Decomp. by acids.

Sol. in H_2O , less sol. in alcohol.
 by acids and alkalies. (Treadwell
 wald, Z. anorg. 1904, **39**. 90.)

N , $\text{Cu}_2(\text{CN})_2$. Sl. sol. in H_2O , but
 by long boiling therewith. Sol. in
 q. (Dufau, A. **88**. 278.)

m cuprous cyanide ammonia,
 CN , $2\text{Cu}_2(\text{CN})_2$, NH_3 .

decomp. (Treadwell and Girse-
 anorg. 1904, **39**. 90.)

Insol. in cold, **decomp.** by boiling
 in $\text{NH}_4\text{OH} + \text{Aq.}$ (Fleurent, C. R.
 191.)

I , $\text{Cu}_2(\text{CN})_2$, 3NH_3 . Insol. in cold,
 boiling H_2O without **decomp.** Sol.
 $\text{I} + \text{Aq.}$ (Fleurent, C. R. 1891, **113**.)

I , $2\text{Cu}_2(\text{CN})_2$, $2\text{NH}_3 + 2\text{H}_2\text{O}$. (Fleu-
5. 498R.)

m gold (aurous) cyanide NH_4CN ,

sol. in cold or warm H_2O or in al-
sol. in ether.

m gold (auric) mercuric cyanide,
 I , $3\text{NH}_4\text{CN}$, $2\text{Au}_2\text{O}$, $\text{Hg}(\text{CN})_2$, HgO .
dt, Ch. Z. 1896, **20**. 633.)

m mercuric silver cyanide, basic,
 I , $2\text{Ag}_2\text{O}$, 3AgCN , $4\text{Hg}(\text{OH})\text{CN} +$
 O .

dt, Z. anorg. 1895, **9**. 431.)

m nickel cyanide, $2\text{NH}_4\text{CN}$, $(\text{CN})_2$.

decomposed.

m tungsten cyanide.

ngstocyanide, ammonium.

m zinc cyanide, $2\text{NH}_4\text{CN}$, $\text{Zn}(\text{CN})_2$, H_2O .

m cyanide mercuric nitrate silver
ide basic, $2\text{Hg}(\text{OH})\text{NO}_3$, $3\text{NH}_4\text{CN}$, CN .

dt, Z. anorg. 1895, **9**. 431.)

ricyanide, $\text{As}(\text{CN})_3$.

p. by H_2O . Not attacked by cold
 O_4 . **Decomp.** on heating. (Guenez,
2, **114**. 1188.)

Barium cyanide, $\text{Ba}(\text{CN})_2$.

Rather sl. sol. in H_2O , more easily in KCN
 $+ \text{Aq.}$ (Schulz, J. pr. **68**. 257.)

10 pts. H_2O dissolve 8 pts., and 10 pts. 70%
 alcohol dissolve 1.8 pts. $\text{Ba}(\text{CN})_2$ at 14.
 (Joannis, A. ch. (5) **26**. 489.)

Insol. in methyl acetate. (Naumann, B.
 1909, **42**. 3790.)

$+ 2\text{H}_2\text{O}$. Very deliquescent.

$\text{Ba}(\text{CN})_2$, BaO . (Drechsel, J. pr. (2) **21**.
 84.)

**Barium cadmium cyanide, $\text{Ba}(\text{CN})_2$,
 $\text{Cd}(\text{CN})_2 + \text{H}_2\text{O}$.**

Easily sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq.}$

Sl. sol. in alcohol. (Loebe, Dissert, **1902**.)

$2\text{Ba}(\text{CN})_2$, $3\text{Cd}(\text{CN})_2 + 10\text{H}_2\text{O}$. Sol. in
 H_2O . (Weselsky, B. **2**. 590.)

Barium cobaltous cobaltic cyanide.

See Cobaltocobalticyanide, barium.

Barium cuprous cyanide, $\text{Ba}(\text{CN})_2$, $\text{Cu}_2(\text{CN})_2$.

Sol. in H_2O without **decomp.** (Traube, Z.
 anorg. 1894, **8**. 21.)

$+ \text{H}_2\text{O}$. (Weselsky, B. **2**. 590.)

Could not be obtained. (Grossmann, Z.
 anorg. 1905, **43**. 101.)

$+ 4\text{H}_2\text{O}$. **Decomp.** by H_2O . (Grossmann,
 Z. anorg. 1905, **43**. 101.)

$2\text{Ba}(\text{CN})_2$, $\text{Cu}_2(\text{CN})_2 + 6\text{H}_2\text{O}$. **Decomp.**
 by H_2O . (Grossmann, Z. anorg. 1905, **43**.
 105.)

**Barium gold (aurous) cyanide, $\text{Ba}(\text{CN})_2$,
 $2\text{AuCN} + 2\text{H}_2\text{O}$.**

Sl. sol. in cold but easily sol. in hot H_2O .
 Sl. sol. in alcohol. (Lindbom, Lund Univ.
 Arsk. **12**. No.6.)

Barium iridium cyanide.

See Iridicyanide, barium.

**Barium manganous cyanide, $\text{Ba}(\text{CN})_2$,
 $2\text{Mn}(\text{CN})_2$.**

Ppt. (Descamps.)

*See also Manganocyanide and Mangani-
 cyanide, barium.*

**Barium mercuric cyanide, $(\text{Ba}(\text{CN})_2$,
 $\text{Hg}(\text{CN})_2 + 3\text{H}_2\text{O}$.**

Very hygroscopic. Very sol. in H_2O .
 (Grossmann, B. 1904, **37**. 4142.)

**Barium mercuric cyanide iodide, $\text{Ba}(\text{CN})_2$,
 $\text{HgI}_2 + 6\text{H}_2\text{O}$.**

(Varet, C. R. 1895, **121**. 499.)

**Barium palladium cyanide, $\text{Ba}(\text{CN})_2$,
 $\text{Pd}(\text{CN})_2 + 4\text{H}_2\text{O}$.**

See Palladocyanide, barium.

**Barium nickel cyanide, $\text{Ba}(\text{CN})_2$, $\text{Ni}(\text{CN})_2 +$
 $3\text{H}_2\text{O}$.**

Sol. in H_2O ; **decomp.** by acids with pptn.
 of $\text{Ni}(\text{CN})_2$. (Weselsky, B. **2**. 590.)

Barium silver cyanide, Ba(CN)_2 , $2\text{AgCN} + \text{H}_2\text{O}$.

Sol. in H_2O . (Weselsky, B. 2. 589.)

Barium zinc cyanide, Ba(CN)_2 , $\text{Zn(CN)}_2 + 2\text{H}_2\text{O}$.

Sol. in H_2O .

Cadmium cyanide, basic, CdO_2H_2 , $2\text{Cd(CN)}_2 + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O ; insol. in alcohol. (Loebe, Dissert, 1902.)

Cadmium cyanide, Cd(CN)_2 .

Sl. sol. in H_2O . 100 pts. H_2O dissolve 1.7 pts. Cd(CN)_2 at 15° . (Joannis.)

Easily sol. in acids; sol. in $\text{KCN} + \text{Aq}$. Sol. in warm $\text{NH}_4\text{OH} + \text{Aq}$, but insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Wittstein.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Cadmium calcium cyanide, Cd(CN)_2 , $4\text{Ca(CN)}_2 + 20\text{H}_2\text{O}$.

Sol. in H_2O and in alcohol. (Loebe, Dissert, 1902.)

Cadmium chromic cyanide.

See Chromicyanide, cadmium.

Cadmium cobaltic cyanide.

See Cobalticyanide, cadmium.

Cadmium cuprous cyanide, 2Cd(CN)_2 , $\text{Cu}_2(\text{CN})_2$.

Permanent. Insol. in H_2O . Sl. sol. in cold, easily in warm $\text{HCl} + \text{Aq}$ without decomp., except by long boiling. Insol. in NH_4OH , or NH_4 salts + Aq . (Schüler.)

Cadmium cupric cyanide, Cd(CN)_2 , Cu(CN)_2 .

Very unstable.

Cadmium gold (aurous) cyanide, Cd(CN)_2 , 2AuCN .

Nearly insol. in cold H_2O . Sl. sol. in boiling H_2O . Insol. in alcohol. (Lindbom.)

Cadmium mercuric cyanide, 2Cd(CN)_2 , 3Hg(CN)_2 .

Permanent. Readily sol. in cold H_2O . (Schüler.)

Cadmium mercuric cyanide mercuric iodide, Cd(CN)_2 , Hg(CN)_2 , $\text{HgI}_2 + 8\text{H}_2\text{O}$.

Very sol. in H_2O . (Varet, Bull. Soc. (3) 5. 8.)

+ $7\text{H}_2\text{O}$. Sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq}$. (Varet, C. R. 1890, 111, 679.)

Cadmium mercuric cyanide mercuric iodide, Cd(CN)_2 , Hg(CN)_2 , $\text{HgI}_2 + 8\text{H}_2\text{O}$.

Very sol. in H_2O . (Varet, Bull. Soc. (3) 5. 8.)

+ $7\text{H}_2\text{O}$. Sol. in H_2O and in NH_4OH (Varet, C. R. 1890, 111. 679.)

Cadmium mercuric cyanide mercuric ammonia, Cd(CN)_2 , Hg(CN)_2 , 4NH_3 .

Very easily decomp. (Varet, Bull. S. 6. 22.)

Cadmium molybdenum cyanide.

See Molybdocyanide, cadmium.

Cadmium potassium cyanide, Cd(CN)_2 , 2KCN .

Sol. in 3 pts. cold, and 1 pt. boiling H_2O . Insol. in absolute alcohol. (Rammelsberg.)

Cadmium sodium cyanide, $\text{Na}_2\text{Cd(CN)}_2$, $+3\text{H}_2\text{O}$.

Sol. in H_2O and in alcohol. (Loebert, 1902.)

Cadmium strontium cyanide, Cd(CN)_2 , $2\text{Sr(CN)}_2 + 3\text{H}_2\text{O}$.

Sol. in H_2O and in alcohol. (Loebert, 1902.)

Cadmium tungsten cyanide, $\text{Cd}_2\text{W(CN)}_8$, $8\text{H}_2\text{O}$.

Nearly insol. in H_2O .

Sl. sol. in dil. HCl . Sol. in conc. NH_4OH .

Insol. in organic solvents. (Olsson anorg. 1914, 88. 68.)

Cadmium cyanide dihydrazine, Cd(CN)_2 , $(\text{N}_2\text{H}_4)_2$.

Easily sol. in dil. acids. (Franse anorg. 1911, 70. 152.)

Cæsium cuprous cyanide, CsCN , CuCN , $1\frac{1}{2}\text{H}_2\text{O}$.

H_2O separates CuCN . (Grossman anorg. 1905, 43. 98.)

2CsCN , $\text{CuCN} + \text{H}_2\text{O}$. Sol. in H_2O . (Grossmann, Z. anorg. 1905, 43. 98.)

2CsCN , 3CuCN . Insol. in, and decomp. by H_2O . (Grossmann, Z. anorg. 43. 98.)

Cæsium tungsten cyanide.

See Tungstocyanide, cæsium.

Calcium cyanide, Ca(CN)_2 .

Sol. in H_2O , but the solution is very unstable. (Schulz.)

Ca(CN)_2 , $3\text{CaO} + 15\text{H}_2\text{O}$. Decomp. by H_2O . (Joannis, A. ch. (5) 28. 496.)

Calcium cuprous cyanide, Ca(CN)_2 , CuCN , $4\text{H}_2\text{O}$.

Easily decomp. by H_2O . (Grossmann, Z. anorg. 1905, 43. 106.)

Ca(CN)_2 , $3\text{CuCN} + 8\text{H}_2\text{O}$. Immed.

decomp. by H_2O . (Grossmann, Z. anorg. 1905, 43. 99.)

Calcium gold (aurous) cyanide, $Ca(CN)_2$, $2AuCN + 3H_2O$.

Easily sol. in hot or cold H_2O or in alcohol. (Lindbom.)

Calcium manganous cyanide, $Ca(CN)_2$, $2Mn(CN)_2$.

Ppt. (Descamps.)

See also **Manganocyanide, calcium**.

Calcium mercuric cyanide, $Ca(CN)_2$, $2Hg(CN)_2 + 8H_2O$.

Very deliquescent. (Grossmann, B. 1904, 37. 4143.)

$2Ca(CN)_2$, $3Hg(CN)_2 + 6H_2O$. Very sol. in H_2O . (Grossmann, B. 1904, 37. 4143.)

Calcium mercuric cyanide iodide, $Ca(CN)_2$, HgI_2 , $Hg(CN)_2 + 7H_2O$.

(Varet, C. R. 1895, 121. 499.)

Calcium nickel cyanide, $Ca(CN)_2$, $Ni(CN)_2 + xH_2O$.

Sol. in H_2O .

Calcium tungsten cyanide.

See **Tungstocyanide, calcium**.

Calcium zinc cyanide, $(Ca(CN)_2, Zn(CN)_2 + 3\frac{1}{2}H_2O$.

Sol. in H_2O and in alcohol. (Loebe, Dissert. 1902.)

Cerous cyanide (?).

Ppt. Very easily decomp. (Behringer, A. 42. 139.)

Chromic cyanide, with MCN.

See **Chromicyanide, M**.

Chromous potassium cyanide.

See **Chromocyanide, potassium**.

Cobaltous cyanide, $Co(CN)_2 + H_2O$.

Insol. in H_2O . Easily sol. in $NH_4OH + Aq$, and $KCN + Aq$; also in $(NH_4)_2CO_3$, or NH_4 succinate + Aq ; insol. in NH_4NO_3 , or $NH_4Cl + Aq$. (Wittstein.)

Cobaltous cyanide with 4MCN.

See **Cobaltocyanide, M**.

Cobaltic cyanide with 3MCN.

See **Cobaltocyanide, M**.

Cobalt gold (aurous) cyanide, $Co(CN)_2$, $2AuCN$.

Insol. in H_2O or cold $HCl + Aq$.

Cobalt hydrazine cyanide, $(N_2H_4)_4Co(CN)_6$.

Deliquescent. (Franzen, Z. anorg. 1911, 70. 155.)

Cobaltous cyanide ammonia, $Co(CN)_2$, $2NH_3$.

Unstable. (Peters, B. 1908, 41. 3178.)

Cuprous cyanide, $Cu_2(CN)_2$.

Insol. in H_2O and dil. acids. Sol. in NH_4OH , $(NH_4)_2SO_4$, or NH_4 succinate + Aq , and in hot NH_4Cl , or $NH_4NO_3 + Aq$. Sol. in conc. $HCl + Aq$. Sol. in $KCN + Aq$.

Easily sol. in conc. NH_4SCN or $KSCN + Aq$. Sl. sol. in $NaSCN + Aq$. (Grossmann, Z. anorg. 1903, 37. 408.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)

Very sl. sol. in pyridine. (Schroeder, Dissert. 1902.)

Mol. weight determined in pyridine. (Werner, Z. anorg. 1897, 15. 20.)

Cupric cyanide, $Cu(CN)_2$.

Easily decomp. Insol. in H_2O .

Sol. in pyridine. (Schroeder, Dissert. 1901.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Cuprocupric cyanide, $Cu(CN)_2$, $Cu_2(CN)_2 + 5H_2O$.

Insol. in H_2O , but decomp. by boiling. Sol. in cold conc. $HCl + Aq$. Sol. in $NH_4OH + Aq$. $(NH_4)_2CO_3 + Aq$, and in hot NH_4 salts + Aq . Easily sol. in $KCN + Aq$.

+ H_2O . Ppt. (Dufau.)

+ $Cu(CN)_2$, $2Cu_2(CN)_2 + H_2O$. Ppt.

Cuprous hydrazine cyanide, $Cu_2(CN)_2$, N_2H_5CN .

Insol. in alcohol and H_2O . (Ferratini, C. C. 1912, I. 1281.)

Cupric iridium cyanide.

See **Iridicyanide, cupric**.

Cuprous lithium cyanide, $Cu_2(CN)_2$, $LiCN + H_2O$.

Gradually decomp. by H_2O . (Grossmann, Z. anorg. 1905, 43. 97.)

Cuprous magnesium cyanide, $Cu_2(CN)_2$, $Mg(CN)_2 + 11H_2O$.

Decomp. by H_2O . (Grossmann, Z. anorg. 1905, 43. 103.)

Cuprous mercuric cyanide bromide, $Cu(CN)_2$, $2Hg(CN)_2$, $HgBr_2$.

Sol. in H_2O . (Varet, C. R. 1890, 110. 148.)

Cupric molybdenum cyanide ammonia.

See **Molybdocyanide ammonia, cupric**.

Cuprous potassium cyanide, $\text{Cu}_2(\text{CN})_2, 2\text{KCN}$.

Sl. sol. in H_2O , with partial decomp. Decomp. by acids, but not by alkalis.

Decomp. by boiling H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Fleurent, C. R. 1893, 116. 191.)

Sol. without decomp. in conc. KSCN . (Grossmann, Z. anorg. 1903, 37. 407.)

Sol. without decomp. in $\text{KCN} + \text{Aq}$. (Treadwell and Girsewald, Z. anorg. 1904, 38. 94.)

$\text{Cu}_2(\text{CN})_2, \text{KCN} + \text{H}_2\text{O}$. Almost insol. in cold H_2O . 100 cc. H_2O dissolve 0.0594 g. at 15° . Decomp. by much hot H_2O with separation of $\text{Cu}_2(\text{CN})_2$. Sol. in $\text{KCN} + \text{Aq}$ or in $\text{NH}_4\text{OH} + \text{Aq}$. (Treadwell and Girsewald, Z. anorg. 1904, 38. 93.)

$3\text{Cu}_2(\text{CN})_2, 4\text{KCN}$. Sol. in H_2O .

$\text{Cu}_2(\text{CN})_2, 6\text{KCN}$. Sol. in H_2O .

Cuprous potassium cyanide ammonia, $\text{Cu}_2(\text{CN})_2, \text{KCN}, \text{NH}_3$.

(Treadwell and Girsewald, Z. anorg. 1904, 39. 88.)

Cuprous potassium cyanide potassium sulphocyanide, $\text{Cu}_2(\text{CN})_2, 4\text{KCN}, 2\text{KSCN}, \text{H}_2\text{O}$.

Easily sol. in cold H_2O . (Itzig, B. 1902, 35. 108.)

Cupric potassium cyanide, $\text{Cu}(\text{CN})_2, 2\text{KCN}$.

Sol. in $\frac{3}{4}$ pt. H_2O at 15° and $\frac{1}{2}$ pt. at 100° . (Buignet, J. Pharm. 1859, (3), 35. 168.)

Cuprocupric potassium cyanide, $\text{Cu}_2(\text{CN})_2, \text{Cu}(\text{CN})_2, 2\text{KCN}$.

(Straus, Z. anorg. 1895, 9. 15.)

Cuprous rubidium cyanide, $\text{Cu}_2(\text{CN})_2, 2\text{RbCN}$.

Sl. sol. in H_2O . Pure H_2O separates CuCN . (Grossmann, Z. anorg. 1905, 43. 100.)

$3\text{Cu}_2(\text{CN})_2, 4\text{RbCN}$. Sl. sol. in H_2O . Pure H_2O separates CuCN . (Grossmann, Z. anorg. 1905, 43. 98.)

Cuprous silver cyanide, $\text{Cu}_2(\text{CN})_2, 2\text{AgCN}$.

Ppt.

$\text{Cu}_2(\text{CN})_2, 6\text{AgCN}$. Sol. in excess of $\text{Cu}_2(\text{CN})_2, \text{KCN} + \text{Aq}$. (Rammelsberg.)

Cuprous sodium cyanide, $\text{Cu}_2(\text{CN})_2, 2\text{NaCN}$.

(Traube, Z. anorg. 1894, 8. 21.)

$+ 4\text{H}_2\text{O}$. Decomp. by H_2O . Sol. in excess of $\text{NaCN} + \text{Aq}$. (Grossmann, Z. anorg. 1905, 43. 96.)

$\text{Cu}_2(\text{CN})_2, \text{NaCN} + 2\text{H}_2\text{O}$. Decomp. by H_2O . (Grossmann, Z. anorg. 1905, 43. 96.)

$\text{Cu}_2(\text{CN})_2, 4\text{NaCN} + 6\text{H}_2\text{O}$. Very sol. in H_2O without decomp. (Grossmann, Z. anorg. 1905, 43. 96.)

$\text{Cu}_2(\text{CN})_2, 6\text{NaCN} + 6\text{H}_2\text{O}$. Very sol. in

H_2O without decomp. (Grossmann, Z. anorg. 1905, 43. 96.)

Cuprous strontium cyanide, $\text{Cu}_2(\text{CN})_2, \text{Sr}(\text{CN})_2 + 8\text{H}_2\text{O}$.

H_2O separates $\text{Cu}_2(\text{CN})_2$. (Grossmann, Z. anorg. 1905, 43. 103.)

Cuprous cyanide ammonia, $\text{Cu}_2(\text{CN})_2, 2\text{NH}_3$.

Nearly insol. in cold H_2O . Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$ in absence of oxygen. Insol. in alcohol and ether. Decomp. by hot H_2O and acids. (Treadwell and Girsewald, Z. anorg. 1904, 39. 87.)

Cuprocupric cyanide ammonia, $\text{Cu}_2(\text{CN})_2, \text{Cu}(\text{CN})_2, 2\text{NH}_3$.

(Malmberg, Arch. Pharm. 1898, 236. 256.) $+ \text{H}_2\text{O}$. Sl. sol. in cold, decomp. by boiling H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Dufau, A. 88. 278.)

$\text{Cu}(\text{CN})_2, \text{Cu}_2(\text{CN})_2, 3\text{NH}_3$. (Mills, Z. Ch. 1867. 545.)

Sl. decomp. by boiling H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ and can be recryst. therefrom. Insol. in alcohol and ether. Decomp. by alkalis and acids. (Treadwell and Girsewald, Z. anorg. 1904, 39. 96.)

$\text{Cu}(\text{CN})_2, \text{Cu}_2(\text{CN})_2, 4\text{NH}_3$. Insol. in cold, decomp. by hot H_2O . Sol. in NH_4OH , or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Treadwell and Girsewald, Z. anorg. 1904, 39. 92.)

$2\text{Cu}_2(\text{CN})_2, \text{Cu}(\text{CN})_2, 2\text{NH}_3$. Insol. in H_2O , alcohol and ether. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by boiling acids and alkalis. (Treadwell and Girsewald, Z. anorg. 1904, 39. 92.)

$+ \text{H}_2\text{O}$. (Monthier, J. Pharm. 11. 257.)

$\text{Cu}(\text{CN})_2, 2\text{Cu}_2(\text{CN})_2, 4\text{NH}_3$. (Hillenkamp, A. 97. 218.)

$\text{Cu}(\text{CN})_2, 2\text{Cu}_2(\text{CN})_2, 6\text{NH}_3$. (Schiff and Becchi, A. 134. 33.)

$2\text{Cu}(\text{CN})_2, \text{Cu}_2(\text{CN})_2, 2\text{NH}_3 + 3\text{H}_2\text{O}$. (Fleurent, C. R. 114. 1060.)

$2\text{Cu}(\text{CN})_2, \text{Cu}_2(\text{CN})_2, 4\text{NH}_3 + \text{H}_2\text{O}$. Correct formula for $\text{Cu}(\text{CN})_2, \text{Cu}_2(\text{CN})_2, 4\text{NH}_3$. (Bouveault, Bull. Soc. (3) 4. 641.)

Cuprous cyanide ammonium sulphocyanide, $\text{Cu}_2(\text{CN})_2, 3\text{NH}_4\text{SCN}$.

Decomp. by H_2O . (Grossmann, Z. anorg. 1903, 37. 409.)

Cupric cyanide hydrazine, $\text{Cu}(\text{CN})_2(\text{N}_2\text{H}_4)$.

Insol. in H_2O and cold dil. acids.

Sol. in warm dil. acids. (Fransen, Z. anorg. 1911, 70. 154.)

Cuprous cyanide mercuric iodide, $\text{Cu}_2(\text{CN})_2, \text{HgI}_2$.

Sol. in H_2O . (Varet, Bull. Soc. (3) 4. 484.)

Cuprous cyanide potassium sulphocyanide, $\text{Cu}_2(\text{CN})_2, 3\text{KSCN}$.

Decomp. by H_2O . (Grossmann, Z. anorg. 1903, 37. 409.)

Gold (aurous) cyanide, AuCN .

Insol. in H_2O , alcohol, or ether. Not attacked by dil., or conc. acids, even boiling *qua regia*.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$, also in soluble cyanides + Aq .

Slowly decomp. by boiling $\text{KOH} + \text{Aq}$, also by $(\text{NH}_4)_2\text{S} + \text{Aq}$.

Sol. in $\text{K}_4\text{Fe}(\text{CN})_6 + \text{Aq}$. (Bentel, Z. anorg. 1912, 78. 152.)

Gold (auric) cyanide with MCN.

See Auricyanide, M.

Gold (auroauric) mercuric cyanide auric mercuric chloride, $4\text{AuCN}, \text{Au}(\text{CN})_3, 5\text{Hg}(\text{CN})_2, 7\text{AuCl}_3, 5\text{HgCl}_2$.

(Schmidt, Ch. Z. 1896, 20. 633.)

Gold (aurous) potassium cyanide, AuCN, KCN .

Sol. in 7 pts. cold, and less than 0.5 pt. boiling H_2O . Sl. sol. in cold, and somewhat more sol. in boiling alcohol. Insol. in ether. (Himly, A. 42. 160.)

Decomp. by warm acids, even tartaric, and acetic acids.

Gold (aurous) sodium cyanide, AuCN, NaCN .

Sl. sol. in cold, more easily in hot H_2O . Sl. sol. in alcohol. (Lindbom.)

Gold (aurous) strontium cyanide, $2\text{AuCN}, \text{Sr}(\text{CN})_2 + 3\text{H}_2\text{O}$.

As the Na salt.

Gold (aurous) zinc cyanide, $2\text{AuCN}, \text{Zn}(\text{CN})_2$.

Nearly insol. in hot or cold H_2O .

Insol. in cold $\text{HCl} + \text{Aq}$.

Gold (auric) cyanide auric mercuric chloride, $\text{Au}(\text{CN})_3, \text{AuCl}_3, 2\text{HgCl}_2$.

(Schmidt, Ch. Z. 1896, 20. 633.)

Gold (auroauric) cyanide aurous mercuric chloride, $12\text{AuCN}, 3\text{Au}(\text{CN})_3, 4\text{AuCl}_3, 2\text{HgCl}_2$.

(Schmidt, Ch. Z. 1896, 20. 633.)

Gold (auroauric) cyanide mercuric chloride. $15\text{AuCN}, 2\text{Au}(\text{CN})_3, 5\text{HgCl}_2$.

(Schmidt, Ch. Z. 1896, 20. 633.)

Iridium cyanide, $\text{Ir}(\text{CN})_3$.

Insol. in H_2O . Sol. in $\text{HCN} + \text{Aq}$.

Iridium cyanide with MCN.

See Iridicyanide, M.

Lanthanum cyanide, $\text{La}(\text{CN})_3$.

Ppt. (Frerichs and Smith, B. 11. 910, 1151.)

Lead cyanide, $\text{Pb}(\text{CN})_2$.

Sl. sol. in cold, more in hot H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$, and $\text{KCN} + \text{Aq}$. Partially sol. in $\text{NH}_4\text{OH} + \text{Aq}$, and NH_4 salts + Aq . Not pptd. in presence of Na citrate.

Above compound is $2\text{PbO}, \text{Pb}(\text{CN})_2 + \text{H}_2\text{O}$. (Joannis, A. ch. (5) 26. 204.)

$2\text{PbO}, \text{Pb}(\text{CN})_2 + \text{H}_2\text{O}$. Insol. in H_2O .

Lead tungsten cyanide.

See Tungstocyanide, lead.

Lead zinc cyanide, $\text{Pb}(\text{CN})_2, 2\text{Zn}(\text{CN})_2$.

Ppt. (Rammelsberg.)

Lead cyanide chloride, $2\text{Pb}(\text{CN})_2, \text{PbCl}_2$.

Insol. in H_2O . (Grissom and Thorp, Am. Ch. J. 10. 229.)

Lithium mercuric cyanide mercuric iodide, $2\text{Li}(\text{CN})_2, \text{Hg}(\text{CN})_2, \text{HgI}_2 + 7\text{H}_2\text{O}$.

Deliquescent; sol. in H_2O . (Varet, C. R. 111. 526.)

Magnesium cyanide, $\text{Mg}(\text{CN})_2$.

Known only in aqueous solution which decomposes on evaporation. (Schulz.)

Magnesium mercuric cyanide, $2\text{Mg}(\text{CN})_2, 3\text{Hg}(\text{CN})_2 + 5\text{H}_2\text{O}$.

(Grossmann, B. 1904, 37. 4143.)

Magnesium mercuric cyanide mercuric bromide, $\text{Mg}(\text{CN})_2, \text{Hg}(\text{CN})_2, \text{HgBr}_2 + 8\text{H}_2\text{O}$.

Very sol. in H_2O . (Varet, Bull. Soc. (3) 7. 170.)

Magnesium mercuric cyanide mercuric iodide, $\text{Mg}(\text{CN})_2, \text{Hg}(\text{CN})_2, \text{HgI}_2 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Varet, Bull. Soc. (3) 7. 170.)

Magnesium platinum cyanide.

See Platinocyanide, magnesium.

Magnesium tungsten cyanide.

See Tungstocyanide, magnesium.

Manganous and manganic cyanides.

See Manganocyanhydric, and Mangani-cyanhydric acids.

Manganous strontium cyanide, $2\text{Mn}(\text{CN})_2, \text{Sr}(\text{CN})_2$.

Ppt. (Descamps.)

See also Manganocyanide, strontium.

Manganous tungsten cyanide.

See Tungstocyanide, manganous.

Manganic cyanide, with MCN.

See Manganicyanide, M.

Manganous cyanide with MCN.

See Manganocyanide, M.

Mercuric cyanide, basic, $\text{Hg}(\text{CN})_2$, HgO .

Sl. sol. in cold, moderately sol. in hot H_2O . Sol. with decomp. in KOH , KCN , or $\text{KCl} + \text{Aq}$. (Johnston.)

Decomp. by H_2O over 80° . (Holdermann, Arch. Pharm. 1906, 244. 135.)

Cold H_2O dissolves about 1%, boiling H_2O about 5%. (Borelli, Gazz. ch. it. 1908, 38. (1), 361.)

1.1% dissolves in H_2O at ord. temp. (Richard, J. Chim. Phys. (6) 18. 555.)

At 0° 1/100 mol. dissolve in 1 l. H_2O .

At 25° 1/32 " " " " "

At 90° 1/10 " " " " "

(Borelli, Gazz. ch. it. 1908, 38. (1), 361.)

1000 cc. cold H_2O dissolve 1.35g. (Holdermann, Arch. Pharm. 1906, 244. 135.)

Less sol. in cold H_2O than $\text{Hg}(\text{CN})_2$. (Pieverling, J. B. 1899, 783.)

Somewhat sol. in dil. alcohol.

Practically insol. in alcohol, ether, C_6H_6 , and all organic solvents. (Borelli, Gazz. ch. it. 1908, 38, (1), 361.)

Sol. in 110 pt. alcohol of 90° Bé. (Richard, J. Chim. Phys. (6), 18. 555.)

$3\text{Hg}(\text{CN})_2$, HgO . (Joannis, A. ch. (5) 26. 469.)

Moderately sol. in H_2O . (Barthe, J. Pharm. 1896, (6), 3. 186.)

Very sol. in hot, less sol. in cold H_2O . (Holdermann, Arch. Pharm. 1904, 242. 32.)

Easily sol. in HCl . (Joannis, A. ch. 1882, (5) 26. 511.)

$\text{Hg}(\text{CN})_2$, 3HgO . More sol. in H_2O than $\text{Hg}(\text{CN})_2$, HgO .

Mercuric cyanide, $\text{Hg}(\text{CN})_2$.

Moderately sol. in H_2O .

100 pts. $\text{Hg}(\text{CN})_2 + \text{Aq}$ sat. at 101.1° contain 35 pts. $\text{Hg}(\text{CN})_2$, or 100 pts. H_2O dissolve 53.85 pts. $\text{Hg}(\text{CN})_2$ at 101.1° . (Griffiths.)

Sol. in 8 pts. H_2O at 15° . (Abl.)

Sol. in 11 pts. cold, and 2.5 pts. boiling H_2O . (Wittstein.)

8 g. are sol. in 100 g. H_2O at -0.45° . (Guthrie, Phil. Mag. 1878, (5) 6. 40.)

100 g. H_2O dissolve 9.3 g. at 13.5° . (Timofeiev, Dissert. 1894.)

100 cc. sat. solution contain 9.3 g. at 20° . (Konowalow, J. russ. Soc. 1898, (4) 30. 367.)

Solubility in H_2O at $25^\circ = 0.44$ mol. l. (Sherrill, Z. phys. Ch. 1903, 43. 735.)

1 l. H_2O dissolves 0.3956 mol. (Hofmann and Wagner, Z. Elektrochem. 1909, 15. 444.)

100 g. H_2O dissolve 12.5 g. at 15° . (Marsh and Struthers, Chem. Soc. 1905, 87. 1879.)

100 g. H_2O dissolve 11.27 g. at 25° . Sp. gr. of solution = 1.0813. (Herz and Anders, Z. anorg. 1907, 52. 164.)

$\text{Hg}(\text{CN})_2 + \text{Aq}$ containing 7.23% $\text{Hg}(\text{CN})_2$ has sp. gr. $20^\circ/20^\circ = 1.0572$.

$\text{Hg}(\text{CN})_2 + \text{Aq}$ containing 9.07% $\text{Hg}(\text{CN})_2$ has sp. gr. $20^\circ/20^\circ = 1.0743$.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 12. 282.)

Sp. gr. at $16^\circ/4^\circ$ of $\text{Hg}(\text{CN})_2 + \text{Aq}$ containing 7.8921% $\text{Hg}(\text{CN})_2 = 1.06376$; containing 5.4037% = 1.04246; containing 7.5009% = 1.06049. (Schönrock, Z. phys. Ch. 1893, 11. 770.)

Not decomp. by acids except hot conc. H_2SO_4 .

Sol. without decomp. in $\text{HNO}_3 + \text{Aq}$. (Berzelius.)

1 l. $\text{NH}_4\text{OH} + \text{Aq}$ (5.2% NH_3) dissolves 204.3 g. at about 25° . (Konowalow.)

Solubility in bases.

1 l. H_2O containing 0.3286 mols. KOH dissolves 0.5179 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.2350 mols. NaOH dissolves 0.4840 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.4775 mols. NaOH dissolves 0.5977 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.9475 mols. NaOH dissolves 0.79603 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.970 mols. LiOH dissolves 0.6543 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.480 mols. LiOH dissolves 0.5500 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.243 mols. LiOH dissolves 0.4840 mols. $\text{Hg}(\text{CN})_2$.

(Hofmann and Wagner, Z. Elektrochem. 1909, 15. 444.)

Solubility in $\text{KCN} + \text{Aq}$ at 25° .

Concentration of KCN Mols. per litre	Solubility of $\text{Hg}(\text{CN})_2$ Mols. per litre
0.0493	0.4855
0.0985	0.5350
0.1970	0.627.

(Sherrill, Z. phys. Ch. 1903, 43. 719.)

Solubility in $\text{Na}_2\text{CO}_3 + \text{Aq}$.

1 l. H_2O containing 0.4923 mols. Na_2CO_3 dissolves 0.4956 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.2443 mols. Na_2CO_3 dissolves 0.4464 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.1250 mols. Na_2CO_3 dissolves 0.4147 mols. $\text{Hg}(\text{CN})_2$.

1 l. H_2O containing 0.0000 mols. Na_2CO_3 dissolves 0.3952 mols. $\text{Hg}(\text{CN})_2$.

(Hofmann and Wagner, Z. Elektrochem. 1909, 15. 444.)

Solubility in $\text{KNO}_3 + \text{Aq}$ at 25° .
1 l. H_2O containing 0.9574 mols. KNO_3 dissolves 0.5383 mols. $\text{Hg}(\text{CN})_2$.
1 l. H_2O containing 0.4614 mols. KNO_3 dissolves 0.4619 mols. $\text{Hg}(\text{CN})_2$.
1 l. H_2O containing 0.0000 mols. KNO_3 dissolves 0.3956 mols. $\text{Hg}(\text{CN})_2$.
(Iofmann and Wagner, Z. Elektrochem. 1909, 15. 444.)

Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54. 674.)
Very easily sol. in liquid NH_3 . (Franklin, m. Ch. J. 1898, 20. 829.)

Solubility of $\text{Hg}(\text{CN})_2$ in ethyl alcohol at t° .

t°	% HgCl_2
0	8.3
10	8.8
20	9.25
30	9.8
40	10.3

(Timofeiev, Dissert. 1894.)

Solubility of $\text{Hg}(\text{CN})_2$ in methyl alcohol at t° .
 $\text{Hg}(\text{CN})_2 = \text{g. Hg}(\text{CN})_2$ in 100 g. of the solution.

t°	$\text{Hg}(\text{CN})_2$
0.0	26.10
14.7	29.17
23.4	32.01
27.4	31.77
31.7	32.53
38.1	33.29
44.5	34.05

(Dukelski, Z. anorg. 1907, 53. 337.)

100 pts. methyl alcohol dissolve 44.2 pts. $\text{Hg}(\text{CN})_2$ at 19.5° ; 100 pts. ethyl alcohol dissolve 2.09 pts. at 19.5° . (de Bruyn, Z. phys. Ch. 1892, 10. 784.)
Sol. in 2.5 pts. methyl alcohol at 14° ; in 10 pts. ethyl alcohol at 15° . (Marsh, Chem. Soc. 1905, 87. 1878.)

Solubility of $\text{Hg}(\text{CN})_2$ in methyl alcohol + Aq at 25° .

P = g. alcohol in 100 g. alcohol + Aq.
 $\text{Hg}(\text{CN})_2 = \text{millimols. Hg}(\text{CN})_2$ in 10 cc. of the solution.

P	$\text{Hg}(\text{CN})_2$	Sp gr.
0	4.34	1.0813
10.60	4.37	1.0642
30.77	4.94	1.0484
37.21	5.40	1.0430
47.06	6.49	1.0426
64.00	8.13	1.0441
78.05	9.75	1.0484
100	13.60	1.0762

Herz and Anders, Z. anorg. 1907, 52. 165.)

Solubility of $\text{Hg}(\text{CN})_2$ in ethyl alcohol + Aq at 25° .

P = g. alcohol in 100 g. alcohol + Aq.
 $\text{Hg}(\text{CN})_2 = \text{millimols Hg}(\text{CN})_2$ in 10 cc. of the solution.

P	$\text{Hg}(\text{CN})_2$	Sp. gr.
0	4.34	1.0813
20.18	3.47	1.0339
40.69	3.58	1.0006
70.01	3.80	0.9419
100	3.25	0.8552

(Herz and Anders, l. c.)

Solubility of $\text{Hg}(\text{CN})_2$ in mixtures of methyl and ethyl alcohol at 25° .

P = % methyl alcohol in the solvent.
 $\text{Hg}(\text{CN})_2 = \text{g. Hg}(\text{CN})_2$ in 10 ccm. of the solution.
 $S\ 25^\circ/4^\circ = \text{Sp. gr. of the sat. solution.}$

P	$\text{Hg}(\text{CN})_2$	S $25^\circ/4^\circ$
0	0.819	0.8552
4.37	0.902	0.8618
10.4	1.01	0.8707
41.02	1.67	0.9267
80.69	2.82	1.024
84.77	2.96	1.034
91.25	3.09	1.052
100	3.43	1.076

(Herz and Kuhn, Z. anorg. 1908, 58. 166.)

100 g. propyl alcohol dissolve 3.79 g. $\text{Hg}(\text{CN})_2$ at 13.5° . (Timofeiev, Dissert. 1894.)

Solubility in mixtures of propyl and methyl alcohol at 25° .

P = % propyl alcohol in the solvent.
G = g. $\text{Hg}(\text{CN})_2$ in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S $25^\circ/4^\circ$
0	3.43	1.0760
11.11	2.952	1.0327
23.8	2.448	0.9891
65.2	1.048	0.8800
91.8	0.504	0.8376
93.97	0.423	0.8335
96.6	0.398	0.8322
100	0.344	0.8283

(Herz and Kuhn, Z. anorg. 1908, 60.158.)

Solubility in mixtures of propyl and ethyl alcohol at 25°.

P = % propyl alcohol in the solvent.

G = g. Hg(CN)₂ in 10 ccm. of the solution.

S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	0.819	0.8552
8.1	0.790	0.8549
17.85	0.730	0.8527
56.6	0.521	0.8386
88.6	0.387	0.8311
91.2	0.384	0.8306
95.2	0.364	0.8293
100	0.344	0.8283

(Herz and Kuhn, l. c.)

Sp. gr. at 16°/4° of Hg(CN)₂+alcohol, containing 8.2206 %Hg(CN)₂=0.85273; containing 5.8652% =0.8348+.

Sp. gr. of 16°/4° of Hg(CN)₂+ pyridine containing 29.6018% Hg(CN)₂=1.28155; containing 23.2275% =1.20198.

(Schönrock, Z. phys. Ch. 1893, 11. 771.)

1 l. ether dissolves 0.01 mol. at 25°. (Sherrill, Z. phys. Ch. 1903, 43. 735.)

Easily sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 84.)

100 g. glycerol dissolve 27 g. Hg(CN)₂ at 15.5°. (Ossendowski, Pharm. J. 1907, 79. 575.)

Nearly insol. in C₆H₆. (Sherrill, Z. phys. Ch. 1903, 43. 735.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

100 g. boiling methyl acetate dissolve 3.2 g. (Steiner, Dissert, 1906.)

Solubility of Hg(CN)₂ in ethyl acetate+Aq at 25°.

P=g. ethyl acetate in 100 g. ethyl acetate +Aq.

Hg(CN)₂=millimols Hg(CN)₂ in 10 cc. of the solution.

P	Hg(CN) ₂	Sp. gr.
0	4.34	1.0810
4.39	4.295	1.0797
96.76	1.056	1.9374
100	0.714	0.09097

(Herz and Anders, Z. anorg. 1907, 52. 165.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Solubility in organic solvents at 18-20°.

100 g. tetrachlormethane dissolve 0.001 g. Hg(CN)₂.

100 g. bromoform dissolve 0.005 g. Hg(CN)₂.

100 g. ethyl bromide dissolve 0.013 g. Hg(CN)₂.

100 g. ethylene dibromide dissolve 0.001 g. Hg(CN)₂.

(Sule, Z. anorg. 1900, 25. 401.)

100 g. acetonitrile dissolve 9.58 g. Hg(CN)₂ at 18°. (Naumann and Schier, B. 1914, 249.)

Solubility in benzonitrile at 18°=1.00 in 100 g. (Naumann, B. 1914, 47. 1370.)

Sl. sol. in ethyl amine. (Shinn, J. p Chem. 1907, 11. 538.)

Very sol. in liquid methyl amine. (G J. Am. Chem. Soc. 1906, 28. 1419.)

Sol. in paratoluidine. (Werner, Z. anorg. 1897, 15. 7.)

Mol. weight determined in pyridine benzonitrile. (Werner, Z. anorg. 1897, 20 and 32.)

100 g. pyridine dissolve 64.8 g. Hg(CN)₂ at 18°. (Schroeder, Z. anorg. 1905, 44. 1370.)

Solubility in pyridine.

Mols. per 100 Hg(CN) ₂	Temp. of Solidification	Mols. per 100 Hg(CN) ₂	Temp. of Solidification
7.1	9	22.9	45
8.7	11	23.7	46
10.1	12.3	25.3	53
10.4	12.2	26.0	54
11.3	13	26.6	56
12.9	13.5	27.5	68
13.8	14.5	27.7	70
15.8	16.5	29.0	86
15.9	20.5	32.0	111
17.3	22.5	33.8	122
18.4	28.5	34.4	125
19.3	32	38.3	141
20.6	38
22.3	42

(Staronka, Ans. Ak. Wiss. Krakau, 1907, 372.)

Solubility in quinoline.

Mols. per 100 Hg(CN) ₂	Temp of Solidification	Mols. per 100 Hg(CN) ₂	Temp of Solidification
4.2	45°	13.2	13
6.0	54	17.4	16
8.2	89(61)	22.5	19
9.2	99(61)	27.1	19

(Staronka, l. c.)

Solubility in aniline.

Mols. per 100 Hg(CN) ₂	Temp. of Solidification	Mols. per 100 Hg(CN) ₂	Temp. of Solidification
3.7	14.2	77°
4.9	26° (?)	18.2	83
5.7	30.5(?)	19.7	84
7.7	35 (?)	23.4	88
9.2	38.5(?)

(Staronka, l. c.)

Mercuric nickel cyanide ammonia, 2Hg(CN)₂.4Ni(CN)₂.5NH₃.+2H₂O.

(Papiermeister, Dissert. 1898.)

5Hg(CN)₂.18Ni(CN)₂.8NH₃.+15H₂O (Papiermeister, Dissert. 1898.)

Potassium cyanide, $\text{Hg}(\text{CN})_2$,

pts. cold H_2O ; sl. sol. in alcohol;
acids.

100 dissolve 22.7 g. (Fronmüller,
92.)

Very sol. in liquid NH_3 . (Frank-
Ch. 1909, 69. 295.)

**Mercuric cyanide, basic, $\text{Hg}(\text{CN})_2$,
 AgCN .**

Exam, B. 16. 2669.)

**Mercuric cyanide mercuric sulphate,
 $\text{Hg}(\text{CN})_2$, 2AgCN , $\text{HgSO}_4 + \text{H}_2\text{O}$.**

**Lithium cyanide, $\text{Hg}(\text{CN})_2$, NaCN
 O .**

O . (Grossmann, B. 1904, 37.

**Strontium cyanide,
 $\text{Sr}(\text{CN})_2 + 5\text{H}_2\text{O}$.**

Microscopic. Sol. in H_2O . (Gross-
M., 37. 4142.)

**Strontium cyanide iodide, $\text{Sr}(\text{CN})_2$,
 $\text{Hg}(\text{CN})_2 + 7\text{H}_2\text{O}$.**

R. 1895, 121. 499.)

**Thallium cyanide, $\text{Hg}(\text{CN})_2$, 2TlCN .
in H_2O . 100 pts. H_2O dissolve
, and 10.3 pts. at 10° . (Fron-
. 92.)**

**Zinc cyanide, $4\text{Zn}(\text{CN})_2$, $\text{Hg}(\text{CN})_2$.
 I_2O . (Dunstan, Chem. Soc. 6.**

**Mercuric cyanide mercuric bromide,
 $\text{Hg}(\text{CN})_2$, $\text{Zn}(\text{CN})_2$, HgBr_2 ,**

Very H_2O . Sl. sol. in cold NH_4OH
t, C. R. 1889, 109. 810.)

**Cyanide ammonia, $\text{Hg}(\text{CN})_2$, NH_3 .
 H_2O , $\text{NH}_4\text{OH} + \text{Aq}$, and alcohol.
1889, 109. 903.)**

H_2O . (Schmidt, B. 1894, 27.

$2\text{NH}_3 + \frac{1}{2}\text{H}_2\text{O}$. Easily de-
t, Bull. Soc. (3) 6. 221.)

Cyanide bromide, $\text{Hg}(\text{CN})_2$,

. even in boiling H_2O . (Prussia,
1898, 28, (2), 114.)

**Cyanide barium bromide, $2\text{Hg}(\text{CN})_2$,
 $5\text{H}_2\text{O}$.**

Especially in hot H_2O and al-
t, C. R. 1895, 121. 398.)

**Mercuric cyanide cadmium bromide,
 $\text{Hg}(\text{CN})_2$, $\text{CdBr}_2 + 3\text{H}_2\text{O}$.**

Sol. in H_2O and $\text{NH}_4\text{OH} + \text{Aq}$. (Varet,
Bull. Soc. (3) 5. 8.)

$2\text{Hg}(\text{CN})_2$, $\text{CdBr}_2 + 4.5 \text{H}_2\text{O}$. Sol. in H_2O
and in $\text{NH}_4\text{OH} + \text{Aq}$. (Varet, C. R. 1890,
111. 680.)

**Mercuric cyanide cadmium bromide ammonia
 $2\text{Hg}(\text{CN})_2$, CdBr_2 , $4\text{NH}_3 + 2\text{H}_2\text{O}$.**

Decomp. by H_2O .

Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Varet, C. R.
1891, 112. 535.)

**Mercuric cyanide calcium bromide,
 $2\text{Hg}(\text{CN})_2$, $\text{CaBr}_2 + 5\text{H}_2\text{O}$.**

Sol. in 1 pt. cold, and 0.25 pt. boiling H_2O ;
also in 2 pts. cold, and 1 pt. boiling 90%
alcohol. (Custer.)

$+7\text{H}_2\text{O}$. (Varet, C. R. 1895, 121. 399.)

**Mercuric cyanide cupric bromide ammonia,
 $2\text{Hg}(\text{CN})_2$, CuBr_2 , 4NH_3 .**

Decomp. by H_2O ; sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
(Varet, Bull. Soc. (3) 6. 221.)

**Mercuric cyanide lithium bromide, $2\text{Hg}(\text{CN})_2$,
 $2\text{LiBr} + 7\text{H}_2\text{O}$.**

Deliquescent. (Varet, C. R. 111. 526.)

Mercuric cyanide magnesium bromide.

See Magnesium mercuric cyanide mercuric
bromide.

$\text{Hg}(\text{CN})_2$, 2KBr . Very sol. in H_2O .
(Harth, Z. anorg. 1897, 14. 351.)

**Mercuric cyanide potassium bromide,
 $\text{Hg}(\text{CN})_2$, $\text{KBr} + 2\text{H}_2\text{O}$.**

Sol. in 13.34 pts. H_2O at 18° , and less than
1 pt. boiling H_2O . (Brett.)

Sol. without decomp. in hot dil. H_2SO_4 ,
 HNO_3 , or $\text{HCl} + \text{Aq}$. (Brett.)

Contains $1\frac{1}{2}\text{H}_2\text{O}$. (Berthelot, A. ch. (5)
29. 226.)

**Mercuric cyanide sodium bromide, $\text{Hg}(\text{CN})_2$,
 $\text{NaBr} + 1\frac{1}{2}\text{H}_2\text{O}$.**

Sol. in H_2O and alcohol.

**Mercuric cyanide strontium bromide,
 $2\text{Hg}(\text{CN})_2$, $\text{SrBr}_2 + 6\text{H}_2\text{O}$.**

Sol. in H_2O and in alcohol. (Varet, C. R.
1895, 121. 399.)

**Mercuric cyanide zinc bromide, HgBr_2 ,
 $\text{Hg}(\text{CN})_2$, $\text{Zn}(\text{CN})_2 + 8\text{H}_2\text{O}$.**

Sol. in H_2O and $\text{NH}_4\text{OH} + \text{Aq}$. (Varet,
Bull. Soc. (3) 5. 8.)

**Mercuric cyanide zinc bromide ammonia,
 HgBr_2 , $\text{Hg}(\text{CN})_2$, $\text{Zn}(\text{CN})_2$, 4NH_3 .**

As the corresponding chloride. (Varet.)

Mercuric cyanide chloride, $\text{Hg}(\text{CN})_2, \text{HgCl}_2$.
Sol. in H_2O . Decomp. by alcohol, which dissolves out HgCl_2 .

Mercuric cyanide ammonium chloride,
 $\text{Hg}(\text{CN})_2, \text{NH}_4\text{Cl}$.

Sol. in H_2O and alcohol. (Poggiale.)
 $\text{Hg}(\text{CN})_2, 4\text{NH}_4\text{Cl}$.

Mercuric cyanide barium chloride, $2\text{Hg}(\text{CN})_2, \text{BaCl}_2 + 4\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O and alcohol
 $+ 6\text{H}_2\text{O}$. (Dexter.)

Mercuric cyanide barium chloride ammonia,
 $2\text{Hg}(\text{CN})_2, \text{BaCl}_2, 4\text{NH}_3$.

Decomp. by H_2O . Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
(Varet, Bull. Soc. (3) 6. 221.)

Mercuric cyanide cadmium chloride,
 $\text{Hg}(\text{CN})_2, \text{CdCl}_2 + 2\text{H}_2\text{O}$.

Sol. in H_2O and $\text{NH}_4\text{OH} + \text{Aq}$. (Varet,
Bull. Soc. (3) 5. 8.)

Mercuric cyanide calcium chloride,
 $2\text{Hg}(\text{CN})_2, \text{CaCl}_2 + 6\text{H}_2\text{O}$.

Efflorescent. Very sol. in H_2O . (Varet,
C. R. 1895, 121. 349.)

Mercuric cyanide cerium chloride, $3\text{Hg}(\text{CN})_2, \text{CeCl}_3 + 8\text{H}_2\text{O}$.

Very sol. in H_2O . (Ahlén, Bull. Soc. (2)
27. 365.)

Mercuric cyanide cobaltous chloride,
 $\text{Hg}(\text{CN})_2, 2\text{CoCl}_2 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Poggiale.)

$2\text{Hg}(\text{CN})_2, \text{CoCl}_2 + 7\text{H}_2\text{O}$. (Dexter.)

Mercuric cyanide cupric chloride, $\text{Hg}(\text{CN})_2, \text{CuCl}_2 + 6\text{H}_2\text{O}$.

Efflorescent.

Sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq}$. (Varet,
C. R. 1888, 107. 1002.)

$2\text{Hg}(\text{CN})_2, \text{CuCl}_2 + 6\text{H}_2\text{O}$. Efflorescent.

Very sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq}$.
(Varet, C. R. 1888, 107. 1002.)

Mercuric cyanide cupric chloride ammonia,
 $2\text{Hg}(\text{CN})_2, \text{CuCl}_2, 4\text{NH}_3$.

Decomp. by H_2O . Sl. sol. in cold NH_4OH
 $+ \text{Aq}$. (Varet, Bull. Soc. (3) 6. 221.)

Mercuric cyanide didymium chloride,
 $3\text{Hg}(\text{CN})_2, \text{DiCl}_3 + 8\text{H}_2\text{O}$.

Very sol. in H_2O . (Ahlén.)

Mercuric cyanide erbium chloride, $3\text{Hg}(\text{CN})_2, \text{ErCl}_3 + 8\text{H}_2\text{O}$.

Easily sol. in H_2O . (Ahlén.)

Mercuric cyanide hydrazine chloride,
 $\text{Hg}(\text{CN})_2, \text{N}_2\text{H}_4, \text{HCl}$.

Very sol. in H_2O .

Nearly insol. in alcohol and ether
ratini, Gazz. ch. it. 1912, 42. (1), 154

Mercuric cyanide ferric chloride, $2\text{Hg}(\text{CN})_2, \text{FeCl}_3 + 3\frac{1}{2}\text{H}_2\text{O}$.

(Dexter.)

Mercuric cyanide lanthanum chlor
 $3\text{Hg}(\text{CN})_2, \text{LaCl}_3 + 8\text{H}_2\text{O}$.

Very sol. in H_2O . (Ahlén.)

Mercuric cyanide magnesium chlor
 $2\text{Hg}(\text{CN})_2, \text{MgCl}_2 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O and dil. alcohol.
(Poggiale.)

Mercuric cyanide manganous
 $\text{Hg}(\text{CN})_2, \text{MnCl}_2 + 3\text{H}_2\text{O}$.

Efflorescent. Very sol. in H_2O . (Poggiale.)

Mercuric cyanide nickel chloride, $\text{Hg}(\text{CN})_2, \text{NiCl}_2 + 6\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . (Poggiale.)
 $2\text{Hg}(\text{CN})_2, \text{NiCl}_2 + 7\text{H}_2\text{O}$. (Dexter.)

Mercuric cyanide chloride nickel oxychloride, $11\text{Hg}(\text{CN})_2, 8\text{HgCl}_2, 8\text{Ni}(\text{OH})\text{Cl} + 76\text{H}_2\text{O}$.

(Papiermeister, Dissert. 1896.)

Mercuric cyanide potassium chlor
 $\text{Hg}(\text{CN})_2, \text{KCl} + \text{H}_2\text{O}$.

Sol. in 6.75 pts. H_2O at 18° .
Sol. in alcohol.

Mercuric cyanide sodium chloride, $\text{Hg}(\text{CN})_2, \text{NaCl}$.

Easily sol. especially in hot H_2O ;
alcohol. (Poggiale.)

Mercuric cyanide strontium chlor
 $2\text{Hg}(\text{CN})_2, \text{SrCl}_2 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O and dil. alcohol.
C. R. 1895, 121. 349.)

Mercuric cyanide yttrium chloride, $3\text{Hg}(\text{CN})_2, \text{YCl}_3 + 8\text{H}_2\text{O}$.

Easily sol. in H_2O . (Ahlén, Bull.
27. 365.)

Mercuric cyanide zinc chloride, $2\text{Hg}(\text{CN})_2, \text{ZnCl}_2 + 6\text{H}_2\text{O}$.

Efflorescent. Sol. in H_2O . (Kahlbaum,
 $\text{HgCl}_2, \text{Hg}(\text{CN})_2, \text{Zn}(\text{CN})_2 + 7\text{H}_2\text{O}$,
efflorescent. Very sol. in H_2O . (Varet,
Bull. Soc. (3) 5. 8.)

Mercuric cyanide zinc chloride a
 $\text{HgCl}_2, \text{Hg}(\text{CN})_2, \text{ZnCl}_2, 4\text{NH}_3$.

Decomp. by H_2O . Sol. in NH_4OH .
(Varet, Bull. Soc. (3) 6. 221.)

$\text{Hg}(\text{CN})_2, \text{Zn}(\text{CN})_2, \text{HgCl}_2, 6\text{NH}_3$.
C. R. 106. 1080.)

cyanide potassium chromate.
chrome mercuric cyanide, potassium.

cyanide potassium ferrocyanide,
 $\text{K}_4\text{Fe}(\text{CN})_6 + 4\text{H}_2\text{O}$.
 sol. in H_2O .

cyanide hydrazine, $\text{Hg}(\text{CN})_2$,
 sol. in H_2O with partial decomp.
 and Marburg, A. 1899, 305. 215.)
 N_2H_4 . Ppt. (Franzen, Z. anorg.
 154.)

cyanide potassium hydroxide,
 N_2 , KOH .
 and Wagner, B. 1908, 41. 321.)
 H_2O . (Hofmann and Wagner, B.
 1630.)
 N_2 , $\text{KOH} + \text{H}_2\text{O}$. Very sol. in H_2O .
 and Wagner, B. 1908, 41. 320.)

cyanide sodium hydroxide,
 N_2 , $\text{NaOH} + 1\frac{1}{2}\text{H}_2\text{O}$ or H_2O .
 and Wagner, B. 1908, 41. 1631.)

cyanide barium iodide, $2\text{Hg}(\text{CN})_2$,
 $+4\text{H}_2\text{O}$.
 deliquescent. Sol. in 16.5 pts. cold,
 at boiling H_2O . Sol. in 22.5 pts.
 1.6 pts. hot 90% alcohol. Solution
 on boiling. (Custer.)

cyanide cadmium iodide, $\text{Hg}(\text{CN})_2$,
 N_2 , $\text{HgI}_2 + 8\text{H}_2\text{O}$.
cadmium mercuric cyanide mercuric

cyanide caesium iodide, $\text{Hg}(\text{CN})_2$,
 t. from H_2O without decomp.
 p. by acids. (Mathewson and
 n. Ch. J. 1903, 30. 433.)

cyanide calcium iodide, $2\text{Hg}(\text{CN})_2$,
 $+6\text{H}_2\text{O}$.
 fluorescent. More sol. in H_2O than
 corresponding Sr. comp. (Custer.)

cyanide lithium iodide, $\text{Hg}(\text{CN})_2$,
 CN_2 , $\text{HgI}_2 + 7\text{H}_2\text{O}$.
cyanide, lithium mercuric mercuric

cyanide magnesium iodide,
 CN_2 , $\text{Mg}(\text{CN})_2$, $\text{HgI}_2 + 8\text{H}_2\text{O}$.
cyanide, magnesium mercuric mercuric

cyanide potassium iodide, $\text{Hg}(\text{CN})_2$,
 16 pts. cold, and less hot H_2O . Sol.
 in cold alcohol of 34° Baumé. (Cail-
 sol. in ether. Decomp. by acids.
 N_2 , $2\text{KI} + \frac{1}{2}\text{H}_2\text{O}$. (Berthelot.)

Mercuric cyanide sodium iodide, $\text{Hg}(\text{CN})_2$,
 $\text{NaI} + 2\text{H}_2\text{O}$.

Sol. in $4\frac{1}{2}$ pts. H_2O at 18°, and $\frac{6}{7}$ pt.
 boiling H_2O .

Sol. in 2 pts. boiling, and $6\frac{1}{2}$ pts. cold
 90% alcohol. (Custer.)

Mercuric cyanide strontium iodide,
 $2\text{Hg}(\text{CN})_2$, $\text{SrI}_2 + 6\text{H}_2\text{O}$.

Sol. in 7 pts. H_2O at 18°, and $\frac{1}{2}$ pt. at b.-pt.
 Sol. in 4 pts. 90% alcohol at 18°, and $\frac{1}{2}$ pt.
 at b.-pt. (Custer.)

Mercuric cyanide zinc iodide, $2\text{Hg}(\text{CN})_2$,
 $\text{ZnI}_2 + 6\text{H}_2\text{O}$.

Efflorescent; sol. in H_2O .

Mercuric cyanide iodide potassium cyanide,
 HgI_2 , $\text{Hg}(\text{CN})_2$, 2KCN .

Easily decomp. by dil. acids. (Rupp.
 Apoth. Ztg., 23. 374.)

Mercuric cyanide cadmium nitrate,
 $2\text{Hg}(\text{CN})_2$, $\text{Cd}(\text{NO}_3)_2 + 7\text{H}_2\text{O}$.

Decomp. by H_2O , not by alcohol. (Ny-
 lander, J. B. 1859 271.)

Mercuric cyanide cobalt nitrate, $2\text{Hg}(\text{CN})_2$,
 $\text{Co}(\text{NO}_3)_2 + 7\text{H}_2\text{O}$.

Decomp. by H_2O , not by alcohol. (Ny-
 lander.)

Mercuric cyanide copper nitrate, $\text{Hg}(\text{CN})_2$,
 $\text{Cu}(\text{NO}_3)_2 + 5\text{H}_2\text{O}$.

Decomp. by H_2O , not by alcohol. (Ny-
 lander.)

Mercuric cyanide ferrous nitrate, $2\text{Hg}(\text{CN})_2$,
 $\text{Fe}(\text{NO}_3)_2 + 7\text{H}_2\text{O}$.

Decomp. by H_2O , not by alcohol. (Ny-
 lander.)

Mercuric cyanide manganous nitrate,
 $\text{Hg}(\text{CN})_2$, $\text{Mn}(\text{NO}_3)_2 + 5\text{H}_2\text{O}$.

Decomp. by H_2O , not by alcohol. (Ny-
 lander.)

$2\text{Hg}(\text{CN})_2$, $\text{Mn}(\text{NO}_3)_2 + 7\text{H}_2\text{O}$. As above.

Mercuric cyanide nickel nitrate, $2\text{Hg}(\text{CN})_2$,
 $\text{Ni}(\text{NO}_3)_2 + 7\text{H}_2\text{O}$.

Decomp. by H_2O , not by alcohol. (Ny-
 lander.)

Mercuric cyanide silver nitrate, $2\text{Hg}(\text{CN})_2$,
 $\text{AgNO}_3 + 2\text{H}_2\text{O}$.

Sl. sol. in cold, more readily in hot H_2O .
 Sol. with decomp. in $\text{HNO}_3 + \text{Aq}$.

As sol. in alcohol as in H_2O .

Mercuric cyanide zinc nitrate, $2\text{Hg}(\text{CN})_2$,
 $\text{Zn}(\text{NO}_3)_2 + 7\text{H}_2\text{O}$.

Sol. in H_2O with decomp. Not decomp. by
 alcohol. (Nylander, J. B. 1859. 271.)

- Mercuric cyanide nitrate silver cyanide, basic,**
 $\text{Hg}(\text{NO}_3)\text{CN}$, 10AgCN , $\text{Hg}(\text{OH})\text{NO}_3$.
 (Schmidt, Z. anorg. 1895, 9. 431.)
- Mercuric cyanide potassium selenocyanide,**
 $\text{Hg}(\text{CN})_2$, KSeCN .
 Sl. sol. in cold, much more easily sol. in hot H_2O or alcohol. Traces dissolve in ether.
 (Cameron and Davy, C. N. 44. 63.)
- Mercuric cyanide nickel sulphate,**
 $\text{Hg}(\text{CN})_2$, $\text{NiSO}_4 + 9\text{H}_2\text{O}$.
 (Papiermeister, Dissert. 1898.)
- Mercuric cyanide ammonium sulphocyanide,**
 $\text{Hg}(\text{CN})_2$, NH_4SCN .
 Easily sol. in hot H_2O . (Cleve, Bull. Soc. (2) 23. 71.)
- Mercuric cyanide barium sulphocyanide,**
 $2\text{Hg}(\text{CN})_2$, $\text{Ba}(\text{SCN})_2 + 4\text{H}_2\text{O}$.
 Permanent. Sol. in hot H_2O . (Cleve.)
- Mercuric cyanide cadmium sulphocyanide,**
 $2\text{Hg}(\text{CN})_2$, $\text{Cd}(\text{SCN})_2 + 4\text{H}_2\text{O}$.
 Permanent. Sol. in hot H_2O . (Cleve.)
- Mercuric cyanide calcium sulphocyanide,**
 $2\text{Hg}(\text{CN})_2$, $\text{Ca}(\text{SCN})_2 + 8\text{H}_2\text{O}$.
 Sol. in H_2O . (Cleve.)
- Mercuric cyanide cerium sulphocyanide,**
 $3\text{Hg}(\text{CN})_2$, $\text{Ce}(\text{SCN})_3 + 12\text{H}_2\text{O}$.
 Easily sol. in hot H_2O . (Jolin.)
- Mercuric cyanide didymium sulphocyanide,**
 $3\text{Hg}(\text{CN})_2$, $\text{Di}(\text{SCN})_3 + 6\text{H}_2\text{O}$.
 Sl. sol. in cold, easily in hot H_2O . (Cleve.)
- Mercuric cyanide erbium sulphocyanide,**
 $3\text{Hg}(\text{CN})_2$, $2\text{Er}(\text{SCN})_3 + 12\text{H}_2\text{O}$.
 Sl. sol. in cold, easily in hot H_2O . (Cleve.)
- Mercuric cyanide lanthanum sulphocyanide,**
 $3\text{Hg}(\text{CN})_2$, $\text{La}(\text{SCN})_3 + 12\text{H}_2\text{O}$.
 Very sol. in H_2O . (Cleve.)
- Mercuric cyanide magnesium sulphocyanide,**
 $2\text{Hg}(\text{CN})_2$, $\text{Mg}(\text{SCN})_2 + 4\text{H}_2\text{O}$.
 Permanent. Easily sol. in hot H_2O .
 (Cleve.)
- Mercuric cyanide potassium sulphocyanide,**
 $\text{Hg}(\text{CN})_2$, KSCN .
 Permanent. Easily sol. in hot H_2O .
 (Cleve.)
 $+ 2\text{H}_2\text{O}$. (Philip, Z. Ch. 1867. 552.)
- Mercuric cyanide rubidium sulphocyanide,**
 $\text{Hg}(\text{CN})_2$, $\text{Rb}(\text{SCN})$.
 Sol. in hot H_2O without decomp. (Grossmann, B. 1904, 37. 1259.)
- Mercuric cyanide samarium sulphocyanide,**
 $3\text{Hg}(\text{CN})_2$, $\text{Sm}(\text{SCN})_3 + 12\text{H}_2\text{O}$.
 Easily sol. in H_2O . (Cleve.)
- Mercuric cyanide sodium sulphocyanide,**
 $\text{Hg}(\text{CN})_2$, $\text{NaSCN} + 2\text{H}_2\text{O}$.
 Efflorescent. Sol. in H_2O . (Cleve, Soc. (2) 23. 71.)
- Mercuric cyanide strontium sulphocyanide,**
 $2\text{Hg}(\text{CN})_2$, $\text{Sr}(\text{SCN})_2 + 4\text{H}_2\text{O}$.
 Efflorescent. (Cleve.)
- Mercuric cyanide yttrium sulphocyanide,**
 $3\text{Hg}(\text{CN})_2$, $\text{Y}(\text{SCN})_3 + 12\text{H}_2\text{O}$.
 Sl. sol. in warm, much less in cold
 (Cleve.)
- Mercuric cyanide zinc sulphocyanide,**
 $2\text{Hg}(\text{CN})_2$, $\text{Zn}(\text{SCN})_2 + 4\text{H}_2\text{O}$.
 Sl. sol. in H_2O . (Cleve.)
- Mercuric cyanide zinc sulphocyanide monia,**
 $2\text{Hg}(\text{CN})_2$, $\text{Zn}(\text{SCN})_2$, 3N
 Not efflorescent. Decomp. by H_2O .
- Mercuric cyanide potassium thiosulphate,**
 $\text{Hg}(\text{CN})_2$, $\text{K}_2\text{S}_2\text{O}_3$.
 Permanent. Sol. in H_2O . (Kessler.)
 $+ \text{H}_2\text{O}$. (Fock and Klüss, B. 24. 134.)
- Molybdenum hydroxyl potassium cyanide,**
 $\text{K}_2\text{Mo}(\text{OH})_2(\text{CN})_2$.
 (Rosenheim and Koss, Z. anorg. 1901, 155.)
 $\text{K}_2\text{Mo}(\text{OH})_2(\text{CN})_2$. Very sol. in
 (Rosenheim and Koss.)
- Molybdenum cyanide with MCN.**
 See Molybdocyanide M.
- Molybdenyl potassium cyanide,**
 $\text{MoO}_2(\text{CN})_2$, 2KCN .
 Very sol. in H_2O . Aqueous solution stable in presence of alkalis.
 Insol. in alcohol. (Péchar, C. R. 118. 805.)
 $\text{MoO}_2(\text{CN})_2$, 3KCN . Sol. in H_2O . Insol. in alcohol. (Hofmann, Z. anorg. 1896, 287.)
 $+ \text{H}_2\text{O}$. Sol. in H_2O . Insol. in alk. (Hofmann.)
 $+ 4\text{H}_2\text{O}$. Sol. in H_2O . Insol. in alk. (Hofmann.)
- Nickel cyanide, $\text{Ni}(\text{CN})_2 + x\text{H}_2\text{O}$.**
 Insol. in H_2O . Insol. in conc. HCl , H or $\text{HNO}_3 + \text{Aq}$, but decomp. by heating with. Sol. in NH_4OH , warm (NH_4 or NH_4 succinate $+ \text{Aq}$; also in KCN Sl. sol. in NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (stein.)
 Insol. in methyl acetate. (Nauman 1909, 42. 3790.)

$\text{Ni(CN)}_2 \cdot 3\frac{1}{4}\text{H}_2\text{O} + 3\frac{3}{4}\text{H}_2\text{O} + 4\frac{1}{4}\text{H}_2\text{O}$, $\frac{1}{4}\text{H}_2\text{O}$. (Papiermeister, Dissert.

1. (Hofmann and Höchtlen, B. 1149.)

potassium cyanide, $\text{Ni(CN)}_2 \cdot 2\text{KCN}$

O . Decomp. by acids with residue Ni(CN)_2 .

O . (Rammelsberg.)

potassium cyanide, $\text{Ni(CN)}_2 \cdot 2\text{NaCN} +$

O ; decomp. by acids with residue Ni(CN)_2 .

potassium cyanide, $\text{Ni(CN)}_2 \cdot \text{Sr(CN)}_2 \cdot \text{O}$.

O . (Handl, J. B. 1859. 273.)

potassium cyanide ammonia, $\text{Ni(CN)}_2 \cdot \text{NH}_3 +$

is attacked by H_2O or dil. acids. Conc. H_2SO_4 . Sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{NH}_4\text{OH} + \text{Aq}$. $(\text{NH}_4)_2\text{C}_2\text{O}_4 + \text{Aq}$, Aq , and $\text{KCN} + \text{Aq}$. Decomp. with NaOH or KOH . (Bernoulli, Ch. Z. 1901, 25. 436.)

potassium cyanide trihydrazine, $\text{Ni(CN)}_2 \cdot 3\text{N}_2\text{H}_4$.

vanzen, Z. anorg. 1911, 70. 155.)

potassium cyanide, $\text{Os(CN)}_2(?)$.

H_2O ; not attacked by acids.

osmium cyanhydric acid.

potassium cyanide.

rhodocyanide, potassium.

rhodium cyanide, Pd(CN)_2 .

in H_2O . Insol. in dil. acids. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$, also in conc. HCN .

potassium cyanide, Pt(CN)_2 .

H_2O , alkalis, or acids. Sol. in H_2O . When freshly pptd., sol. in H_2O .

potassium cyanide with MCN.

potassium cyanide, M.

potassium cyanide, KCN .

cent. Very sol. in H_2O .

$\text{KCN} + \text{Aq}$, sat. at b.-pt. 103.3° , 100 pts. KCN , i. e. 100 pts. H_2O dis-
solves 100 pts. KCN at 103.3° . (Griffiths.)
Aq containing 3.25% KCN has
sp. gr. 1.054; 6.5% KCN , 1.0316. (Kohl-
Ann. 1879. 1.)

Aq containing 9.64% KCN has
sp. gr. $20^\circ = 1.0514$.

$\text{KCN} + \text{Aq}$ containing 14.42% KCN has
sp. gr. $20^\circ/20^\circ = 1.0768$. (Le Blanc and Roh-
land, Z. phys. ch. 1896, 19. 278.)

Moderately sol. in liquid NH_3 . (Franklin,
Am. Ch. J. 1898, 20. 829.)

Almost insol. in absolute alcohol.

Sol. in 80 pts. 95% alcohol when boiling,
and easily sol. in 35% alcohol. (Geiger, A. 1.
50.)

100 pts. absolute methyl alcohol dissolve
4.91 pts. at 19.5° ; 100 pts. absolute ethyl
alcohol dissolve 0.87 pt. at 19.5° . (de Bruyn,
Z. phys. Ch. 10. 783.)

Insol. in methyl acetate (Naumann,
B. 1909, 42. 3790); ethyl acetate. (Naumann,
B. 1904, 37. 3601.)

100 g. glycerol dissolve 32 g. KCN at 15.5° .
(Ossendowski, Pharm. J. 1907, 79. 575.)

Sol. in CS_2 when pure. (Loughlin, J. B.
1875. 234.)

Wholly insol. in CS_2 . (Moldenhauer, Z.
anal. 16. 199.)

Sl. sol. in benzonitrile. (Naumann, B.
1914, 47. 1369.)

Potassium chromium tetroxide pentacyanide,
 $\text{K}_2[(\text{CrO}_4)_2(\text{CN})_5] + 5\text{H}_2\text{O}$.

Very hygroscopic.

Sol. in H_2O . (Riesenfeld, B. 1908, 41.
3548.)

Potassium chromium tetroxide dicyanide ammonia,
 $\text{K}_2[\text{CrO}_4(\text{CN})_2\text{NH}_3] + 5\text{H}_2\text{O}$.

Hygroscopic in the air.

Easily sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq}$.
(Riesenfeld, B. 1908, 41. 3545.)

Potassium rhodium cyanide.

See Rhodocyanide, potassium.

Potassium ruthenium cyanide.

See Ruthenocyanide, potassium.

Potassium silver cyanide, KCN , AgCN .

Sol. in 4.7 pts. H_2O at 15° , 4 pts. at 20° ,
and in much less at higher temp. Sol. in 25
pts. 85% alcohol. (Baup, A. ch. (3) 53. 464.)

Potassium silver sodium cyanide, 2KCN ,
 NaCN , 3AgCN .

Sol. in 4.4 pts. H_2O at 15° , and 22 pts. 85%
alcohol at 17° . (Baup.)

Potassium tungsten cyanide.

See Tungstocyanide, potassium.

Potassium uranyl cyanide,

$(\text{UO}_2)(\text{CN})_2 \cdot 2\text{KCN}$.

Ppt. Sol. in H_2O . Sl. sol. in presence of
large excess of KCN . (Aloy, A. ch. 1901, (7)
24. 417.)

Potassium vanadium cyanide, $\text{K}_2\text{V(CN)}_6$.

Readily sol. in H_2O ; decomp. slowly in neu-
tral aq. solution, rapidly in acid aq. solution;

insol. in alcohol. (Locke, Am. Ch. J. 1898, 20. 601.)

$K_4V(CN)_6 + 3H_2O$. Sol. in H_2O .

Insol. in alcohol and ether. (Petersen, Z. anorg. 1904, 38. 345.)

Potassium zinc cyanide, $2KCN, Zn(CN)_2$.

100 pts. H_2O dissolve 11 pts. at 20° . (Sharwood, Eng. Min. J. 1904, 77. 845.)

Potassium cyanide molybdenum dioxide, $4KCN, MoO_3 + 5H_2O$.

Sol. in H_2O . Insol. in alcohol. (Hofmann, Z. anorg. 1896, 12. 287.)

$+6H_2O$. "Potassium dioxotetracyanomolybdate."

Very sol. in H_2O . (Winkler, Dissert. 1909.)

$+8H_2O$. (Rosenheim, Kohn and Garfunkel, Z. anorg. 1910, 65. 174.)

$+10H_2O$. Decomp. by conc. HCl , HNO_3 , and H_2SO_4 .

Not acted upon by cold dil. acids. (v. der Heide and Hofmann, Z. anorg. 1896, 12. 285.)

$5KCN, MoO_3 + 8H_2O$. Sol. in H_2O . Insol. in alcohol. (Kalischer, Dissert. 1902.)

Potassium cyanide molybdenum dioxide hydroxylamine, $4KCN, MoO_3, NH_2OH + H_2O$.

Sol. in H_2O .

Decomp. by dil. acids. (v. der Heide and Hofmann, Z. anorg. 1896, 12. 282.)

Potassium cyanide molybdenum sulphide, $6KCN, Mo_2S_3 + 5H_2O$.

Easily sol. in H_2O . Decomp. by dil. acids. (Hofmann, Z. anorg. 1896, 12. 289.)

Very sol. in H_2O . Slowly decomp. in the cold by dil. mineral acids. (v. der Heide and Hofmann, Z. anorg. 1896, 12. 289.)

Potassium cyanide molybdenum sulphocyanide, $2KCN, MoS_2(CN)_2$.

(Péchar, C. R. 1894, 118, 806.)

$5KCN, Mo_2S_4(CN)_2 + 7H_2O$. Sol. in H_2O . Stable toward dil. acids and alkalies. (Hofmann, Z. anorg. 1896, 12. 289.)

Potassium cyanide molybdenum sulphonyl cyanide, $4KCN, Mo_2SO(CN)_2 + 4H_2O$.

Sol. in H_2O . Stable toward dil. acids. (Hofmann, Z. anorg. 1896, 12. 289.)

Potassium cyanide nitrite, $KCN, KNO_2 + \frac{1}{2}H_2O$.

Sol. in H_2O ; decomp. slowly by H_2O ; explosive. (Hofmann, Z. anorg. 1895, 10. 260-261.)

Potassium cyanide sulphur dioxide, $KCN, SO_2 + H_2O$.

Much more sol. in hot than cold H_2O . (Étard, C. R. 88. 649.)

$KCN, HCN, 2SO_2 + 3H_2O$. Very sl. in cold H_2O ; decomp. by hot H_2O . (Étard, C. R. 88. 649.)

Rubidium tungsten cyanide.

See Tungstocyanide, rubidium.

Rhodium cyanide, $Rh(CN)_3$.

Ppt. Not decomp. by acids. So $KCN + Aq$. (Martius, A. 117. 361.)

Rhodium cyanide with 3KCN.

See Rhodicyanide, potassium.

Ruthenium cyanide with 4MCN.

See Ruthenocyanide, M.

Silver cyanide, $AgCN$.

Sl. sol. in H_2O .

$2.2 + 10^{-4}$ g. sol. in 1 liter of H_2O at 17.5° . (Böttger, Z. phys. ch. 1903, 46. 603.)

1 l. solution in H_2O contains 0.000 $AgCN$ at 17.5° . (Abegg and Cox, Z. Ch. 1903, 46. 11.)

Solubility in H_2O at $25^\circ = 2.22$ mol. per l. (Lucas, Z. anorg. 1904, 41. 1.)

Insol. in dil. acids. Decomp. by acids. Not sol. to any extent in HCN .

Freshly pptd. $AgCN$ is not dissolved by cold dil. HNO_3 , but is attacked by HNO_3 on boiling. From dry $AgCN$ solved 5% by boiling 1 hour with 1% $+ Aq$. Conc. HNO_3 dissolves more. (E. B. 1901, 34. 1605.)

Sol. in $NH_4OH + Aq$. Sol. in $KCl, NaCl, CaCl_2, BaCl_2$, or $MgCl_2 + Aq$.

very slowly sol. therein at ord. temp. $Na_2S_2O_3, K_4Fe(CN)_6, (NH_4)_2CO_3, (NH_4)_2SO_4$, and NH_4 succinate $+ Aq$, large amt. of hot $NH_4Cl + Aq$. (Witt, Z. anorg. 1901, 34. 1605.)

Sol. in $KCN, NaCN, Ba(CN)_2, Ca(CN)_2$, or $Sr(CN)_2 + Aq$. Insol. in KOH , or $+ Aq$. Sol. in conc. boiling $AgNO_3$ (Wöhler.)

Sol. in 431.7 pts. 5% $NH_4OH + Aq$ (sp. gr. 0.998) at 12° ; in 184.5 pts. 10% $NH_4OH + Aq$ (sp. gr. 0.96) at 18° . (Longi, G. it. 13. 87.)

Sl. sol. in Na citrate $+ Aq$. Sol. in $Hg(NO_3)_2 + Aq$.

1 l. of a 3-N solution of $AgNO_3$ dissolves 1.216 g. $AgCN$ at 25° . (Hellwig, Z. anorg. 1900, 25. 177.)

Very sol. in $(NH_4)_2S_2O_8 + Aq$. (Rosenheim and Steinhäuser, Z. anorg. 1905, 46. 2.)

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Sl. sol. in liquid HF . (Franklin, Z. anorg. 1905, 46. 2.)

Abundantly sol. in quinoline at 60° . (C. R. 1893, 116. 60.)

Sl. sol. in ethyl amine. (Shinn, J. Chem. 1907, 11. 538.)

Insol. in methyl acetate. (Bezold, Z. anorg. 1906; Naumann, B. 1909, 42. 3790.)

Hamers, Dissert. 1906; Naumann, B. 314.)

rogen cyanide, AgCN , HCN .
B. 1903, 36. 1859.)

ium cyanide, AgCN , NaCN .
pts. H_2O at 20° and in much less
Sol. in 24 pts. 85% alcohol at
mp, A. ch. (3) 53. 468.)

lous cyanide, AgCN , TlCN .
ol. in H_2O . 100 pts. H_2O dissolve
at 0° , and 7.4 pts. at 16° . (Fronmüller, B. 11. 92.)

gsten cyanide.
gstocyanide, silver.

ide ammonia, AgCN , NH_3 .
ent. Decomp. on air.
l. in ammonia at -10° . (Joannis,
, 118. 1151.)

ide hydrazine, AgCN , N_2H_4 .
. in the air.
. by H_2O . (Franzen, Z. anorg.
153.)

ide nitrate, 2AgCN , AgNO_3 .
. by H_2O .

anide, NaCN .
 H_2O and 75% alcohol.
, and $2\text{H}_2\text{O}$. Very sol. in H_2O ; sl.
hol. (Joannis, A. ch. (5) 26. 484.)

ngsten cyanide.
gstocyanide, sodium.

inc cyanide, NaCN , $\text{Zn}(\text{CN})_2 + \text{O}$.
ore sol. in H_2O than the correspond-
salt. (Rammelsberg.)
(Loebe, Dissert. 1902.)

cyanide molybdenum dioxide,
 N , $\text{MoO}_3 + 6\text{H}_2\text{O}$.
r, Dissert. 1909.)
\text{H}_2\text{O}. (Rosenheim,
and Kohn, Z. anorg. 1910, 65. 174.)

yanide molybdenum dioxide hy-
amine, 4NaCN , MoO_3 , NH_2OH
mp. (Winkler, Dissert. 1909.)

cyanide, $\text{Sr}(\text{CN})_2 + 4\text{H}_2\text{O}$.
stable; very deliquescent, and sol.
Joannis, A. ch. (5) 26. 496.)

tungsten cyanide.
gstocyanide, strontium.

Strontium zinc cyanide, $2\text{Sr}(\text{CN})_2$,
 $3\text{Zn}(\text{CN})_2 + \text{H}_2\text{O}$.
Sol. in H_2O and alcohol. (Loebe, Dissert.
1902.)

Thallous cyanide, TlCN .
100 pts. H_2O dissolve 16.8 pts. at 28.5° .
(Fronmüller, B. 6. 1178.)

Thallothallic cyanide, $\text{Tl}_2(\text{CN})_4 = \text{TlCN}$,
 $\text{Tl}(\text{CN})_3$.
Easily sol. in H_2O .
100 pts. H_2O dissolve 27.3 pts. at 30° , 15.3
pts. at 12° , 9.7 pts. at 0° . (Fronmüller, B. 11.
92.)

Thallous tungsten cyanide.
See Tungstocyanide, thallous.

Thallous zinc cyanide, 2TlCN , $\text{Zn}(\text{CN})_2$.
Easily sol. in H_2O . 100 pts. H_2O dissolve
8.7 pts. at 0° ; 15.2 pts. at 14° ; and 29.6 pts.
at 31° . (Fronmüller, B. 11. 92.)

Tungsten cyanide with MCN .
See Tungstocyanide, M.

Zinc cyanide, $\text{Zn}(\text{CN})_2$.
Insol. in H_2O and alcohol. Sol. in alkalies.
Easily sol. in $\text{KCN} + \text{Aq}$. Sol. in hot NH_4
salts + Aq . (Wittstein.)
Easily sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Gore.)
Sol. in $\text{KOH} + \text{Aq}$. Solution is stable
when less than 1 mol. $\text{Zn}(\text{CN})_2$ to 2 mols.
 KOH is present. When proportion is 1:1,
 ZnO_2H_2 soon separates.
Sol. in dil. $\text{KCN} + \text{Aq}$. (Sharwood, J.
Am. Chem. Soc. 1903, 25. 587.)
Sl. sol. in conc. Zn salts + Aq . 1 l. conc.
 $\text{Zn}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$ dissolves 4 g., and 1 l.
conc. $\text{ZnSO}_4 + \text{Aq}$ dissolves 2 g. $\text{Zn}(\text{CN})_2$.
Insol. in $\text{HCN} + \text{Aq}$. Easily sol. in dil.
acids. (Joannis.)

Very sol. in liquid NH_3 . (Franklin, Am.
Ch. J. 1898, 20. 830.)
Insol. in methyl acetate. (Naumann, B.
1909, 42. 3790); ethyl acetate. (Naumann,
B. 1910, 43. 314.)

Zinc cyanide ammonia, $\text{Zn}(\text{CN})_2$, 2NH_3 .
Decomp. on air. (Varet, C. R. 105. 1070.)
 $+ \text{H}_2\text{O}$. Decomp. on air. Decomp. by
 H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Varet.)

Zinc cyanide dihydrazine, $\text{Zn}(\text{CN})_2$, $2\text{N}_2\text{H}_4$.
Decomp. by H_2O . (Franzen, Z. anorg
1911, 70. 153.)

Cyanogen, CN .
 H_2O absorbs $4\frac{1}{2}$ vols. CN gas at 20° . Alco-
hol absorbs 23 vols., and ether 5 vols. at the
same temperature. (Gay-Lussac.)
The solution gradually decomposes, but
this is prevented by traces of acids.

0.221 mol. litre are dissolved in H_2O at 0° . (Naumann, Z. Electrochem. 1910, 16. 177.)

Oil of turpentine absorbs 5 vols. (Gay-Lussac.) Absorbed by many essential oils.

Very sol. in $CuCl_2 + Aq$.

Absorbed with decomp. by $NH_4OH + Aq$ and other alkaline liquids.

Absorbed by aniline. (Jacquemain, C. R. 100. 1006.)

Cyclotriborene, B_3H_3 .

Insol. in H_2O . (Ramsay and Hatfield, Proc. Chem. Soc. 1901, 17. 152.)

Decamine cobaltic sulphite,



Sol. in H_2O . (Vortmann and Magdeburg, B. 22. 2636.)

Decamine cobaltisulphurous acid.

Cobaltic decamine cobaltisulphite,



Ppt. (Vortmann and Magdeburg, B. 22. 2635.)

Sodium decamine cobaltisulphite,



Sol. in H_2O . (Vortmann and Magdeburg, B. 22. 2635.)

Diamide, N_2H_4 .

See Hydrazine.

Diamine chromium sulphocyanhydric acid, $Cr(NH_3)_3(SCN)_3, HSCN + H_2O$.

Sol. in H_2O . (Nordenskiöld, Z. anorg. 1. 130.)

Diamine chromium diaquo sulphocyanide, $Cr(NH_3)_2(SCN)_3 + 2H_2O$.

Sol. in H_2O , from which it is pptd. by con. $HCl + Aq$. (Nordenskiöld, Z. anorg. 1. 137.)

Ammonium diamine chromium sulphocyanide, $Cr(NH_3)_2(SCN)_3, NH_4SCN$.

(Reinecke's salt.) Quite easily sol. in H_2O , less in alcohol, and insol. in benzene. Slowly decomp. by boiling H_2O or dil. acids. (Nordenskiöld, Z. anorg. 1. 130.)

+ H_2O . Insol. in absolute ether. (Christensen, J. pr. (2) 45. 218.)

Ammonium diamine chromium sulphocyanide iodide, $Cr(NH_3)_2(SCN)_3, NH_4SCN, I$.

Barium ———, $[Cr(NH_3)_2(SCN)_3]_2, Ba(SCN)_2$.

Sol. in H_2O and alcohol. (N.)

Cadmium diamine chromium sulphocyanide, $Cd(SCN)_2, [Cr(NH_3)_2(SCN)_3]_2 + H_2O$.

Nearly insol. in cold, sl. sol. in hot H_2O . Sl. sol. in boiling alcohol. (Christensen, J. pr. (2) 45. 371.)

Cupric ———, $Cu(SCN)_2, [Cr(NH_3)_2(SCN)_3]_2$.

Insol. in H_2O or dil. acids. (Reinecke, Z. anorg. 126. 116.)

Ferric ———, $[Cr(NH_3)_2(SCN)_3]_2, Fe(SCN)_3$. (N.)

Luteocobaltic ———, $Co(NH_3)_6(SCN)_3[Cr(NH_3)_2(SCN)_3]_2$.

As good as insol. in cold H_2O . Sl. sol. in hot H_2O and alcohol. (Christensen, J. pr. (2) 45. 370.)

Mercuric ———, $[Cr(NH_3)_2(SCN)_3]_2, Hg(SCN)_2$.

Insol. in H_2O . (N.)

Insol. in H_2O and dil. acids. (Reinecke, Z. anorg. 126. 116.)

Potassium ———, $Cr(NH_3)_2(SCN)_3, KSCN$.

Properties as the NH_4 salt. (N.)

$Cr(NH_3)_2(SCN)_3, KSCN, I$. As the salt. (N.)

Sodium ———, $NaSCN, Cr(NH_3)_2(SCN)_3$.

Sol. in H_2O , alcohol, and ether. (Reinecke, Z. anorg. 126. 116.)

Diamine cobaltic nitrite ammonium nitrite, $Co(NH_3)_2(NO_2)_3, NH_4NO_2$.

Sol. in H_2O . (Erdmann.)

——— nitrite lead nitrite, $2Co(NH_3)_2(NO_2)_3, Pb(NO_2)_2$.

Sol. in hot H_2O with partial decomp.

——— nitrite mercurous nitrite, $2Co(NH_3)_2(NO_2)_3, Hg_2(NO_2)_2$.

Ppt. Not sol. in hot H_2O without decomp.

——— nitrite potassium nitrite, $Co(NH_3)_2(NO_2)_3, KNO_2$.

Sol. in H_2O . (Erdmann, J. pr. 97.)

——— nitrite silver nitrite, $Co(NH_3)_2(NO_2)_3, AgNO_2$.

Ppt. Crystallises out of hot H_2O . (Erdmann.)

——— nitrite thallium nitrite, $Co(NH_3)_2(NO_2)_3, TlNO_2$.

Crystallises out of hot H_2O without decomp.

praseobaltic carbonate,
 $(\text{H}_2)_3(\text{OH})\text{CO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.
 H_2O . (Vortmann, B. 15. 1901.)

praseobaltic chloride, $\text{Co}(\text{NH}_3)_3\text{Cl}_3 + \text{H}_2\text{O}$.
 sol. in cold H_2O , dil. acids, conc.
 dil. alcohol.
 solution in conc. H_2SO_4 , the salt is
 precipitated by much $\text{HCl} + \text{Aq}$. Composi-
 $\text{Co}(\text{NH}_3)_3(\text{OH})_2\text{Cl}_3$. (Jorgensen, Z.
 189.)

praseobaltate, $\text{Co}(\text{NH}_3)_3(\text{NO}_3)_3 + 4\text{H}_2\text{O}$.
 efflorescent. Sol. in H_2O . More sol. in
 $\text{H}_2\text{O} + \text{Aq}$ than praseocobaltic nitrate.
 (Jorgensen, B. 15. 1897.)
 praseobaltous. Insol. in H_2O as such, but
 dissolves into above salt thereby. (Jorgen-
 berg, org. 5. 186.)

praseobaltite, $\text{Co}(\text{NH}_3)_3(\text{NO}_3)_3$.
 slightly sol. in cold, but rather easily sol.
 in H_2O .

praseobaltophosphate, $[\text{Co}(\text{NH}_3)_3]_2(\text{SO}_4)_3 + 6\text{H}_2\text{O}$.
 sol. in H_2O . (Vortmann, B. 15.

praseobaltic white, $[\text{Co}(\text{NH}_3)_3]_2(\text{SO}_4)_3 + \text{H}_2\text{O}$.
 insol. in cold, slowly decomp. by hot
 comp. by acids or $\text{KOH} + \text{Aq}$. Insol.
 sol. in warm $\text{NH}_4\text{OH} + \text{Aq}$. (Kunzel,
 72. 209.) According to Geuther (A.
 189), is a double salt—
 $[\text{I}_3]_2(\text{SO}_4)_3, \text{Co}_2(\text{SO}_4)_3 + 2\text{H}_2\text{O}$.

praseodymium, Di.
 decomp. by H_2O . Insol. in cold
 O_2 . Sol. in dil. acids.
 composed of two elements, neodymium
 and praseodymium. (v. Welsbach, W. A.
 7.)

praseodymium bromide, $\text{DiBr}_3 + 6\text{H}_2\text{O}$.
 deliquescent, and sol. in H_2O .

praseodymium nickel bromide, $2\text{DiBr}_3, 3\text{NiBr}_2 +$
 $3\text{H}_2\text{O}$.
 efflorescent. Very sol. in H_2O . (Frerichs
 and Smith, A. 191. 342.)

praseodymium zinc bromide, $\text{DiBr}_3, 3\text{ZnBr}_2 +$
 $3\text{H}_2\text{O}$.
 very deliquescent. (Cleve, Bull.
 13. 361.)
 $3\text{ZnBr}_2 + 36\text{H}_2\text{O}$. (F. and S.)

praseodymium chloride, DiCl_3 .
 praseodymous. Deliquescent. Sol. in H_2O
 and alcohol. (Marignac.)
 praseodymic. Deliquescent. Easily sol. in
 alcohol. (Marignac.)

Didymium mercuric chloride, $2\text{DiCl}_3,$
 $9\text{HgCl}_2 + 24\text{H}_2\text{O}$.

More sol. in H_2O than the corresponding
 La salt. (Marignac.)
 $\text{DiCl}_3, 4\text{HgCl}_2 + 11\text{H}_2\text{O}$. Not deliquescent.
 Easily sol. in H_2O .

Didymium stannic chloride.
 See Chlorostannate, didymium.

Didymium fluoride, $\text{DiF}_3 + \frac{1}{2}\text{H}_2\text{O}$.
 Precipitate. (Cleve.)

Didymium hydrogen fluoride, $2\text{DiF}_3, 3\text{HF}$.
 Precipitate. (Smith.)
 Does not exist. (Cleve.)

Didymium potassium fluoride, $\text{DiF}_3, \text{KF} +$
 H_2O .
 Sol. in H_2O . (Brauner, B. 15. 114.)
 $+ \frac{1}{2}\text{H}_2\text{O}$. As above. (B.)
 $2\text{DiF}_3, 3\text{KF} + \text{H}_2\text{O}$. As above. (B.)

Didymium hydroxide, $\text{Di}_2\text{O}_3\text{H}_6$.
 Insol. in KOH , or $\text{NaOH} + \text{Aq}$, but is sl.
 sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Rose.)
 See also Di_2O_3 .

Didymium penthydroxide, $\text{DiO}_4\text{H}_8 = \text{Di}_2\text{O}_5,$
 $3\text{H}_2\text{O}$.
 Precipitate. (Brauner, B. 15. 113.)

Didymium zinc iodide, $2\text{DiI}_3, 3\text{ZnI}_2 + 24\text{H}_2\text{O}$.
 Very deliquescent. (Frerichs and Smith.)

Didymium oxide, Di_2O_3 .
 With H_2O slowly forms $\text{Di}_2\text{O}_3\text{H}_6$.
 Sol. in conc., or dil. mineral acids (Marig-
 nac), and in acetic acid (Hermann). Sol. in
 ammonium salts + Aq .
 Slightly more slowly sol. in conc. NH_4NO_3
 + Aq than La_2O_3 . (Damour and Deville.)
 A solution of NH_4NO_3 in H_2O that can
 dissolve 2.9 mols. La_2O_3 dissolves 1 mol.
 Di_2O_3 . (Brauner, B. 15. 114.)
 Insol. in acetone. (Naumann, B. 1904,
 37. 4329.)

Didymium peroxide, Di_4O_9 .
 Sol. in acids with decomp. (Frerichs, B.
 7. 799.)

Not obtained by Cleve. (B. 11. 910.)
 The contradictory statements concerning
 the composition of Di peroxide are owing to
 the fact that praseodidymium is the only one
 of the constituents of Di which easily forms a
 peroxide. (v. Welsbach.)

Didymium pentoxide, Di_2O_5 .
 Sol. in dil. HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq}$ in the
 cold without evolution of gas, but gas is
 evolved if treated with conc. acids. Insol.
 in $\text{HF} + \text{Aq}$. Sl. sol. in cold $\text{NH}_4\text{NO}_3 + \text{Aq}$.
 $= \text{Di}_4\text{O}_9$. (Cleve.)

Didymium oxybromide, DiOBr .

(Frerichs and Smith.)

Didymium oxychloride, DiOCl .

Anhydrous. Insol. in H_2O . (Smith.)
 $+3\text{H}_2\text{O}$. Sol. in cold dil. $\text{HNO}_3 + \text{Aq}$. (Marignac.)
 Sl. sol. in $\text{HCl} + \text{Aq}$. (Hermann.)

Didymium oxysulphide, $\text{Di}_2\text{O}_3\text{S}$.

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$ without residue. (Marignac.)

Didymium sulphide, Di_2S_3 .

Insol. in H_2O . Decomp. by dil. acids. (Marignac, A. ch. (3) 38. 159.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Disulphuric acid, $\text{H}_2\text{S}_2\text{O}_7$.*See Disulphuric acid.***Dithionic acid (Hyposulphuric acid), $\text{H}_2\text{S}_2\text{O}_6$.**

Known only in aqueous solution, which is stable only when dil. Can be evaporated in vacuo until sp. gr. = 1.347, but decomp. upon further evaporation. (Welter and Gay-Lussac, A. ch. 10. 312.)

Dithionates.

All dithionates are sol. in H_2O .

Aluminum dithionate, $\text{Al}_2(\text{S}_2\text{O}_6)_3 + 18\text{H}_2\text{O}$.

Extremely deliquescent. Easily sol. in H_2O or absolute alcohol. (Klüss, A. 246. 218.)

Aluminum ammonium dithionate, $\text{Al}_2(\text{S}_2\text{O}_6)_3, (\text{NH}_4)_2\text{S}_2\text{O}_6 + 27\text{H}_2\text{O}$.

Sl. deliquescent. Sol. in H_2O . (Klüss, A. 246. 303.)

Ammonium dithionate, $(\text{NH}_4)_2\text{S}_2\text{O}_6$.

Very sol. in H_2O . Sol. in 0.79 pt. H_2O at 16° , with reduction of temp. Not decomp. on boiling. Insol. in absolute alcohol. (Heeren, Pogg. 7. 172.)

Contains $\frac{1}{2}\text{H}_2\text{O}$. Sol. in 0.56 pt. H_2O at 19° . (Klüss, A. 246. 194.)

Ammonium cadmium dithionate, $2(\text{NH}_4)_2\text{S}_2\text{O}_6, \text{CdS}_2\text{O}_6 + 4\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . (Klüss, A. 246. 298.)

Ammonium cobalt dithionate, $9(\text{NH}_4)_2\text{S}_2\text{O}_6, 2\text{CoS}_2\text{O}_6 + 16\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . (Klüss.)

Ammonium cupric dithionate, $(\text{NH}_4)_2\text{S}_2\text{O}_6, 2\text{CuS}_2\text{O}_6 + 8\text{H}_2\text{O}$.

Sol. in H_2O .

Ammonium ferrous dithionate, $3(\text{NH}_4)_2\text{FeS}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Klüss, A. 246. 300.)
 $9(\text{NH}_4)_2\text{S}_2\text{O}_6, 2\text{Fe}_2\text{S}_2\text{O}_6 + 16\frac{1}{2}\text{H}_2\text{O}$.
 in H_2O . (Klüss.)

Ammonium manganous dithionate, $9(\text{NH}_4)_2\text{S}_2\text{O}_6, 2\text{MnS}_2\text{O}_6$.

Sol. in H_2O . (Klüss, A. 246. 301.)

Ammonium nickel dithionate, $9(\text{NH}_4)_2\text{NiS}_2\text{O}_6 + 16\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . (Klüss.)

Ammonium zinc dithionate, $5(\text{NH}_4)_2\text{ZnS}_2\text{O}_6 + 9\text{H}_2\text{O}$.

Easily sol. in H_2O . (Klüss, A. 246. 301.)
 $9(\text{NH}_4)_2\text{S}_2\text{O}_6, 2\text{ZnS}_2\text{O}_6 + 16\frac{1}{2}\text{H}_2\text{O}$.
 sol. in H_2O . (Klüss.)

Ammonium dithionate chloride, $(\text{NH}_4)_2\text{S}_2\text{O}_6, \text{NH}_4\text{Cl}$.

Sol. in H_2O . (Fock and Klüss, 3017.)

Barium dithionate, $\text{BaS}_2\text{O}_6 + 2\text{H}_2\text{O}$.

Not efflorescent. Sol. in 7.17 pts. H_2O at 8° , 4.04 pts. at 18° , and 1.1 pts. at 25° .
 Insol. in alcohol. (Gay-Lussac, 1807.)
 Sol. in 0.994 pt. H_2O at 102° , the point of the sat. solution. (Baker, Soc. (2) 44. 166.)

Insol. in methyl acetate. (Naumann, 1909, 42. 3790.)

$+4\text{H}_2\text{O}$. Very efflorescent. (Heeren, 1807.)

Barium magnesium dithionate, $\text{BaMgS}_2\text{O}_6 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Schiff, A. 118. 97.)

Barium rubidium dithionate, $\text{BaRbS}_2\text{O}_6 + \text{H}_2\text{O}$.

Sol. in H_2O . Solubility is diminished in presence of excess of Rb_2SO_4 , but increased in presence of BaS_2O_6 . (Bodlander, Chem. Ztg. 14. 186.)

Barium sodium dithionate, $\text{BaNa}_2\text{S}_2\text{O}_6 + 4\text{H}_2\text{O}$.

Sol. in H_2O . Decomp. by recrystallization. (Kraut, A. 118. 95.)
 $+6\text{H}_2\text{O}$. (Schiff.)

Barium dithionate chloride, $\text{BaS}_2\text{O}_6, \text{HCl} + 4\text{H}_2\text{O}$.

(Fock and Klüss, B. 23. 3001.)

Bismuth dithionate, basic, $\text{Bi}_2\text{O}_3, \text{S}_2\text{O}_6 + 5\text{H}_2\text{O}$.

Efflorescent. Insol. in H_2O , but soluble in dil. acids, especially $\text{HCl} + \text{Aq}$. (Klüss, A. 246. 183.)

$4\text{Bi}_2\text{O}_3, 3\text{S}_2\text{O}_6 + 5\text{H}_2\text{O}$. Insol. in H_2O , but soluble in dil. acids. (Klüss.)

dithionate.	Didymium dithionate, $\text{Di}_2(\text{S}_2\text{O}_6)_3 + 24\text{H}_2\text{O}$. Extremely sol. in H_2O . (Cleve.)
insol. in moist air; very sol. in H_2O . Pogg. 7. 183.)	
dithionate ammonia, CdS_2O_6 ,	Erbium dithionate, $\text{Er}_2(\text{S}_2\text{O}_6)_3 + 18\text{H}_2\text{O}$. Very sol. in H_2O or alcohol; insol. in ether. (Höglund.)
insol. by alcohol; sol. in $\text{NH}_4\text{OH} + \text{Aq}$, decomp. on heating. (Rammelsberg, 298.)	Glucinum dithionate, basic, $5\text{GfO}, 2\text{S}_2\text{O}_6 + 14\text{H}_2\text{O}$. Easily sol. in H_2O and absolute alcohol. (Klüss, A. 246. 196.)
dithionate, $\text{Cs}_2\text{S}_2\text{O}_6$. sol. in H_2O . (Chabrié, C. R. 1901,	Iron (ferrous) dithionate, $\text{FeS}_2\text{O}_6 + 5\text{H}_2\text{O}$. Very sol. in H_2O . Insol. in alcohol. De- comp. in aqueous solution into FeSO_4 by boiling. (Heeren, Pogg. 7. 181.) $+ 7\text{H}_2\text{O}$. Sol. in 0.59 pt. H_2O at 18.5° . (Klüss, A. 246. 198.)
dithionate, $\text{CaS}_2\text{O}_6 + 4\text{H}_2\text{O}$. 1.46 pts. H_2O at 19° ; 0.8 pt. at 100° . sol. in alcohol. (Heeren, Pogg. 7. 178.) insol. in acetone. (Naumann, B. 1904,	Iron (ferric) dithionate, basic, $8\text{Fe}_2\text{O}_3, \text{S}_2\text{O}_6 + 20\text{H}_2\text{O}$. Insol. in H_2O or alcohol. Very sl. sol. in $\text{H}_2\text{S}_2\text{O}_6 + \text{Aq}$; easily sol. in $\text{HCl} + \text{Aq}$. (Heeren.) Contains $14\text{H}_2\text{O}$. (Klüss, A. 246. 200.) $3\text{Fe}_2\text{O}_3, \text{S}_2\text{O}_6 + 8\text{H}_2\text{O}$. Insol. in H_2O . Easily sol. in acids. (Klüss, A. 246. 201.)
dithionate, $\text{Ce}_2(\text{S}_2\text{O}_6)_3 + 24\text{H}_2\text{O}$. insol. in H_2O . (Jolin.) $+ 5\text{H}_2\text{O}$. (Wyruboff.)	Lanthanum dithionate, $\text{La}_2(\text{S}_2\text{O}_6)_3 + 16\text{H}_2\text{O}$, and $24\text{H}_2\text{O}$. Sol. in H_2O . (Cleve.)
dithionate, $\text{Cr}_2(\text{S}_2\text{O}_6)_3 + 18\text{H}_2\text{O}$. sol. in H_2O and alcohol. (Klüss, A. 246.	Lead dithionate, basic, $2\text{PbO}, \text{S}_2\text{O}_6 + 2\text{H}_2\text{O}$. Very difficultly sol. in H_2O . (Heeren, Pogg. 7. 171.) $10\text{PbO}, \text{S}_2\text{O}_6 + 2\text{H}_2\text{O}$. Sl. sol. in H_2O . (Heeren.)
$4\text{S}_2\text{O}_6 + 24\text{H}_2\text{O}$. Easily sol. in H_2O Insol. in ether. (Klüss.)	Lead dithionate, $\text{PbS}_2\text{O}_6 + 4\text{H}_2\text{O}$. Easily sol. in H_2O . (Heern.) Sol. in 0.869 pt. H_2O at 20.5° . (Baker, C. N. 36. 203.)
dithionate, $\text{CoS}_2\text{O}_6 + 6\text{H}_2\text{O}$. efflorescent. Very sol. in H_2O .	Lead strontium dithionate, $(\text{Pb}, \text{Sr})\text{S}_2\text{O}_6 + 4\text{H}_2\text{O}$. (Rammelsberg.)
Sol. in 0.49 pt. H_2O at 19° . Sol. in alcohol. (Klüss, A. 246. 203.)	Lithium dithionate, $\text{Li}_2\text{S}_2\text{O}_6 + 2\text{H}_2\text{O}$. Sl. deliquescent, and easily sol. in H_2O . Insol. in alcohol. (Rammelsberg.)
dithionate basic, $4\text{CuO}, \text{S}_2\text{O}_6 + 4\text{H}_2\text{O}$. sol. in H_2O . (Heeren, Pogg. 7.	Magnesium dithionate, $\text{MgS}_2\text{O}_6 + 6\text{H}_2\text{O}$. Sol. in 0.85 pt. H_2O at 13° . Solution can be boiled without decomp. (Heeren, Pogg. 7. 179.) Sol. in 0.692 pt. H_2O at 17° . (Baker, C. N. 36. 203.)
1 pt. H_2O ; easily sol. in dil. acids. (Klüss, A. 246. 208.)	Manganous dithionate, $\text{MnS}_2\text{O}_6 + 3\text{H}_2\text{O}$. Sol. in H_2O . (Kraut, A. 118. 98.) $+ 6\text{H}_2\text{O}$. Efflorescent. Sol. in H_2O . (Mar- ignac, J. B. 1855. 380.)
Insol. in H_2O and $\text{NaC}_2\text{H}_3\text{O}_2 +$ in traces in conc. $\text{CuS}_2\text{O}_6 + \text{Aq}$. insol. in dil. acids, even $\text{HC}_2\text{H}_3\text{O}_2$, or in aq . (Klüss.)	
dithionate, $\text{CuS}_2\text{O}_6 + 4\text{H}_2\text{O}$. efflorescent. Very sol. in H_2O . Insol. (Heeren.) (Efflorescent. Sol. in 0.64 pt. H_2O at 5° . (Klüss, A. 246. 204.)	
dithionate ammonia, $\text{CuS}_2\text{O}_6, 4\text{NH}_3$. slightly sol. in cold H_2O , moderately sol. at 40° . Decomp. by much H_2O or by solution above 60° . Decomp. by (Heeren.) recryst. from $\text{NH}_4\text{OH} + \text{Aq}$. in liquid NH_3 . (Horn, Am. Ch. J. 1813.) 9NH_3 . Decomp. at ord. temp.	
insol. in liquid NH_3 . (Horn, Am. Ch. J. 1813.)	

Mercurous dithionate, $\text{Hg}_2\text{S}_2\text{O}_6$.

Sl. sol. in cold, decomp. by hot H_2O . (Rammelsberg.)

Mercuric dithionate, basic, 5HgO , $2\text{S}_2\text{O}_6$.

Sl. sol. in cold, decomp. by hot H_2O . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Rammelsberg, Pogg. 59. 472.)

Mercuric dithionate, $\text{HgS}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Decomp. by H_2O or on standing. (Klüss, A. 246. 216.)

Nickel dithionate, $\text{NiS}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Topsøe.)
Sol. in 0.897 pt. H_2O at 12° . (Baker, C. N. 36. 203.)

Nickel dithionate ammonia, NiS_2O_6 , 6NH_3 .

Can be recryst. from warm $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by H_2O . (Rammelsberg, Pogg. 58. 295.)

Nickel dithionate hydrazine, NiS_2O_6 , $3\text{N}_2\text{H}_4$.

Unstable.
Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Franzen, Z. anorg. 1908, 60. 267.)

Potassium dithionate, $\text{K}_2\text{S}_2\text{O}_6$.

Not deliquescent. Sol. in 16.5 pts. H_2O at 16° , and 1.58 pts. at 100° . Insol. in alcohol. (Heeren.)

Sol. in 2.65 pts. H_2O at 16° . (Dumas.)
Sol. in 16.5 pts. H_2O at 16° ; in 1.58 pts. boiling H_2O .

Insol. in alcohol. (Heeren, Pogg. 1826, 7. 72.)

Praseodymium dithionate, $\text{Pr}_2(\text{S}_2\text{O}_6)_3 + 12\text{H}_2\text{O}$.

Deliquescent; very sol. in H_2O . (von Schule, Z. anorg. 1898, 18. 361.)

Rubidium dithionate, $\text{Rb}_2\text{S}_2\text{O}_6$.

Sol. in H_2O . (Topsøe and Christiansen.)

Ruthenium dithionate, RuS_2O_6 .

Ppt. from aq. sol. by alcohol. (Antony, Gazz. ch. it. 1898, 28. 139-142.)

Silver dithionate, $\text{Ag}_2\text{S}_2\text{O}_6 + 2\text{H}_2\text{O}$.

Sol. in 2 pts. H_2O at 16° . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Heeren, Pogg. 7. 191.)

Silver sodium dithionate, AgS_2O_6 , $\text{Na}_2\text{S}_2\text{O}_6 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Kraut, A. 118. 96.)

Silver dithionate ammonia, $\text{Ag}_2\text{S}_2\text{O}_6$, 4NH_3 .

Sol. in H_2O without decomp. (Rammelsberg, Pogg. 58. 298.)

Sodium dithionate, $\text{Na}_2\text{S}_2\text{O}_6 + 2\text{H}_2\text{O}$.

Sol. in 2.1 pts. H_2O at 16° , and in 1.1 pt. boiling H_2O . Insol. in alcohol.

Fuming $\text{HCl} + \text{Aq}$ precipitates the salt from aqueous solution. (Heeren, Pogg. 76.)

+ $6\text{H}_2\text{O}$. (Kraut, A. 117. 97.)

Strontium dithionate, $\text{SrS}_2\text{O}_6 + 4\text{H}_2\text{O}$.

Sol. in 4.5 pts. H_2O at 16° , 1.5 pts. boiling H_2O . Insol. in alcohol. (Heeren, Pogg. 7. 177.)

Thallous dithionate, $\text{Tl}_2\text{S}_2\text{O}_6$.

Very easily sol. in H_2O . (Werther.)

Thallous dithionate sulphate, $3\text{Tl}_2\text{S}_2\text{O}_6$, Tl_2SO_4 .

Sol. in H_2O . (Wyruboff, Ann. Phys. Beibl. 8. 802.)

Thorium dithionate, $\text{Th}(\text{S}_2\text{O}_6)_3 + 4\text{H}_2\text{O}$ (?).

Very unstable. (Klüss, A. 246. 188.)

Tin (stannous) dithionate, SnS_2O_6 .

Known only in solution.

8SnO , $\text{S}_2\text{O}_6 + 9\text{H}_2\text{O}$. Insol. in H_2O . Sol. in dil. acids, even dithionic acid + Aq . (Klüss, A. 246. 186.)

Uranous dithionate, 6UO_2 , $\text{S}_2\text{O}_6 + 10\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in warm $\text{HCl} + \text{Aq}$. (Klüss, A. 246. 191.)

7UO_2 , $\text{S}_2\text{O}_6 + 8\text{H}_2\text{O}$. As above.

8UO_2 , $\text{S}_2\text{O}_6 + 21\text{H}_2\text{O}$. As above.

Divanadyl dithionate, $(\text{VO}_2)_2\text{S}_2\text{O}_6$.

Sol. in H_2O . (Bevan, C. N. 33. 294.)

Yttrium dithionate, $\text{Y}_2(\text{S}_2\text{O}_6)_3 + 18\text{H}_2\text{O}$.

Not deliquescent. Easily sol. in H_2O , but difficultly sol. in alcohol. Insol. in ether. (Cleve, Bull. Soc. (2) 21. 344.)

Zinc dithionate, $\text{ZnS}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Very sol. in H_2O ; decomp. on boiling. (Heeren, Pogg. 7. 183.)

Zinc dithionate ammonia, ZnS_2O_6 , 4NH_3 .

Decomp. with H_2O ; sol. in warm, less so in cold $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg, Pogg. 58. 297.)

+ H_2O . Ppt. (Ephraim, B. 1915, 4. 640.)

Dysprosium, Dy.

(Lecoq de Boisbaudran, C. R. 102. 1006)

Dysprosium chloride, $\text{DyCl}_3 + 6\text{H}_2\text{O}$.

Deliquescent, sol. in H_2O . (Urbain, C. R. 1908, 146. 129.)

m.

1 chloride, EuCl_2 .

H_2O . Stable in very dil. aqueous but decomp. when the solution is heated at 100° . (Urbain, C. R. 1911, .)

Er.

poses H_2O . (Höglund.)

so-called element "erbium" can be decomp. into simple substances. *J. anorg. 3.* 353.)

bromide, $\text{ErBr}_3 + 9\text{H}_2\text{O}$.

deliquescent.

chloride, $\text{ErCl}_3 + 6\text{H}_2\text{O}$.

deliquescent. Sol. in H_2O and alcohol. (.)

mercuric chloride, $\text{ErCl}_3, 5\text{HgCl}_2 +$

deliquescent. (Cleve.)

fluoride, ErF_3 .

insol. in H_2O . Very sl. sol. in $\text{HF} + \text{Aq}$. *J. Bull. Soc. (2)* 18. 193.)

hydroxide, $\text{Er}_2\text{O}(\text{OH})_4$.insol. in KOH , or $\text{NaOH} + \text{Aq}$.

sol. in acids. Decomp. ammonium oiling therewith.

iodide, ErI_3 .

deliquescent. Very sol. in H_2O and insol. in ether. (Höglund.)

oxide, Er_2O_3 .

slightly but completely sol. in warm H_2SO_4 , or $\text{HCl} + \text{Aq}$. Decomp. NH_4 oiling therewith.

peroxide, Er_2O_5 .

stable. (Cleve, *Bull. Soc. (2)* 43.)

sulphide.

decomp. in moist air and with acids.

chromium bromide,

 $\text{Cr}_2(\text{NH}_3)_{10}\text{Br}_5 + \text{H}_2\text{O}$.

easily sol. in H_2O . Insol. in $\text{HBr} + \text{Aq}$. in $\text{NH}_4\text{OH} + \text{Aq}$. (Jörgensen, *J.* 398.)

nitride, basic, $\text{HOCr}_2(\text{NH}_3)_{10}(\text{OH})\text{Br}_4$ O .

insol. in H_2O . (Jörgensen.)

triiodide, $\text{HOCr}_2(\text{NH}_3)_{10}\text{I}_3 + \text{H}_2\text{O}$.

insol. in H_2O and in alcohol. (Jörgensen.)

Erythrochromium chloroplatinate,

 $[\text{HOCr}_2(\text{NH}_3)_{10}]_2(\text{PtCl}_6)_5 + 10\text{H}_2\text{O}$.

Nearly insol. in H_2O . (Jörgensen.)

— dithionate, basic,

 $\text{HOCr}_2(\text{NH}_3)_{10}(\text{S}_2\text{O}_6)_2(\text{OH}) + 2\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in very dil. HNO_3 , HBr , $\text{HCl} + \text{Aq}$. Sol. in conc. $\text{NH}_4\text{Cl} + \text{Aq}$. (Jörgensen.)

— nitrate, $\text{HOCr}_2(\text{NH}_3)_{10}(\text{NO}_3)_5 + \text{H}_2\text{O}$.

Easily sol. in H_2O . Insol. in dil. $\text{HNO}_3 + \text{Aq}$. Sol. in conc. HNO_3 with decomp. Very sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in alcohol. (Jörgensen.)

— nitrate, basic, $\text{HOCr}_2(\text{NH}_3)_{10}(\text{NO}_3)_4\text{OH} + 3\frac{1}{2}\text{H}_2\text{O}$.

Sol. in cold H_2O . (Jörgensen.)

— sulphate, $[\text{HOCr}_2(\text{NH}_3)_{10}]_2(\text{SO}_4)_5$.

Nearly insol. in H_2O . (Jörgensen.)

Tetraferriammonium, Fe_2N .

See Iron nitride.

Ferric acid.

Barium ferrate, $\text{BaFeO}_4 + \text{H}_2\text{O}$.

Ppt. Can be boiled for some time with H_2O without decomp. Decomp. by mineral acids. Sol. in dil. acetic acid. (Fremy, *A. ch. (3)* 12. 373.)

Insol. in H_2O ; not readily acted upon by acids when dry. (Rosen, *J. Am. Chem. Soc.* 1895, 17. 766.)

Ppt. Easily decomp. by acids. (Moeser, *Arch. Pharm.* 1895, 233. 526.)

Insol. in acetone. (Naumann, *B.* 1904, 37. 4329.)

Calcium ferrate, CaFe_2O_4 .

Sol. in H_2O . (Rosell, *J. Am. Chem. Soc.* 1895, 17. 760–69.)

Potassium ferrate, K_2FeO_4 .

Very deliquescent. Easily sol. in cold H_2O with evolution of much heat. Decomp. by standing or warming. Decomp. by acids or alkalis. (Fremy, *A. ch. (3)* 12. 369.)

Sol. in H_2O ; insol. in alcohol. (Moeser, *Arch. Pharm.* 1895, 233. 524.)

Quickly decomp. by potassium tartrate or racemate, sugar, or albumen without separation of $\text{Fe}_2\text{O}_3\text{H}_2$, by alcohol with separation of $\text{Fe}_2\text{O}_3\text{H}_2$. Potassium oxalate, acetate, formate, and benzoate, also citrate decomp. much more slowly. Insol. in conc. $\text{KOH} + \text{Aq}$. (Wackenroder, *A.* 33. 41.)

Sodium ferrate, Na_2FeO_4 .

Sol. in H_2O and in conc. $\text{NaOH} + \text{Aq}$. (Fremy, *l. c.*)

Strontium ferrate, SrFeO_4 .

Sl. sol. in H_2O by which it is decomp.

Decomp. by acids.

Sol. in aqueous solutions of Na and K salts with partial decomp.

Insol. in sat. $\text{SrBr}_2 + \text{Aq.}$, alcohol and ether. (Eidmann, B. 1903, 36, 2290.)

Ferricomolybdic acid.

Ammonium ferricomolybdate, $3(\text{NH}_4)_2\text{O}, \text{Fe}_2\text{O}_3, 12\text{MoO}_3 + 19\text{H}_2\text{O}$.

Ppt. (Hall, J. Am. Chem. Soc. 1907, 29, 697.)

Ferricyanhydric acid,

$\text{H}_2\text{Fe}(\text{CN})_6$, (or $\text{H}_2\text{Fe}_2(\text{CN})_{12}$).

Easily sol. in H_2O or alcohol. Solution decomposes slowly by standing, more rapidly by heating. Insol. in ether.

Ferricyanides.

The alkali, and alkaline-earth ferricyanides are sol. in H_2O ; the others are insol. The ferricyanides of metals, the oxides of which are sol. in NH_4OH , or $\text{KOH} + \text{Aq.}$ are themselves sol. in those reagents.

Ammonium ferricyanide, $(\text{NH}_4)_3\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Permanent. Readily sol. in H_2O (and alcohol?).

Ammonium ferrous ferricyanide, $\text{NH}_4\text{FeFe}(\text{CN})_6 + 1\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O and not pptd. by alcohol from aqueous solution. More stable than the corresponding K salt.

Ammonium lead ferricyanide, $\text{NH}_4\text{PbFe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Ammonium potassium ferricyanide, $(\text{NH}_4)_2\text{KFe}(\text{CN})_6$.

Sol. in H_2O . (Schaller, Bull. Soc. (2) 1, 275.)

Barium ferricyanide, $\text{Ba}_3[\text{Fe}(\text{CN})_6]_2 + 20\text{H}_2\text{O}$.

Easily sol. in H_2O ; insol. in alcohol. (Schuler, W. A. B. 77, 692.)

Barium potassium ferricyanide, $\text{BaKFe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Permanent. Easily sol. in H_2O , less in alcohol.

Barium ferricyanide bromide, $\text{Ba}_3[\text{Fe}(\text{CN})_6]_2, 2\text{BaBr}_2 + 20\text{H}_2\text{O}$.

Easily sol. in H_2O . Boiling alcohol does not dissolve out BaBr_2 . (Rammelsberg, J. pr. (2) 39, 463.)

Bismuth ferricyanide, $\text{Bi}_3[\text{Fe}(\text{CN})_6]_2$.

Insol. in H_2O , but decomp. by therewith. (Muir, Chem. Soc. 33, 40.)

Cadmium ferricyanide ammonia, $\text{Cd}_3[\text{Fe}(\text{CN})_6]_2, 6\text{NH}_3 + 3\text{H}_2\text{O}$.

Effloresces to form—

$\text{Cd}_3[\text{Fe}(\text{CN})_6]_2, 4\text{NH}_3 + 2\text{H}_2\text{O}$. In H_2O . (Wyrouboff, A. hc. (5) 10, 413.)

Calcium ferricyanide, $\text{Ca}_3[\text{Fe}(\text{CN})_6]_2$ or $12\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O and dil. a

Calcium potassium ferricyanide, $\text{CaKFe}(\text{CN})_6$.

Sol. in H_2O .

Cerous ferricyanide, $\text{CeFe}(\text{CN})_6 + 4\text{H}_2\text{O}$.
Sol. in H_2O ; easily decomp. (Jolin.)

Chromic ferricyanide (?).

Ppt.

Cobaltous ferricyanide, $\text{Co}_3[\text{Fe}(\text{CN})_6]_2$.
Insol. in H_2O and $\text{HCl} + \text{Aq.}$ 8 $\text{NH}_4\text{OH} + \text{Aq.}$

Cobaltous ferricyanide ammonia, $\text{Co}_3[\text{Fe}(\text{CN})_6]_2, 4\text{NH}_3 + 6\text{H}_2\text{O}$.

Cobaltic ferricyanide ammonia.

See Luteo, - purpureo, - etc. cobaltic cyanide.

Cuprous ferricyanide, $(\text{Cu}_2)_3[\text{Fe}(\text{CN})_6]_2$.
Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$; insol. in $\text{NH}_4\text{Aq.}$ (Wittstein.)

Cupric ferricyanide, $\text{Cu}_3[\text{Fe}(\text{CN})_6]_2$.
Insol. in H_2O or NH_4 salts + Aq. 1 NH_4OH , and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ (Wittstein.)
Insol. in $\text{HCl} + \text{Aq.}$

Iron (ferrous) ferricyanide, $\text{Fe}_3[\text{Fe}(\text{CN})_6]_2, x\text{H}_2\text{O}$.

(Turnbull's blue.) Properties as ferrocyanide (Prussian blue), with which perhaps identical. (Gintl, Z. anal. 21)

Iron (ferrosoferric) ferricyanide, $\text{Fe}_3(\text{CN})_{12} = \text{Fe}^{\text{III}}_2\text{Fe}^{\text{II}}[\text{Fe}(\text{CN})_6]_2$.

(Prussian green.) Insol. in H_2O or $\text{HCl} + \text{Aq.}$, but slowly decomp. by therewith.

$\text{Fe}_3(\text{CN})_{12} + 4\text{H}_2\text{O} = \text{Fe}^{\text{III}}_2\text{Fe}^{\text{II}}[\text{Fe}(\text{CN})_6]_2 + 2\text{H}_2\text{O}$. Properties as above. (Roy, Chem. Soc. 54, 767.)

Iron (ferrous) potassium ferricyanide, $\text{KFe}_2(\text{CN})_{12} = \text{KFeFe}(\text{CN})_6 + 4$, or
(Soluble Prussian blue.) Sol. in H_2O insol. in salts + Aq. or alcohol.

the same composition, called "Wil-
lue," is insol. in H₂O.

ferrocyanide, basic, Pb₃[Fe(CN)₆]₂,
H₂+11H₂O.
)

ferrocyanide, Pb₃[Fe(CN)₆]₂+16H₂O.
1 H₂O; more sol. in hot, than cold
decomp. on boiling. (Gmelin.)
Easily sol. in H₂O; sl. sol. in
Schuler, W. A. B. 77. 692.)

potassium ferricyanide, PbKFe(CN)₆.
)
75 pts. H₂O at 16°, and the solu-
p. on standing. (Schuler.)
) Efflorescent. Much more sol.
in the Pb salt. Insol. in alcohol.
(.)

ferrocyanide nitrate, Pb₃[Fe(CN)₆]₂,
(NO₃)₂+12H₂O.
3.31 pts. H₂O at 16°. (Schuler.)
(Joannis, A. ch. (5) 26. 528.)

potassium ferricyanide, Mg₃[Fe(CN)₆]₂.
H₂O.

potassium ferricyanide,
K₃Fe(CN)₆.
(J. pr. 103. 166.)

potassium ferricyanide, Mn₃[Fe(CN)₆]₂.
H₂O, acids, NH₄OH, or NH₄ salts

potassium ferricyanide, Hg₃Fe(CN)₆.
(Mernekes, J. Am. Chem. Soc. 1906,

potassium ferricyanide, Hg₃[Fe(CN)₆]₂.
in H₂O. Solution quickly decomp.
J. Am. Chem. Soc. 1906, 28. 603.)

potassium ferricyanide ammonia, Ni₃[Fe(CN)₆]₂,
+H₂O.
[H₄OH+Aq. (Reynoso, A. ch.(3)

potassium ferricyanide, Ni₃[Fe(CN)₆]₂(?).
sol. in HCl+Aq.

potassium ferricyanide, K₃Fe(CN)₆, (or
(CN)₁₂).
Easily sol. in H₂O.

H₂O dissolve pts. K₃Fe(CN)₆ at t°.

s. l.t	t°	Pts. salt	t°	Pts. salt
0	15.6	40.8	100	77.5
6	37.8	58.8	104.4	82.6

(Wallace, Chem. Soc. 7. 80.)

100 pts. H₂O at 13° dissolve 38 pts., and the
solution has sp. gr. = 1.1630. (Schiff, A. 113.
350.)

1 l. sat. solution in H₂O at 25° contains
385.5 g. K₃Fe(CN)₆. (Grube, Z. Electrochem.
1914, 20. 342.)

Sp. gr. of K₃Fe(CN)₆+Aq at 13°.

% salt	Sp. gr.	% salt	Sp. gr.	% salt	Sp. gr.
1	1.0051	11	1.0595	21	1.1202
2	1.0103	12	1.0653	22	1.1266
3	1.0155	13	1.0712	23	1.1331
4	1.0208	14	1.0771	24	1.1396
5	1.0261	15	1.0831	25	1.1462
6	1.0315	16	1.0891	26	1.1529
7	1.0370	17	1.0952	27	1.1596
8	1.0426	18	1.1014	28	1.1664
9	1.0482	19	1.1076	29	1.1732
10	1.0538	20	1.1039	30	1.1802

(Schiff.)

Sp. gr. of K₃Fe(CN)₆+Aq at 25°.

Concentration of K ₃ Fe(CN) ₆ +Aq.	Sp. gr.
1—normal	1.0574
1/2—"	1.0289
1/4—"	1.0143
1/8—"	1.0092

(Wagner, Z. phys. Ch. 1890, 5. 37.)

Sat. K₃Fe(CN)₆+Aq boils at 104.4°.
(Wallace.)

1 l. sat. solution at 25° of K₃Fe(CN)₆+
K₄Fe(CN)₆ contains 338.1 g. K₃Fe(CN)₆ and
79.02 g. K₄Fe(CN)₆. (Grube.)

Solubility of K₃Fe(CN)₆+K₄Fe(CN)₆ in
KOH+Aq at 25°.

KOH Normality	g. per l.	
	K ₃ Fe(CN) ₆	K ₄ Fe(CN) ₆
0.4687	309	66.64
0.9628	275.3	55.19
1.949	200.8	35.95

(Grube.)

Solubility in KOH+Aq at 25°.

KOH Normality	g. K ₃ Fe(CN) ₆ per l.
0.4687	342.7
0.9628	302.3
1.949	215.1

(Grube, Z. Electrochem, 1914, 20. 342.)

Insol. in liquid NH₃. (Franklin, Am. Ch.
J. 1898, 20. 828.)

Insol. in absolute alcohol, and only sl. sol. in dil. alcohol.

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Potassium sodium ferricyanide,
 $\text{KNa}_2\text{Fe}(\text{CN})_6$.

Sol. in H_2O .

$\text{K}_2\text{NaFe}(\text{CN})_6$. Sol. in H_2O .

$\text{K}_2\text{Na}_3[\text{Fe}(\text{CN})_6]_2$. Sol. in H_2O .
 $+3\text{H}_2\text{O}$.

Potassium ferricyanide iodide, $\text{K}_3\text{Fe}(\text{CN})_6$,
 KI .

Very unstable.

Silver ferricyanide, $\text{Ag}_3\text{Fe}(\text{CN})_6$.

1 l. H_2O dissolves 0.00066 g. $\text{Ag}_3\text{Fe}(\text{CN})_6$ at 20° . (Whitby, Z. anorg. 1910, 67. 108.)

Sol. in NH_4OH , and hot $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$, but insol. in NH_4 salts + Aq .

Insol. in $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$. (Wackenroder, A. 41. 317.)

Silver ferricyanide ammonia, $2\text{Ag}_3\text{Fe}(\text{CN})_6$,
 $3\text{NH}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Gintl.)

$2\text{Ag}_3\text{Fe}(\text{CN})_6$, 5NH_3 . (Carlo, Gazz. ch. it. 1910, 40. (2) 477.)

Sodium ferricyanide, $\text{Na}_3\text{Fe}(\text{CN})_6 + \text{H}_2\text{O}$.

Deliquescent. Sol. in 5.3 pts. cold, and 1.5 pts. boiling H_2O . Insol. in alcohol, but not pptd. thereby from aqueous solution. (Bette.)

Ferrinitrososulphydic acid.

See Ferroheptanitrososulphydic acid.

Ferrocyanhydric acid, $\text{H}_4\text{Fe}(\text{CN})_6$.

Sol. in H_2O and alcohol.

100 pts. H_2O dissolve 15 pts. acid at 14° . (Joannis, A. ch. (5) 26. 514.)

Insol. in ether, and much less sol. in ether-alcohol than in alcohol. Insol. in conc. $\text{HCl} + \text{Aq}$.

Ferrocyanides.

The ferrocyanides of the alkali and alkaline-earth metals are sol. in H_2O ; the others are insol., but sol. in alkalis + Aq in case the base is sol. therein.

Aluminum ferrocyanide, $\text{Al}_4[\text{Fe}(\text{CN})_6]_3 + 17\text{H}_2\text{O}$.

Sl. sol. in H_2O .

Sl. sol. in $\text{HCl} + \text{Aq}$ with partial decomp. (Wyrouboff, A. ch. (5) 8. 446.)

Ammonium ferrocyanide, $(\text{NH}_4)_4\text{Fe}(\text{CN})_6 \cdot 3\text{H}_2\text{O}$.

Very sol. in H_2O ; insol. in alcohol.

$+ \text{H}_2\text{O}$. (Berzelius.)

Ammonium cadmium ferrocyanide ammonia,
 $(\text{NH}_4)_2\text{Cd}_2[\text{Fe}(\text{CN})_6]_2 \cdot 2\text{NH}_3 + \text{H}_2\text{O}$.

Sol. in H_2O . (Wyrouboff, A. ch. (5) 413.)

Ammonium calcium ferrocyanide,
 $(\text{NH}_4)_2\text{CaFe}(\text{CN})_6$.

Sl. sol. in H_2O . (Kunheim and Zimman, Dingl. 252. 478.)

100 g. sat. solution in H_2O contain 0 g. at 16° . (Brown, J. phys. Ch. 1898, 2.

Ammonium cuprous ferro cyanide,
 $(\text{NH}_4)_2\text{Cu}_2\text{Fe}(\text{CN})_6$.

Insol. in H_2O and alcohol.

Decomp. in the air. (Messner, Z. anorg. 1895, 8. 382.)

Ammonium cupric ferrocyanide,
 $(\text{NH}_4)_2\text{CuFe}(\text{CN})_6$.

Ppt.

$+ x\text{H}_2\text{O}$. Very unstable. Insol. in H_2O .
decomp. by boiling H_2O . (Messner, Z. anorg. 1895, 8. 384.)

Ammonium lithium ferrocyanide,
 $(\text{NH}_4)_2\text{Li}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Wyrouboff, A. ch. (4) 270.)

Ammonium magnesium ferrocyanide,
 $(\text{NH}_4)_2\text{MgFe}(\text{CN})_6$.

1 l. sat. solution at 17° contains 2.4 g. $(\text{NH}_4)_2\text{MgFe}(\text{CN})_6$. (Robinson, Chem. 1909, 95. 1353.)

Ammonium manganous ferrocyanide,
 $(\text{NH}_4)_2\text{MnFe}(\text{CN})_6$.

Ppt. (Blum, Z. anal. 30. 284.)

Ammonium potassium ferrocyanide,
 $\text{NH}_4\text{K}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Easily sol. in cold, more easily in hot H_2O .
Insol. in alcohol.

$(\text{NH}_4)_2\text{K}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$. Sol. in H_2O .

Ammonium potassium ferrocyanide ammonium chloride, $(\text{NH}_4)_3\text{KFe}(\text{CN})_6$,
 $2\text{NH}_4\text{Cl}$.

Sol. in H_2O . (Étard, J. pr. (2) 31. 43.)

Ammonium ferrocyanide bromide,
 $(\text{NH}_4)_4\text{Fe}(\text{CN})_6 \cdot 2\text{NH}_4\text{Br}$.

Permanent. Very sol. in H_2O .

Ammonium ferrocyanide chloride,
 $(\text{NH}_4)_4\text{Fe}(\text{CN})_6 \cdot 2\text{NH}_4\text{Cl} + 3\text{H}_2\text{O}$.

Permanent. Very sol. in H_2O , but less so than NH_4Cl . (Bunsen.)

ferrocyanide, $\text{Sb}_4[\text{Fe}(\text{CN})_6]_3 + 3\text{H}_2\text{O}$. (Atterberg.)

ferrocyanide, $\text{Ba}_2\text{Fe}(\text{CN})_6 + 6\text{H}_2\text{O}$.

ent. Sl. sol. in H_2O .
184 pts. cold, and 116 pts. boiling
los, 1832); sol. in 1800 pts. cold
ett, 1814); sol. in 1920 pts. cold,
100 pts. boiling H_2O (Thomson);
10 pts. cold, and 100 pts. boiling
e's Dict.)

1000 pts. H_2O at 15° , and 100 pts. at
rouboff, A. ch. (4) 16. 292.)

HNO_3 , HCl , or conc. $\text{H}_2\text{SO}_4 + \text{Aq}$.

pric ferrocyanide, $\text{BaCuFe}(\text{CN})_6$.

1 H_2O . (Messner, Z. anorg. 1895,

potassium ferrocyanide,

$\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

8 pts. cold, and 9.5 pts. boiling H_2O
832); in 36.4 pts. H_2O at 14° , and
t b.-pt. (Mosander.)

re sol. in $\text{NH}_4\text{Cl} + \text{Aq}$ than in H_2O .
, insol. in conc. $\text{HCl} + \text{Aq}$. (Rose.)
. Sol. in 300 pts. H_2O at ord. temp.
ff.)

ferrocyanide, $\text{Bi}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

(?).
in pure H_2O . (Wyrouboff.)
 $[\text{Fe}(\text{CN})_6]_3$. Ppt. (Muir, Chem. Soc.

potassium ferrocyanide,
 $\text{Fe}(\text{CN})_6 + 7\text{H}_2\text{O}$, or $4\text{H}_2\text{O}$.

potassium ferrocyanide,
 $\text{Fe}(\text{CN})_6 + \text{H}_2\text{O}$.

1 H_2O .
given by Wyrouboff is
 $[\text{Fe}(\text{CN})_6]_4 + 11\text{H}_2\text{O}$ (?).

ferrocyanide, $\text{Ca}_2\text{Fe}(\text{CN})_6 + 12\text{H}_2\text{O}$.

l. in H_2O . Sol. in 0.66 pt. H_2O at
not pptd. by cooling, and is ap-
ess sol. in warm than cold H_2O .
ff, A. ch. (4) 16. 280.)

cuprous ferrocyanide,
 $\text{Fe}(\text{CN})_6$. (Messner, Z. anorg.
37.)

pric ferrocyanide, $\text{CaCuFe}(\text{CN})_6$.

1 H_2O . (Messner, Z. anorg. 1895

potassium ferrocyanide,
 $\text{Fe}(\text{CN})_6$.

in H_2O . (Kunheim and Zimmer-
gl. 252. 478.)

+ $3\text{H}_2\text{O}$. Sol. in 795 pts. H_2O at 15° , and
145 pts. at b-pt., with decomp. in the latter
case.

Sol. in dil., insol. in conc. $\text{HCl} + \text{Aq}$. Sol.
in HNO_3 of 1.2 sp. gr. (Mosander.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$.

Calcium sodium ferrocyanide,

$\text{CaNa}_4[\text{Fe}(\text{CN})_6]_2$.

Sol. in H_2O .

Calcium strontium ferrocyanide,

$\text{CaSrFe}(\text{CN})_6 + 10\text{H}_2\text{O}$.

Efflorescent. Sol. in about 3 pts. H_2O .
(Wyrouboff, A. ch. (4) 21. 278.)

Cerium ferrocyanide, $\text{Ce}_4[\text{Fe}(\text{CN})_6]_3 + 30\text{H}_2\text{O}$.

Ppt. (Wyrouboff.)

Cerium potassium ferrocyanide,

$\text{CeKFe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Ppt. (Jolin.)

+ $4\text{H}_2\text{O}$. (Wyrouboff.)

Chromic ferrocyanide, $\text{Cr}_2[\text{Fe}(\text{CN})_6]_3 + 20\text{H}_2\text{O}$.

Ppt.

Cobaltous ferrocyanide, $\text{Co}_2\text{Fe}(\text{CN})_6 + 7\text{H}_2\text{O}$.

Wholly insol. in H_2O .

Sol. in H_2SO_4 with decomp. Insol. in
 $\text{HCl} + \text{Aq}$. Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in
(NH_4) $_2\text{CO}_3 + \text{Aq}$. Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$. Sol.
in $\text{KCN} + \text{Aq}$.

Cobaltous ferrocyanide ammonia,

$\text{Co}_2\text{Fe}(\text{CN})_6, 8\text{NH}_3 + 10\text{H}_2\text{O}$.

Ppt. Decomp. on standing. (Curda, Z.
Ch. 1869. 369.)

$\text{Co}_2\text{Fe}(\text{CN})_6, 12\text{NH}_3 + 9\text{H}_2\text{O}$. As above.
(Curda.)

Cobaltous potassium ferrocyanide,

$\text{CoK}_2\text{Fe}(\text{CN})_6$.

Ppt. (Wyrouboff.)

$\text{Co}_2\text{K}_4[\text{Fe}(\text{CN})_6]_4$ (?). Ppt. Insol. only in
presence of an excess of $\text{K}_4\text{Fe}(\text{CN})_6$. (Wy-
rouboff.)

Columbium potassium ferrocyanide,

$\text{Cb}_{16}\text{K}[\text{Fe}(\text{CN})_6]_2 + 67\text{H}_2\text{O}$ (?).

Sol. in H_2O . (Wyrouboff.)

$\text{Cb}_{12}\text{K}_2\text{Fe}(\text{CN})_6 + 39\text{H}_2\text{O}$ (?). Sol. in
 H_2O . (W.)

(CbO) $_3\text{K}_3[\text{Fe}(\text{CN})_6]_6 + 10\text{H}_2\text{O}$ (?). Ppt. (At-
terberg.)

Cuprous ferrocyanide, $\text{Cu}_4\text{Fe}(\text{CN})_6$.

Insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$; insol.
in $\text{NH}_4\text{Cl} + \text{Aq}$.

Cupric ferrocyanide, basic, $\text{CuFe}(\text{OH})_4(\text{CN})_4$.
Ppt. (Bong, Bull. Soc. 23. 231.)

Cupric ferrocyanide, $\text{Cu}_2\text{Fe}(\text{CN})_6 + 7\text{H}_2\text{O}$.

Insol. in H_2O or acids. Insol. in NH_4 salts + Aq. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in $(\text{NH}_4)_2\text{C}_2\text{O}_4 + \text{Aq}$ and in $\text{KCN} + \text{Aq}$.

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 827.)

+ $10\text{H}_2\text{O}$. Sol. in excess of $\text{K}_4\text{Fe}(\text{CN})_6 + \text{Aq}$, especially if hot. (Wyruboff.)

Cupric ferrocyanide ammonia (cuprammonium ferrocyanide), $\text{Cu}_2\text{Fe}(\text{CN})_6, 4\text{NH}_3 + \text{H}_2\text{O}$.

Insol. in H_2O or alcohol. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Bunsen.)

$\text{Cu}_2\text{Fe}(\text{CN})_6, 8\text{NH}_3 + \text{H}_2\text{O}$.

Cuprous magnesium ferrocyanide, $\text{Cu}_2\text{MgFe}(\text{CN})_6$.

Very unstable. Decomp. in air.

Insol. in H_2O . (Messner, Z. anorg. 1895, 8. 385.)

Cupric magnesium ferrocyanide, $\text{CuMgFe}(\text{CN})_6$.

Insol. in H_2O . Decomp. by boiling H_2O . Very unstable. (Messner, Z. anorg. 1895, 8. 387.)

Cuprous potassium ferrocyanide, $\text{Cu}_2\text{K}_2\text{Fe}(\text{CN})_6$.

Insol. in H_2O . Decomp. by boiling H_2O . Decomp. by acids. Insol. in alcohol. (Messner, Z. anorg. 1895, 8. 378.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O , alcohol, or ether. Decomp. by acids. Sol. in $\text{KCN} + \text{Aq}$.

$\text{K}_2\text{Cu}_2\text{Fe}(\text{CN})_6 + 4\text{H}_2\text{O}$.

+ $5\text{H}_2\text{O}$. (Wonfor.)

+ $6\text{H}_2\text{O}$. (Wyruboff.)

Cupric potassium ferrocyanide, $\text{K}_2\text{CuFe}(\text{CN})_6 + \text{H}_2\text{O}$.

Insol. in cold, sl. decomp. by boiling H_2O . $\text{K}_2\text{Cu}_2[\text{Fe}(\text{CN})_6]_2 + 12\text{H}_2\text{O}$. Ppt.

Cuprous sodium ferrocyanide, $\text{Cu}_2\text{Na}_2\text{Fe}(\text{CN})_6$.

Decomp. by boiling H_2O ; insol. in alcohol; insol. in H_2O ; decomp. by acids. (Messner, Z. anorg. 1895, 8. 373.)

Cupric sodium ferrocyanide, $\text{CuNa}_2\text{Fe}(\text{CN})_6$.

Insol. in cold H_2O . Decomp. by boiling H_2O . (Moissan, Z. anorg. 1895, 8. 376.)

Cupric strontium ferrocyanide, $\text{CuSrFe}(\text{CN})_6$.

Insol. in H_2O . (Messner, Z. anorg. 1895, 8. 389.)

Didymium potassium ferrocyanide, $\text{DiKFe}(\text{CN})_6 + 4\text{H}_2\text{O}$.

Ppt. (Cleve.)

+ $2\text{H}_2\text{O}$. (Wyruboff.)

Erbium potassium ferrocyanide, $\text{ErK} + x\text{H}_2\text{O}$.

(Höglund.)

Gallium ferrocyanide.

Sol. in boiling $\text{HCl} + \text{Aq}$. (de Bois C. R. 99.526.)

Glucinum ferrocyanide, $\text{Gl}_2\text{Fe}(\text{CN})_6 + 7\text{H}_2\text{O}$ (?)

Sol. in H_2O . (Atterberg.)

Iron (ferric) ferrocyanide, $\text{Fe}_7(\text{CFe}_4[\text{Fe}(\text{CN})_6]_3 + x\text{H}_2\text{O}$.

(Prussian blue.) Insol. in H_2O , ether, or oils. Decomp. slowly by H_2O . Insol. in dil. mineral acids. conc. $\text{HCl} + \text{Aq}$, and conc. H_2SO_4 will comp. Sol. in $\text{H}_2\text{C}_2\text{O}_4$ or NH_4 tart. Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by or $\text{KOH} + \text{Aq}$. Not pptd. in presence of citrates or citrates.

Iron (ferrous) potassium ferrocyanide, $\text{FeK}_2\text{Fe}(\text{CN})_6$.

Insol. in H_2O . Decomp. on air.

Iron (ferric) potassium ferrocyanide, $\text{FeKFe}(\text{CN})_6$.

Is probably ferrous potassium ferrocyanide which see.

Iron (ferric) ferrocyanide ammonia, $\text{Fe}_4[\text{Fe}(\text{CN})_6]_3, 6\text{NH}_3 + 9\text{H}_2\text{O}$.

Insol. in NH_4 tartrate + Aq.

Lanthanum potassium ferrocyanide, $\text{LaKFe}(\text{CN})_6 + 4\text{H}_2\text{O}$.

Ppt.

Lead ferrocyanide, $\text{Pb}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Insol. in H_2O , acids, or NH_4 (Wyruboff, A. ch. (5) 8. 480.)

Sl. sol. in conc. H_2SO_4 , from which pptd. by H_2O . (Berzelius.)

Sol. in hot NH_4Cl , or NH_4 succin. insol. in other NH_4 salts + Aq. (Wi

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Brett.)

Not pptd. in presence of Na citrate (Spiller.)

Lithium ferrocyanide, $\text{Li}_2\text{Fe}(\text{CN})_6 + \text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O .

Lithium potassium ferrocyanide, $\text{Li}_2\text{K}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Very sol. in H_2O . Sol. in 1.5 pt. ord. temp. (Wyruboff, A. ch. (4)

m ferrocyanide, $Mg_3Fe(CN)_6 + 3$ pts. cold H_2O . (Bette, A. 22.

m potassium ferrocyanide, $K_4Fe(CN)_6$. 575 pts. H_2O at 15° , and 238 pts. at solution is decomp. by boiling. (Dict.) solution at 17° contains 1.95 g. $(CN)_6$. (Robinson, Chem. Soc. 353.)

s ferrocyanide, $Mn_3Fe(CN)_6 + H_2O$. Sol. in $HCl + Aq$. Insol. in $NH_4NO_3 + Aq$.

ferrocyanide, $Mn_3Fe_3(CN)_{12}$. H_2O . Easily decomp. in the air. (Straus, Z. anorg. 1895, 9. 8.)

s potassium ferrocyanide, $K_4Fe(CN)_6$. (Berzelius.) $(CN)_6$, $4K_4Fe(CN)_6 + 4H_2O(?)$. in dil. $HCl + Aq$. (Wyrouboff.)

potassium ferrocyanide, $K_4Fe(CN)_6$. in H_2O . Appreciably sol. in $+ Aq$. (Fernekas, J. Am. Chem. 28. 87.)

um ferrocyanide, $Mo_4Fe(CN)_6 + (?)$. l. in $NH_4OH + Aq$. (Wyrouboff.) $(CN)_6 + 8H_2O (?)$. (W.) $(?)$. Very sol. in H_2O ; insol. in W .)

um potassium ferrocyanide, $[Fe(CN)_6]_2 + 40H_2O (?)$. (Wyrouboff.) $[Fe(CN)_6]_2, 2MoO_3 + 20H_2O (?)$. $[Fe(CN)_6]_2, 2MoO_3 + 12H_2O (?)$.

rocyanide, $Ni_3Fe(CN)_6 + 11H_2O$, or H_2O . Sol. in H_2O or $HCl + Aq$. Sol. in Aq ; insol. in NH_4 salts $+ Aq$. Sol. in Aq .

rocyanide ammonia, $Ni_3Fe(CN)_6 + H_2O$. Slightly insol. in H_2O and not attacked by $NH_4OH + Aq$ to form— $N)_6, 10NH_3 + 4H_2O$. Decomp. by (Reynoso, A. ch. (3) 30. 252.) $N)_6, 2NH_3 + 4$, and $9H_2O$. Hygro-

scopic. Easily decomp. (Gintl, J. B. 1868. 304.)

$Ni_3Fe(CN)_6, 8NH_3 + 4H_2O$. Sol. in $NH_4OH + Aq$. (G.)

$Ni_3Fe(CN)_6, 12NH_3 + 9H_2O$. Sol. in $NH_4OH + Aq$, but less so than the above compounds. (G.)

Nickel potassium ferrocyanide, $NiK_2Fe(CN)_6 + 3H_2O$.

Ppt. (Wyrouboff.)

Osmium ferrocyanide, $Os_2Fe(CN)_6$.

Ppt. (Martius, A. 117. 368.)

Potassium ferrocyanide, $K_4Fe(CN)_6$.

Permanent. Easily sol. in cold, and more easily in hot H_2O .

Sol. in 4.23 pts. H_2O at 15° , or 100 pts. H_2O dissolve 23.6 pts. salt at 15° . (Schiff, A. 113. 350.)

100 pts. H_2O dissolve 27.8 pts. at 12.2° ; 65.8 pts. at 37.7° ; 87.6 pts. at 65.5° ; and 90.6 pts. at 96.3° . (Thomson.)

Sol. in 4 pts. cold, and 2 pts. boiling H_2O . (Wittstein.)

100 pts. H_2O dissolve 29.2 pts. salt at 15° , and solution has sp. gr. = 1.1441. (Michel and Kraft, A. ch. (3) 41. 478.)

Solubility of $K_4Fe(CN)_6$ in H_2O at t° .

-2°	$+7^\circ$	14°	30°	56°
10.8	15.4	17.9	23.0	31.7%
60°	75°	89°	98°	157°
34.0	39.1	41.9	42.6	46.8%

(Étard, A. ch. 1894, (7) 2. 546.)

$K_4Fe(CN)_6 + Aq$ sat. at 8° has sp. gr. = 1.13. (Anthon.)

Sp. gr. of $K_4Fe(CN)_6 + Aq$ at 15° .

% hydrous salt	Sp. gr.	% hydrous salt	Sp. gr.	% hydrous salt	Sp. gr.
1	1.0058	8	1.0479	15	1.0932
2	1.0116	9	1.0542	16	1.0999
3	1.0175	10	1.0605	17	1.1067
4	1.0234	11	1.0669	18	1.1136
5	1.0295	12	1.0734	19	1.1205
6	1.0356	13	1.0800	20	1.1275
7	1.0417	14	1.0866

(Schiff, A. 113. 199.)

Sp. gr. of $K_4Fe(CN)_6 + Aq$ at 25° .

Concentration of $K_4Fe(CN)_6 + Aq$.	Sp. gr.
1—normal	1.0617
$1/2$ —“	1.0300
$1/4$ —“	1.0150
$1/8$ —“	1.0074
$1/16$ —“	1.0037

(Wagner, Z. phys. Ch. 1890, 5. 37.)

Solubility in KOH + Aq at 25°.

KOH Normality	g. $K_4Fe(CN)_6 + 3H_2O$ per l.
0.09984	308.5
0.2496	283.5
0.4963	247.1
0.7036	217.4
0.9415	184.8
1.395	132.1
1.883	86.12

(Grube, Z. Electrochem, 1914, 20. 342.)

$K_4Fe(CN)_6 + NaCl + Aq$ sat. at 20° contains 26.6 g. NaCl and 17.8 g. $K_4Fe(CN)_6$ per 100 g. H_2O ; sat. at 93° it contains 27.4 g. NaCl and 35.9 g. $K_4Fe(CN)_6$ per 100 g. H_2O . (Conroy, J. Soc. Chem. Ind. 1898, 17. 105.)

$K_4Fe(CN)_6 + KCl + Aq$ sat. at 21° contains 27.2 g. KCl and 4.2 g. $K_4Fe(CN)_6$ per 100 g. H_2O ; sat. at 99° it contains 39.6 g. KCl and 17.0 g. $K_4Fe(CN)_6$ per 100 g. H_2O . (Conroy.)

$K_4Fe(CN)_6 + Na_2CO_3 + Aq$ sat. at 22° contains 29.9 g. Na_2CO_3 and 26.7 g. $K_4Fe(CN)_6$ per 100 g. H_2O ; sat. at 97° it contains 42.0 g. Na_2CO_3 and 27.5 g. $K_4Fe(CN)_6$ per 100 g. H_2O . (Conroy.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in alcohol even when dilute.

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

+ $3H_2O$. 1 l. sat. solution in H_2O contains 319.4 g. $K_4Fe(CN)_6 + 3H_2O$. (Grube, Electrochem. Z. 1914, 20. 342.)

Two modifications with different solubilities.

25.0 g. of α modification are contained in 100 g. of solution at 20°.

24.6 g. of β modification are contained in 100 g. of solution at 20°. (Briggs, Chem. Soc. 1911, 99. 1024.)

32.0 g. $K_4Fe(CN)_6$ (anhydrous) are dissolved in 100 g. H_2O at 25°. (Wagner, Z. phys. Ch. 1910, 71. 428.)

Potassium samarium ferrocyanide,
 $KSmFe(CN)_6 + 5H_2O$.

Precipitate. (Cleve.)

Potassium sodium ferrocyanide,
 $KNa_3Fe(CN)_6 + 12H_2O$.

Sol. in H_2O .

$K_2Na_2Fe(CN)_6 + 8H_2O$. Easily sol. in H_2O .

$K_3NaFe(CN)_6 + 3H_2O$. Permanent. Easily sol. in H_2O ; insol. in alcohol.

Potassium sodium ferrocyanide nitrate,
 $K_2Na_2Fe(CN)_6 \cdot 4KNO_3$.

Sol. in H_2O . (Martius.)

Potassium strontium ferrocyanide,
 $K_2SrFe(CN)_6 + 3H_2O$.

Easily decomp. Sol. in H_2O ; sl. sol. alcohol. (Wyrouboff, A. ch. (4) 21. 276.)

Potassium stannic ferrocyanide,
 $KSn_3[Fe(CN)_6]_2 + 68H_2O$ (?).

Ppt. (Wyrouboff.)

$K_4Sn_{10}[Fe(CN)_6]_{11} + 230H_2O$ (?). (Atterberg.)

Potassium titanium ferrocyanide,
 $K_2Ti_2[Fe(CN)_6]_3 + 11H_2O$ (?).

Ppt. Sol. in $K_4Fe(CN)_6 + Aq$. (Wyrouboff.)

$K_4Fe(CN)_6$, $11Ti_2Fe(CN)_6 + 43H_2O$

Ppt. (Wyrouboff.)
 $K_2(TiO)_2[Fe(CN)_6]_2 + 23H_2O$ (?).

(Atterberg.)
 $K_2(TiO)_{11}[Fe(CN)_6]_6 + 110H_2O$ (?).

Potassium tungsten ferrocyanide,
 $KW_2Fe(CN)_6 + 7H_2O$ (?).

Sol. in H_2O . (Wyrouboff.)

$K_2W_2Fe(CN)_6 + 20H_2O$ (?). Sol. in (W.)

Potassium uranium ferrocyanide,
 $K_2U_2[Fe(CN)_6]_2 + 6H_2O$ (?).

Ppt. (Wyrouboff.)

$K_2(UO_2)_2[Fe(CN)_6]_2 + 6H_2O$. Ppt. (Atterberg.)

$K_6(UO_2)_3[Fe(CN)_6]_4 + 12H_2O$. Sol. in (Atterberg.)

Potassium vanadium ferrocyanide,
 $K_{10}V[Fe(CN)_6]_5 + 39H_2O$ (?).

Ppt. Sl. sol. in H_2O . (Wyrouboff.)

$K_6(VO)_3[Fe(CN)_6]_4 + 60H_2O$ (?). Ppt. (Atterberg.)

Potassium ytterbium ferrocyanide,
 $KYbFe(CN)_6 + 3H_2O$.

Ppt. Sol. in excess $K_4Fe(CN)_6$. (Cleve, Z. anorg. 1902, 32. 140.)

Potassium yttrium ferrocyanide,
 $KYFe(CN)_6 + 2H_2O$.

Ppt. (Wyrouboff, A. ch. (5) 8. 444.)

Potassium zinc ferrocyanide,
 $K_4Zn_2[Fe(CN)_6]_3 + 12H_2O$.

Absolutely insol. in H_2O . (Wyrouboff, ch. (5) 8. 485.)

Potassium ferrocyanide carbonyl,
 $K_3Fe(CN)_5(CO) + 3\frac{1}{2}H_2O$.

See Carbonyl ferrocyanide, potassium

Rubidium ferrocyanide, $Rb_4Fe(CN)_6 + 3H_2O$
Sol. in less than 1 pt. H_2O at ord. with great absorption of heat. (Wyrouboff, A. ch. (4) 16. 307.)

Iron ferrocyanide, $\text{Ag}_4\text{Fe}(\text{CN})_6 + \text{H}_2\text{O}$.

Insol. in H_2O or dil. acids. Insol. in LiOH , or NH_4 salts + Aq. Sol. in KCN Aq.

Decomp. by warm $\text{NH}_4\text{OH} + \text{A-}$. (Weith, Ch. (2) 5. 381.)

Iron ferrocyanide ammonia,
 $\text{Ag}_4\text{Fe}(\text{CN})_6, 2\text{NH}_3 + \text{H}_2\text{O}$.

(Wyruboff.)
+ $6\text{H}_2\text{O}$. (Gintl.)

Sodium ferrocyanide, $\text{Na}_4\text{Fe}(\text{CN})_6 + 12\text{H}_2\text{O}$.

Efflorescent. Less sol. in H_2O than $\text{Fe}(\text{CN})_6$. Sol. in 4.5 pts. H_2O at 12° . (ohn.)

100 pts. H_2O at 15.5° dissolve 22 pts. (Weith's Dict.)

100 pts. H_2O dissolve at:

20°	42°	53°	
17.875	30.2	37.1	pts. Na_4FeCN_6
60°	77°	80°	
42.5	54.8	59.2	pts. Na_4FeCN_6
98°	98°	98.5°	
62.1	61.6	6.30	pts. Na_4FeCN_6

(Conroy, J. Soc. Chem. Ind. 1898, 17. 104.)

+ $10\text{H}_2\text{O}$.

100 pts. H_2O dissolve at:

20°	42°	
31.85	58.5	pts. $\text{Na}_4\text{Fe}(\text{CN})_6 + 10\text{H}_2\text{O}$
58°	60°	
88.4	90.2	pts. $\text{Na}_4\text{Fe}(\text{CN})_6 + 10\text{H}_2\text{O}$
80°	96°	
146.0	157.0	pts. $\text{Na}_4\text{Fe}(\text{CN})_6 + 10\text{H}_2\text{O}$
	98.5°	
161.0		pts. $\text{Na}_4\text{Fe}(\text{CN})_6 + 10\text{H}_2\text{O}$

(Conroy.)

Strontium ferrocyanide, $\text{Sr}_2\text{Fe}(\text{CN})_6 + 15\text{H}_2\text{O}$.

Efflorescent. Sol. in 2 pts. cold, and less in 1 pt. boiling H_2O . (Bette.)

Excessively sol. in H_2O . (Wyruboff, A. (4) 16. 280.)

+ $8\text{H}_2\text{O}$. (Wyruboff.)

Thallous ferrocyanide, $\text{Tl}_4\text{Fe}(\text{CN})_6 + 2\text{H}_2\text{O}$.

100 pts. H_2O dissolve 0.37 pt. at 18° , and 3 pts. at 101° . (Lamy.)

Insol. in KCN + Aq. (Kühlmann.)

Thorium ferrocyanide, $\text{ThFe}(\text{CN})_6 + 4\text{H}_2\text{O}$.

Ppt. (Cleve, Bull. Soc. (2) 24. 355.)

(stannous) ferrocyanide, $\text{Sn}_2\text{Fe}(\text{CN})_6 + 4\text{H}_2\text{O}$.

Insol. in H_2O or acids; sl. sol. in $\text{NH}_4\text{OH} + \text{A-}$. (Wyruboff.)

(stannic) ferrocyanide, $\text{Sn}_2[\text{Fe}(\text{CN})_6]_2 + 18\frac{1}{2}\text{H}_2\text{O}$ (?).

(Wyruboff.)

Titanium ferrocyanide, $\text{Ti}_7[\text{Fe}(\text{CN})_6]_2$ (?).

Ppt. (Wyruboff.)

Uranium ferrocyanide, $\text{UFe}(\text{CN})_6 + 10\text{H}_2\text{O}$.

Ppt. (Wyruboff.)

Vanadyl ferrocyanide, $(\text{VO})_2\text{Fe}(\text{CN})_6 + 11\text{H}_2\text{O}$.

Ppt. (Atterberg.)

Yttrium ferrocyanide, $\text{Y}_4[\text{Fe}(\text{CN})_6]_3$.

Easily sol. in H_2O ; insol. in alcohol. (Popp, A. 131. 179.)

Zinc ferrocyanide, $\text{Zn}_2\text{Fe}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Insol. in H_2O or acids.

Insol. in $\text{HCl} + \text{Aq}$. (Lea, Sill. Am. J. (2) 31. 191.)

Sol. in NH_4OH , or NH_4 salts + Aq. (Wittstein.)

Insol. in NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett.)

Sl. sol. in boiling $\text{K}_4\text{Fe}(\text{CN})_6$, or $\text{K}_2\text{Fe}(\text{CN})_6 + \text{Aq}$. (Gore.)

$\text{Na}_4\text{Fe}(\text{CN})_6 + \text{NaCl} + \text{Aq}$ sat. at 21° contains 29.0 g. NaCl and 5.8 g. $\text{Na}_4\text{Fe}(\text{CN})_6$ per 100 g. H_2O ; sat. at 90° it contains 24.7 g. NaCl and 21.3 g. $\text{Na}_4\text{Fe}(\text{CN})_6$ per 100 g. H_2O . (Conroy, J. Soc. Chem. Ind. 1898, 17. 105.)

$\text{Na}_4\text{Fe}(\text{CN})_6 + \text{Na}_2\text{CO}_3 + \text{Aq}$ sat. at 22° contains 22.6 g. Na_2CO_3 and 6.5 g. $\text{Na}_4\text{Fe}(\text{CN})_6$ per 100 g. H_2O ; sat. at 95° it contains 29.8 g. Na_2CO_3 and 36.8 g. $\text{Na}_4\text{Fe}(\text{CN})_6$ per 100 g. H_2O . (Conroy.)

Very sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 830.)

Insol. in alcohol.

+ $4\text{H}_2\text{O}$. Absolutely insol. in H_2O . (Wyruboff, A. ch. (5) 8. 485.)

+ $8\text{H}_2\text{O}$. (Weith, A. 147. 329.)

+ $10\text{H}_2\text{O}$. (Pebal, A. 233. 165.)

Ferrotetranitrososulphydic acid,

$\text{H}_2\text{S}_2(\text{NO})_4\text{Fe}_2$.

Insol. in H_2O ; sl. sol. in alcohol; more easily in ether; very sol. in CS_2 or CHCl_3 . Not obtained in a pure state. (Pawel, B. 15. 2600.)

Ethyl ferrotetranitrososulphide,

$(\text{C}_2\text{H}_5)_2\text{S}_2(\text{NO})_4\text{Fe}_2$.

Insol. in H_2O , difficultly sol. in alcohol, more easily in ether, and very easily in CS_2 , CHCl_3 , $\text{C}_2\text{H}_5\text{I}$, or C_6H_6 . (Pawel, B. 15. 2609.)

Ferrous —, $\text{FeS}_2(\text{NO})_4\text{Fe}_2$.

More difficultly sol. in H_2O and alcohol than the *hepta* salt.

Sol. in ether.

Potassium —, $\text{K}_2\text{S}_2(\text{NO})_4\text{Fe}_2 + 4\text{H}_2\text{O}$.

Sol. in H_2O . Easily sol. in alcohol; insol. in ether. (Pawel, B. 15. 2600.)

True composition of "nitrosulphide of

iron and potassium" of Roussin. (A. ch. (3) 52. 297.) (Pawel, B. 13. 1949.)

Sodium ferrotetranitrososulphide,
 $\text{Na}_4\text{S}_2(\text{NO})_4\text{Fe}_2 + 8\text{H}_2\text{O}$.

Sol. in H_2O ; easily sol. in alcohol; insol. in ether. (Pawel.)

True composition of "nitrosulphide of iron and sodium" of Roussin. (Pawel.)

Thallium —, $\text{Tl}_2\text{S}_2(\text{NO})_4\text{Fe}_2$.

Insol. in H_2O , alcohol, or ether. (Pawel.)

Ferroheptanitrososulphydic acid,
 $\text{HS}_2(\text{NO})_7\text{Fe}_4$.

Insol. in H_2O , alcohol, and ether. Easily sol. in CS_2 or CHCl_3 . (Pawel, B. 15. 2604.)

May be called Ferrinitrososulphydic acid.

Ammonium ferroheptanitrososulphide,
 $\text{NH}_4\text{S}_2(\text{NO})_7\text{Fe}_4 + \text{H}_2\text{O}$.

Less easily sol. in H_2O than the K compound. (Pawel, B. 15. 2600.)

"Binitrosulphide of iron" of Roussin. Sol. in about 2 pts. boiling H_2O ; very sl. sol. in cold H_2O . Very sol. in alcohols, methyl, ethyl, or amyl, and in $\text{HC}_2\text{H}_5\text{O}_2$. Miscible with ether. Insol. in CS_2 or CHCl_3 .

Decomp. by conc. HCl , HNO_3 , or H_2SO_4 . Not attacked by $\text{H}_2\text{C}_2\text{O}_4$, or $\text{H}_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq}$.

Insol. in NH_4OH , and $\text{KOH} + \text{Aq}$. (Roussin, A. ch. (3) 52. 286.)

Sol. in H_2O . Insol. in alcohol. (Hofmann, Z. anorg. 1895, 9. 299.)

Barium —.

Easily sol. in H_2O . (Pawel.)

Cæsium —, $\text{Fe}_4(\text{NO})_7\text{S}_2\text{Cs} + \text{H}_2\text{O}$.

Insol. in H_2O . Difficultly sol. in alcohol and ether. (Pawel.)

Sparingly sol. in H_2O . (Hofmann, Z. anorg. 1895, 9. 298.)

Calcium —.

Easily sol. in H_2O . (Pawel.)

Ferrous —, $\text{Fe}[\text{S}_2(\text{NO})_7\text{Fe}_4]_2 + 8\text{H}_2\text{O}$.

More easily sol. in H_2O than Na salt. (Pawel.)

Lead —.

Difficultly sol. in H_2O . (Pawel.)

Magnesium —.

Easily sol. in H_2O . (Pawel.)

Potassium —, $\text{KS}_2(\text{NO})_7\text{Fe}_4$.

Sol. in H_2O , alcohol, and very sol. in ether with slight decomp. (Pawel, B. 15. 2600.)

Rubidium ferroheptanitrososulphide,
 $\text{RbS}_2(\text{NO})_7\text{Fe}_4$.

Less soluble in H_2O than the NH_4 (Pawel.)

+ H_2O . Ppt. (Hofmann, Z. anorg. 1895, 9. 298.)

Sodium —, $\text{NaS}_2(\text{NO})_7\text{Fe}_4 + 2\text{H}_2\text{O}$.

More sol. in H_2O than the potassium (Pawel.)

Thallium —, $\text{TlS}_2(\text{NO})_7\text{Fe}_4 + \text{H}_2\text{O}$.

Very difficultly sol. in H_2O . More sol. in alcohol. (Pawel.) (Hofmann, anorg. 1895, 9. 297.)

Ferrodinitrosothiosulphonic acid.

Ammonium ferrodinitrosothiosulphonate,
 $\text{Fe}(\text{NO})_2\text{S}_2\text{O}_3\text{NH}_4 + \text{H}_2\text{O}$.

Can be cryst. from warm H_2O without comp. (Hofmann, Z. anorg. 1895, 8. 321)

Cæsium —, $\text{Fe}(\text{NO})_2\text{S}_2\text{O}_3\text{Cs}$.

Sparingly sol. in H_2O . (Hofmann.)

Potassium —, $\text{Fe}(\text{NO})_2\text{S}_2\text{O}_3\text{K} + \text{H}_2\text{O}$.

Sl. sol. in H_2O without decomp. at

Sol. in 50% alcohol.

Sol. in H_2SO_4 without decomp. (Hofmann.)

Rubidium —, $\text{Fe}(\text{NO})_2\text{S}_2\text{O}_3\text{Rb} + \text{H}_2\text{O}$.

Less sol. in H_2O than the corresponding Na salt. (Hofmann.)

Sodium —, $\text{Fe}(\text{NO})_2\text{S}_2\text{O}_3\text{Na} + 2\text{H}_2\text{O}$.

Closely resembles K salt, but is more in H_2O and alcohol. (Hofmann.)

Ferrotungstic acid.

Sol. in H_2O . (Laurent, C. R. 31. 693.)

Ammonium manganous ferrotungstate,
 $12(\text{NH}_4)_2\text{O}, 6\text{MnO}, 2\text{Fe}_2\text{O}_3, 3\text{H}_2\text{O}, 45\text{WO}_3 + 81\text{H}_2\text{O}$.

Sol. in H_2O . (Laurent.)

Barium ferrotungstate, $21\text{BaO}, 2\text{Fe}_2\text{O}_3, 45\text{WO}_3 + 27\text{H}_2\text{O}$.

Sol. in H_2O . (Laurent.)

Potassium ferrotungstate, $9\text{K}_2\text{O}, 2\text{Fe}_2\text{O}_3, 12\text{H}_2\text{O}, 45\text{WO}_3 + 54\text{H}_2\text{O}$.

Sol. in H_2O . (Laurent.)

$18\text{K}_2\text{O}, 2\text{Fe}_2\text{O}_3, 3\text{H}_2\text{O}, 45\text{WO}_3 + 54\text{H}_2\text{O}$ (Laurent.)

Ferrous acid.

Barium ferrite, $\text{BaO}, \text{Fe}_2\text{O}_3$.

Ppt. (List, B. 11. 1512.)

ferrite, $4\text{CaO}, \text{Fe}_2\text{O}_3$.
in H_2O , or sugar + H_2O . Decomp.
weakest acids, but not by boiling
aq. (Pelouze, A. ch. (3) 33. 5.)

Fe_2O_3 . (List.)

Fe_2O_3 . Much less readily attacked
and acids than the silicates. (Hilpert,
42. 4581.)

$2\text{Fe}_2\text{O}_3$. As above. (Hilpert, B.
4581.)

ferrite chloride, $\text{CaO}, \text{Fe}_2\text{O}_3, \text{CaCl}_2$.
decomp. by H_2O . (Chatelier, C. R. 99.)

ferrite, $\text{CuO}, \text{Fe}_2\text{O}_3$.

(List.)

O. (List.)

argentous ferrite, $2\text{FeO}, \text{Ag}_2\text{O}$,
, (?).

decomp. by $\text{HCl} + \text{Aq}$. Not com-
ol. in dil. $\text{HNO}_3 + \text{Aq}$. Easily sol. in
 HNO_3 . Decomp. by acetic acid.
Pogg. 10. 323.)

um ferrite, $\text{MgO}, \text{Fe}_2\text{O}_3$.

in H_2O . Not attacked by boiling
 HNO_3 . (Deville. C. R. 52. 1264.)

Magnesioferrite. Difficultly sol. in
aq. (Rammelsberg, Pogg. 107. 451.)

O. Ppt. (List, B. 11. 1512.)

, $\text{Fe}_2\text{O}_3 + 9\text{H}_2\text{O}$. Ppt.

, H_2O . Min. *Pyroaurite*.

ous ferrite, $\text{MnO}, \text{Fe}_2\text{O}_3$.

(List.)

ferrite, $\text{NiO}, \text{Fe}_2\text{O}_3$.

(List.)

m ferrite, $3\text{K}_2\text{O}, 4\text{Fe}_2\text{O}_3$.

mp. by H_2O , $\text{KOH} + \text{Aq}$, $\text{NaOH} + \text{Aq}$,
only slowly by $\text{NH}_4\text{Cl} + \text{Aq}$. (Salm-
mar, J. pr. 55. 349.)

O_4 . Decomp. by H_2O . (Rousseau
nheim, C. R. 107. 240.)

argentous) ferrite, $\text{Ag}_2\text{O}, \text{Fe}_2\text{O}_3$ (?).
mp. by dil. $\text{HNO}_3 + \text{Aq}$. (Rose, Pogg.

ferrite, $\text{Na}_2\text{O}, \text{Fe}_2\text{O}_3$.

is dissolved out by H_2O . Easily sol.
 $\text{HCl} + \text{Aq}$. Not easily decomp. by
- Aq . (Salm-Horstmar.)

rite, $\text{ZnO}, \text{Fe}_2\text{O}_3$.

n boiling conc. $\text{HCl} + \text{Aq}$. (Ebel-
ch. (3) 33. 47.)

Franklinite.

baltic compounds.

so Xanthocobaltic compounds.

Flavocobaltic chloraurate,
 $(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4\text{AuCl}_4$.

More easily sol. than the chloroplatinate.
Not wholly insol. in absolute alcohol. (Jör-
gensen, Z. anorg. 5. 159.)

— chloroplatinate, $[(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4]_2\text{PtCl}_6$
As the chloroplatinite. (Jörgensen.)

— chloroplatinite, $[(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4]_2\text{PtCl}_4$.
Somewhat sol. in H_2O , and not insol. in
50% alcohol. (Jörgensen.)

— chromate, $[(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4]_2\text{Cr}_2\text{O}_7$.
Ppt. (Jörgensen.)

— nitrate, $\text{Co}(\text{NO}_2)_2(\text{NH}_3)_4\text{NO}_3$.

Sol. in about 33 pts. cold H_2O ; insol. in
 HNO_3 . (Jörgensen.)

$\text{Co}(\text{NO}_2)_2(\text{NH}_3)_4\text{NO}_3$, HNO_3 . Decomp.
by H_2O or alcohol. (Jörgensen.)

— cobaltic nitrite, $3(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4$,
 $\text{Co}_2(\text{NO}_2)_6 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Jörgensen, Z. anorg. 5.
179.)

— diamine cobaltic nitrite,

$(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4$,

$(\text{NO}_2)_2(\text{NH}_3)_2\text{Co}(\text{NO}_2)_2$.

Very sl. sol. in H_2O . (Jörgensen.)

— sulphate, $[(\text{NO}_2)_2\text{Co}(\text{NH}_3)_4]_2\text{SO}_4$.

Sl. sol. in H_2O , more easily in $\text{HC}_2\text{H}_3\text{O}_2 +$
 Aq . (Jörgensen.)

Fluoborhydric acid, HBF_4 .

Decomp. by H_2O very rapidly. (Landolph,
C. R. 86. 603.)

Aluminum fluoboride, $2\text{AlF}_3, 3\text{BF}_3$.

Sol. in H_2O only when acidulated; sol. in
acids. (Berzelius.)

Ammonium fluoboride, NH_4BF_4 .

Easily sol. in H_2O . Sol. in 4 pts. H_2O at
 16° , and 1.02–1.05 pts. boiling H_2O . (Stolba,
Chem. techn. Cent. Anz. 7. 459.) Sl. sol. in
alcohol.

Barium fluoboride, $\text{Ba}(\text{BF}_4)_2 + 2\text{H}_2\text{O}$.

Deliquescent; easily sol. in H_2O ; decomp.
by alcohol. (Berzelius.)

Cæsium fluoboride, CsBF_4 .

100 pts. H_2O dissolve 0.92 pt. CsBF_4 at 20° ,
and 0.04 pt. at 100° . (Godeffroy, B. 9. 1367.)

0.02 pts. are sol. in 100 pts. H_2O at 20° .
(Erdmann, Arch. Pharm. 1894, 232. 21.)

Calcium fluoboride, $\text{Ca}(\text{BF}_4)_2$.

Decomp. by H_2O , with formation of a sol.
acid salt and an insol. basic salt. (Berzelius.)

Cupric fluoboride, $\text{Cu}(\text{BF}_4)_2$.

Deliquescent, and very sol. in H_2O . (Berzelius.)

Lead fluoboride, $\text{Pb}(\text{BF}_4)_2$.

Sol. in H_2O . Decomp. by boiling with H_2O or alcohol into an acid soluble, and a basic insoluble salt. (Berzelius.)

Lithium fluoboride, LiBF_4 .

Hygroscopic. Easily sol. in H_2O . (Berzelius.)

Magnesium fluoboride.

Easily sol. in H_2O . (Berzelius.)

Potassium fluoboride, KBF_4 .

Sol. in 223 pts. H_2O at 20° . (Stolba.)

Sol. in 70.4 pts. cold H_2O . (Berzelius.)

Sol. in 15.94 pts. H_2O at 100° . (Stolba.)

1.43 pts. are sol. in 100 pts. H_2O at 20° . (Erdmann, Arch. Pharm. 1894, 232. 21.)

Not more sol. in $\text{NH}_4\text{OH} + \text{Aq}$ than in H_2O ; sol. in hot KOH , NaOH , or $\text{M}_2\text{CO}_3 + \text{Aq}$. (Berzelius.) More sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Rose, Pogg. 80. 276.) Insol. in 20% $\text{KC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Stromeyer.) Insol. in cold, sl. sol. in boiling alcohol.

Rubidium fluoboride, RbBF_4 .

100 pts. H_2O dissolve 0.55 pt. at 20° , and 1.0 pt. at 100° . (Godeffroy, B. 9. 1337.)

0.55 pts. are sol. in 100 pts. H_2O at 20° . (Erdmann, Arch. Pharm. 1894, 232. 21.)

Sodium fluoboride, NaBF_4 .

Easily sol. in H_2O . Very sl. sol. in alcohol. (Berzelius.)

Yttrium fluoboride.

Sol. in H_2O with excess of acid. (Berzelius.)

Zinc fluoboride, $\text{Zn}(\text{BF}_4)_2$.

Deliquescent. Sol. in H_2O . (Berzelius.)

Fluoboric acid, HBF_4 .

See Fluoborhydric acid.

$\text{H}_4\text{B}_2\text{O}_7$, 3HF and $\text{H}_4\text{B}_2\text{O}_6$, 2HF (?). Fume on air, and are decomp. with H_2O . (Lan-dolph, B. 12. 1583.)

HBO_2 , 3HF . Decomp. by H_2O . (Berzelius, Pogg. 59. 644.)

Is either a mixture, or a solution of HBO_2 in HF , and is decomp. by distillation, and the salts are decomp. by recrystallisation. (Bas-aw, C. R. 78. 1698.)

Potassium fluoborate, $\text{K}_2\text{B}_2\text{O}_5\text{F}_2$ (?).

Sl. deliquescent. Scarcely sol. in boiling alcohol. (Schiff, A. Suppl. 5. 175.)

See Boron trioxide potassium fluoride, B_2O_3 , 2KF .

Fluochromic acid.**Ammonium fluochromate, $\text{NH}_4\text{CrO}_4\text{F}$**

Sol. in H_2O . (Varenne, C. R. 91.

Potassium fluochromate, KCrO_4F .

Efflorescent. Sol. in H_2O , with decomp. (Streng, A. 129. 225.)

Fluocolumbic acid.

See also Fluoxycolumbic acid.

Ammonium fluocolumbate fluoxycol

$(\text{NH}_4)_2\text{CbF}_7$, 2CbOF_2 , NH_4F .

Cadmium fluocolumbate, $\text{Cd}_2\text{H}_2\text{Cl}$

$28\text{H}_2\text{O}$.

Insol. in, and decomp. by H_2O . (

Cobalt fluocolumbate, $\text{Co}_2\text{H}_2\text{Cb}_2\text{F}_{11}$

Insol. in, and decomp. by H_2O . (

Copper fluocolumbate, Cu_2HCbF_7

Insol. in, and decomp. by H_2O .

Ferrous fluocolumbate, $\text{Fe}_2\text{H}_2\text{Cb}_2\text{F}_{11}$

As above.

Manganous fluocolumbate, Mn_2H_2

$28\text{H}_2\text{O}$.

Mercuric fluocolumbate, $\text{Hg}_2\text{CbF}_{11}$

As above.

Nickel fluocolumbate, $\text{Ni}_2\text{H}_2\text{Cb}_2\text{F}_{11}$

As above.

Potassium fluocolumbate, K_2CbF_7

Decomp. by solution in H_2O . (I A. ch. (4) 8. 34.)

Rubidium fluocolumbate, Rb_2CbF_7

Sol. in H_2O and $\text{HF} + \text{Aq}$. Insol. in alcohol. (Pennington, J. Am. Chem. Soc. 1896, 18. 58.)

Zinc fluocolumbate, $\text{Zn}_2\text{H}_2\text{Cb}_2\text{F}_{11} + 2$

Insol. in cold H_2O ; decomp. by H_2O . (Santesson, Bull. Soc. (2) 24. 52.)

Fluodithionic acid.**Cæsium monofluodithionate,**

$\text{S}_2\text{O}_4(\text{OH})\text{FCs}_2 + \text{H}_2\text{O}$.

Easily sol. in H_2O with decomp.

Sol. in HF ; very unstable. (Weinberg, anorg. 1899, 21. 66.)

Potassium difluodithionate, $\text{S}_2\text{O}_4\text{F}_2\text{K}$

Easily sol. in H_2O with decomp.

Sol. in HF ; very unstable. (Weinberg, anorg. 1899, 21. 66.)

1 difluodithionate, $S_2O_3F_2Rb_2 + 3H_2O$.
 sol. in H_2O with decomp.
 HF; very unstable. (Weinland.)

manic acid, H_2GeF_6 .

1 only in solution. (Winkler, J. pr. 77.)

m fluogermanate, K_2GeF_6 .

173.98 pts. H_2O at 18° . (Winkler.)
 184.61 pts. H_2O at 18° . (Krüss and 3. 20. 1696.)

34.07 pts. H_2O at 100° . (Winkler.)
 38.76 pts. H_2O at 100° . (Krüss and

in alcohol.

dic acid.

um difluoiodate, $NH_4IO_2F_2$.

5 salt.

n 40% HF + Aq. (Weinland, Z. 399, 20. 30.)

1 H_2O . Easily decomp. (Weinland, 30. 868.)

difluoiodate, $CsIO_2F_2$.

land, Z. anorg. 1899, 20. 36.)

hydrogen difluoiodate, $O_2F_2, HIO_2F_2 + 2H_2O$.

acent. Sol. in H_2O with decomp.
nd, Z. anorg. 1899, 22. 257.)

m difluoiodate, KIO_2F_2 .

in H_2O . Decomp. in moist air.
nd, B. 1897, 30. 867.

np. in air. Sol. in H_2O with decomp.
out decomp. in 40% HF + Aq.
nd, Z. anorg. 1899, 20. 31.

m difluoiodate, $RbIO_2F_2$.

ibles K salt. Sol. in HF + Aq.
nd, Z. anorg. 1899, 20. 35.)

m hydrogen difluoiodate, $IO_2F_2, HIO_2F_2 + 2H_2O$.

n 40-60% HF + Aq. (Weinland, Z. 899, 22. 260.)

difluoiodate, $NaIO_2F_2$.

np. by H_2O . (Weinland, B. 1897, 30.

1 HF. (Weinland, Z. anorg. 1899, 20.

inorganic acid, H_2MnF_6 .

np. by H_2O . Sol. in alcohol and ether
ice of H_2O . (Nicklès, C. R. 65. 107.)

Ammonium fluomanganate, $(NH_4)_2MnF_6$.

More sol. than the K salt. (Nicklès, C. R. 65. 107.)

True composition is $(NH_4)_4Mn_2F_{10} = 4NH_4F, Mn_2F_6$. (Christensen, J. pr. (2) 34. 41.)

Cobalt fluomanganate, $2CoF_2, Mn_2F_6 + 8H_2O$.

Sol. in H_2O . (Christensen.)

Nickel fluomanganate, $2NiF_2, Mn_2F_6 + 8H_2O$.

Sol. in H_2O . (Christensen.)

Potassium fluomanganate, K_2MnF_6 .

Difficultly sol. in H_2O . Decomp. by much H_2O . (Nicklès, C. R. 65. 107.)

Composition is $K_4Mn_2F_{10} = 4KF, Mn_2F_6$.
Also with $2H_2O$. (Christensen, J. pr. (2) 34. 41.)

Decomp. by H_2O . Sol. in HCl, H_2SO_4 and HNO_3 with decomp. Can be recryst. from 40% HF + Aq. Insol. in acetic acid. (Weinland and Lauenstein, Z. anorg. 1899, 20. 41.)

Rubidium fluomanganate, $Rb_2MnF_6 + 2H_2O$.

As the K salt. (Weinland and Lauenstein, Z. anorg. 1899, 20. 44.)

Silver fluomanganate, $Ag_2Mn_2F_8 + 14H_2O$.

(Christensen, J. pr. (2) 34. 41.)

Sodium fluomanganate, $4NaF, Mn_2F_6$.

Decomp. by much H_2O . (Christensen.)

Zinc fluomanganate, $2ZnF_2, Mn_2F_6 + 8H_2O$.

Sol. in H_2O . (Christensen.)

Fluomolybdic acid.

See Fluoxyhypomolybdic, and Fluoxymolybdic acids.

Fluopalladous acid.

Potassium fluopalladite,

Sl. sol. in H_2O .

Sodium fluopalladite.

Sl. sol. in H_2O . (Berzelius.)

Fluoperboric acid.

Ammonium fluoperborate,

$NH_4OOb(F)OOb(F)OONH_4$.

Ppt. Insol. in ether. (Petrenko, C. C. 1902, I. 1191.)

Potassium fluoperborate, $K_4B_4F_{10} + H_2O$.

Dry salt is rather stable.

Easily sol. in H_2O . Aqueous solution decomp. rapidly when warmed; at ordinary

temp. the decomp. proceeds slowly. Insol. in alcohol. (Melikoff, B. 1899, 32. 3350.)
 $\text{KOOB(F)OOB(F)OK} + 1\frac{1}{2}\text{H}_2\text{O}$. Ppt. Insol. in ether. (Petrenko, C. C. 1902, I. 1191; J. Russ, phys. chem. Soc. 34. 37.)

Fluoperuranic acid.

Potassium fluoperuranate, $\text{K}_4\text{U}_4\text{F}_6\text{O}_{18} + 4\text{H}_2\text{O} = 3\text{UO}_4\text{KF}, \text{UO}_2\text{F}_2, \text{KF} + 4\text{H}_2\text{O}$.
Ppt. (Lordkipanidse, C. C. 1900, II. 525.)
Sodium fluoperuranate, $\text{UO}_4\text{NaF} + 5\text{H}_2\text{O}$.
Ppt. (Lordkipanidse, C. C. 1900, II. 525.)

Fluophosphamide, $\text{PF}_3(\text{NH}_3)_2$.
Sol. in H_2O . (Poulenc, A. ch. (6) 24. 566.)

Fluophosphoric acid.

Monocæsium monofluophosphate, $\text{P(OH)}_2(\text{OCs})\text{F}$.
Like the K salt. (Weinland, Z. anorg. 1899, 21. 48.)
Monopotassium monofluophosphate, $\text{P(OH)}_2(\text{OK})\text{F}$.
Sol. in 40% HF + Aq; decomp. in the air. (Weinland, Z. anorg. 1899, 21. 44.)

Potassium monofluophosphate, $\text{KHF.PO}_3 + \text{H}_2\text{O}$.
Decomp. by H_2O ; unstable. (Weinland, B. 1898, 31. 124-125.)

Monorubidium monofluophosphate, $\text{P(OH)}_2(\text{ORb})\text{F}$.
Sol. in 40% HF + Aq. (Weinland, Z. anorg. 1899, 21. 47.)

Rubidium monofluophosphate, $\text{RbHFPO}_3 + \text{H}_2\text{O}$.
Decomp. by H_2O . (Weinland, B. 1898, 31. 124.)

Fluoplatinic acid.

Ammonium fluoplatinate.
Decomp. by H_2O to a sol. acid, and an insol. basic salt. Insol. in alcohol. (Berzelius.)
Potassium fluoplatinate.
Deliquescent. Insol. in alcohol. Decomp. by H_2O . (Berzelius.)
Sodium fluoplatinate.
Decomp. by H_2O . (Berzelius.)

Fluor- and Fluoro-.
See Fluo-.

Fluorhydric (Hydrofluoric) acid, HF or H_2F_2 .

Attracts H_2O from air with great avidity. Very sol. in H_2O with evolution of much heat. Sat. solution has sp. gr. 1.25. (H. Davy.)
On boiling the aqueous solution an acid of constant composition is obtained, which boils at 120° , has sp. gr. 1.15, and contains 35.37% HF (Bineau, A. ch. (3) 7. 257.) The residual acid after boiling contains 36 to 38% HF, and by standing over CaO gives off HF until an acid containing 32.5 to 32.7% HF is formed. Weaker acids increase their strength to 32.3 to 32.4% HF, while an acid containing 32.5% HF remains unchanged. (Roscoe, A. 118. 218.)

Does not attack gutta-percha. Sol. in H_2SO_4 .

Sp. gr. of HF + Aq at 15° .

Sp. gr.	% HF	Sp. gr.	% HF	Sp. gr.	% HF
1.01	2.90	1.10	29.00	1.19	55.10
1.02	5.80	1.11	31.90	1.20	58.00
1.03	8.70	1.12	34.80	1.21	60.90
1.04	11.60	1.13	37.70	1.22	63.80
1.05	14.50	1.14	40.60	1.23	66.70
1.06	17.40	1.15	43.50	1.24	69.60
1.07	20.30	1.16	46.40	1.25	72.50
1.08	23.20	1.17	49.30
1.09	26.10	1.18	52.20

(Hart, J. Anal. Ch. 3. 372.)

Sp. gr. of HF + Aq at ord. temp.

Deg. Baumé	Sp. gr.	% HF
1	1.0069	2.32
2	1.0139	4.04
3	1.0211	5.76
4	1.0283	7.48
5	1.0356	9.20
6	1.0431	10.92
7	1.0506	12.48
8	1.0583	14.04
9	1.0661	15.59
10	1.074	17.15
11	1.082	18.86
12	1.0901	21.64
13	1.0983	24.42
14	1.1067	27.20
15	1.1152	29.98
16	1.1239	32.78
17	1.1326	35.15
18	1.1415	37.53
19	1.1506	39.91
20	1.1598	42.29
21	1.1691	44.67
22	1.1786	47.04
23	1.1883	49.42
24	1.1981	51.57
25	1.2080	53.72
26	1.2182	55.87
27	1.2285	58.02

of HF + Aq at ord. temp.—Continued.

Sp. gr.	% HF
1.2390	60.17
1.2497	62.32
1.2605	64.47
1.2716	66.61
1.2828	68.76
1.2943	70.91
1.3059	73.06
1.3177	75.21
1.3298	77.36
1.3421	79.51
1.3546	81.66
1.3674	83.81
1.3804	85.96
1.3937	88.10
1.4072	90.24
1.4211	92.39
1.4350	94.54
1.4493	96.69

Eckelt, Ch. Z. 1898, 22. 225.)

Sp. gr. of HF + Aq at 0°.

Sp. gr.	% HF	Sp. gr.
1.005	71.73	1.262
1.009	72.21	1.260
1.012	78.05	1.260
1.017	84.27	1.235
1.035	87.72	1.212
1.065	88.11	1.210
1.097	88.82	1.207
1.110	89.02	1.202
1.120	89.15	1.200
1.130	89.82	1.190
1.145	90.20	1.185
1.155	90.64	1.175
1.155	91.04	1.165
1.157	92.09	1.152
1.182	92.81	1.135
1.187	92.91	1.130
1.200	94.26	1.095
1.217	95.84	1.065
1.220	97.50	1.035
1.230	98.22	1.022
1.245	100.05	1.0005
1.255

Roy. Soc. Proc. 1909, 83. A. 144.)

Sp. gr. of HF + Aq at 18°.

% HF	Sp. gr.
0.484	1.003
1.504	1.005
2.48	1.009
4.80	1.017
7.75	1.028
15.85	1.058
24.47	1.087
29.83	1.103

(Hill.)

Aq. solution of sp. gr. 1.138 at 18° contains 43.2% HF and has a constant bpt. of 111° at 750 mm. (Deussen, Z. anorg. 1906, 49. 297.)

The strongest acid that can be obtained by distillation contains 48.17% HF and boils at 125–125.5°. (Gore.)

Fluorides.

The alkali fluorides, also AgF and SnF₂, are sol. in H₂O; the fluorides of Fe, Sr, and Cd are sl. sol.; the others are insol. in H₂O. Most fluorides are sol. in acids, especially HF + Aq.

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 822.)

See under each element.

Fluorine, F₂.

Decomposes H₂O and all organic solvent with great violence. (Moissan, C. R. 103. 202 and 256.)

Liquified at –185° to a yellowish liquid which does not dissolve glass nor ignite cooled Si, B, C, S, P, or Fe. (Moissan, C. R. 1897, 124. 1202–1204.)

Fluomolybdic acid.

Ammonium fluomolybdate, (NH₄)MoF₄ + H₂O.

Somewhat more sol. in H₂O than the K salt. Hydrolysed by H₂O. (Rosenheim, Z. anorg. 1905, 46. 321.)

(NH₄)₂Mo₂F₇ + 2H₂O. (Rosenheim.)

Potassium fluomolybdate, KMoF₄ + H₂O.

Nearly insol. in H₂O. (Rosenheim.)

Fluoselenic acid.

Ammonium monofluoselenate, SeO₃(OH)F(NH₄)₂.

Not hygroscopic.

Easily sol. H₂O with decomp.

Sol. in HF. (Weinland, Z. anorg. 1899, 21. 58.)

Tripotassium difluodiselenate, Se₂O₇F₂K₃H + H₂O.

Decomp. in the air; sol. in H₂O with decomp.; sol. in HF. (Weinland.)

Trirubidium difluodiselenate, Se₂O₇F₂Rb₃H + H₂O.

Decomp. in the air; sol. in H₂O with decomp.; sol. in HF. (Weinland, Z. anorg. 1899, 21. 57.)

Fluosilicic acid, H_2SiF_6 .

Sp. gr. of $\text{H}_2\text{SiF}_6 + \text{Aq}$ at 17.5° (H_2O at $17.5^\circ = 1.000$).

% H_2SiF_6	Sp. gr.	% H_2SiF_6	Sp. gr.
2	1.0161	20	1.1748
4	1.0324	22	1.1941
6	1.0491	24	1.2136
8	1.0661	26	1.2335
10	1.0834	28	1.2537
12	1.1011	30	1.2742
14	1.1190	32	1.2951
16	1.1373	34	1.3162
18	1.1559		...

(Stolba, J. pr. 90. 193.)

+2 H_2O . Very deliquescent, and sol. in H_2O . (Kessler, C. R. 90. 1285.) Solution decomp. into HF and SiF_4 on evaporation, when it becomes concentrated.

Fluosilicates.

Most of the fluosilicates are sol. in H_2O , but the alkali salts (especially K) and the Ba salt are only sl. sol. in H_2O .

Aluminum fluosilicate, $\text{Al}_2(\text{SiF}_6)_3$.

Easily sol. in H_2O . After evaporating to dryness, the residue is slowly but completely sol. in H_2O . (Deville, A. ch. (3) 61. 327.)

Insol. in acetone. (Naumann, B. 1904, 37. 4328.)

Aluminum fluosilicate silicate, $\text{Al}_2\text{SiF}_{10}$, $5\text{Al}_2\text{SiO}_4$.

Min. *Topaz*. Insol. in acids.

Ammonium fluosilicate, $(\text{NH}_4)_2\text{SiF}_6$.

Sol. in 538 pts. H_2O at 17.5° to form a solution of 1.0961 sp. gr.; sol. in 1.8 pts. hot H_2O ; sol. in 45.5 pts. alcohol of 31%. (Stolba, C. C. 1877. 418.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

$3\text{NH}_4\text{F} \cdot \text{SiF}_4 = (\text{NH}_4)_2\text{SiF}_6 \cdot \text{NH}_4\text{F}$. Sol. in H_2O . (Marignac, Ann. Min. (5) 15. 221.)

Barium fluosilicate, BaSiF_6 .

Sol. in 3802 pts. cold H_2O . (Fresenius, A. 59. 120.)

Sol. in 3731 pts. H_2O at 17.5° ; in 3315 pts. at 21° ; in 1175 pts. at 100° . (Stolba, J. pr. 90. 22.)

Sol. in 640-733 pts. H_2O containing a little HCl . (Fresenius.)

488 pts. $\text{HCl} + \text{Aq}$ containing 4.25% HCl dissolve 1 pt. at 22° . (Stolba.)

More sol. in $\text{HNO}_3 + \text{Aq}$ than in H_2O . (Fresenius.)

272 pts. $\text{HNO}_3 + \text{Aq}$, containing 8% N_2O_5 , dissolve 1 pt. at 22° . (Stolba.)

1 pt. BaSiF_6 dissolves in 428 pts. sat. $\text{NH}_4\text{Cl} + \text{Aq}$; in 589 pts. sat. $\text{NH}_4\text{Cl} + \text{Aq} + 2$ vols. H_2O . (Mallet, Sill. Am. J. (2) 20. 48.)

1 pt. BaSiF_6 dissolves in 306 pts. sat. $\text{NH}_4\text{Cl} + \text{Aq}$ at 22° ; in 361 pts. 15% solution of NH_4Cl ; in 563 pts. sat. boiling $\text{NaCl} + \text{Aq}$; in 349 pts. 10% solution of NaCl at boiling temp.; in 2185 pts. 10% solution of NaCl at 20° ; in 1140 pts. 5% solution of NaCl at 20° . (Stolba.)

Nearly absolutely insol. in alcohol. (Fresenius.)

Solubility in a mixture of H_2O , alcohol (96%), $\text{HCl} + \text{Aq}$ (20%), $\text{H}_2\text{SiF}_6 + \text{Aq}$ (3.7%). 1 pt. BaSiF_6 is sol. in pts. of solutions of given composition.

H_2O	Alcohol	$\text{HCl} + \text{Aq}$	$\text{H}_2\text{SiF}_6 + \text{Aq}$	BaSiF_6
■	50	0	0	37.219
74.1	25	0.9	■	5.263
70.8	25	4.2	0	2.860
77.95	20	0.9	1.15	39.061
73.0	25	0.9	1.1	70.679
97.09	0	1.25	1.66	3.247
75.0	25	■	0	16.914

(Fresenius, Z. anal. 29. 143.)

Cadmium fluosilicate, $\text{CdSiF}_6 + 6\text{H}_2\text{O}$.

Extremely sol. in H_2O . Easily sol. in 50% alcohol. (Engelskirchen, Dissert. 1903.)

Cesium fluosilicate, Cs_2SiF_6 .

Sol. in 166 pts. H_2O at 17° , and much less hot H_2O . Insol. in alcohol. (Preis, J. pr. 103. 410.)

Calcium fluosilicate, $\text{CaSiF}_6 + 2\text{H}_2\text{O}$.

Sl. sol. in, and partly decomp. by H_2O . Sol. in HF and $\text{HCl} + \text{Aq}$. Sol. in fluosilicic acid without decomp. Easily sol. in 60% alcohol. (Fleischer.)

Cerium fluosilicate.

Very difficulty sol. in H_2O , acetic, or fluosilicic acids. Insol. in alcohol. (Stolba, C. C. 1874. 130.)

Chromium fluosilicate.

Deliquescent. (Berzelius.)

Efflorescent. Sol. in H_2O . (Berlin.)

Cobaltous fluosilicate, $\text{CoSiF}_6 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Berzelius.)

Cuprous fluosilicate, Cu_2SiF_6 .

Insol. in H_2O . (Berzelius, Pogg. 1. 199.)

Cupric fluosilicate, $\text{CuSiF}_6 + 6\text{H}_2\text{O}$.

Deliquescent in moist, efflorescent in dry air.

Sol. in 0.428 pt. H_2O at 17° . Sp. gr. of solution sat. at $17^\circ = 1.6241$.

Sol. in 17.5 pts. alcohol of 62 vol. % at 20° ; in 150 pts. of 85% at 20° ; in 617 pts. of 90% at 20° . (Stolba, J. pr. 102. 7.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Contains $6\frac{1}{2}$ H₂O. (Stolba.)

+ $5\frac{1}{2}$ H₂O. (Knop and Wolf.)

Capric fluosilicate phosphate, CuSiF₆, Cu₃(PO₄)₂.

Insol. in H₂O, but easily sol. in dil. HCl + Aq. (Thorpe and Rodger, Chem. Soc. 55. 320.)

Glucinum fluosilicate.

Known only in solution.

Iron (ferrous) fluosilicate, FeSiF₆ + 6H₂O.

Easily sol. in H₂O. (Berzelius.)

Iron (ferric) fluosilicate, Fe₂(SiF₆)₃.

Sol. in H₂O. (Berzelius.)

Lead fluosilicate, PbSiF₆ + 2H₂O.

Deliquescent. Easily sol. in H₂O.

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+4H₂O. (Marignac.)

Lithium fluosilicate, Li₂SiF₆ + 2H₂O.

100 pts. H₂O at 17° dissolve 73 pts. crystalline salt. (Marignac.)

100 pts. cold H₂O dissolve 52.6 pts. crystals.

Sol. in dil. alcohol. (Stolba, J. pr. 91. 456.)

100 pts. alcohol of 46 vol. % dissolve about 4 pts., and 100 pts. alcohol of 79 vol. % dissolve about 0.4 pt. crystals. (Stolba, Z. anal. 8. 311.)

Insol. in ether or benzene.

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Magnesium fluosilicate, MgSiF₆ + 6H₂O.

Efflorescent. Sol. in 1534 pts. cold H₂O, forming a solution of 1.235 sp. gr. at 17.5°. Separates out SiO₂ on warming, which nearly all redissolves on cooling. (Stolba, C. C. 1877. 578.)

Magnesium fluosilicate silicate, Mg₂Si₂F₁₂, xMg₂Si₂O₇.

Min. *Humite*; *Chondrodite*. Gelatinises with HCl, or H₂SO₄ + Aq.

Manganous fluosilicate, MnSiF₆ + 6H₂O.

Sol. in H₂O. (Marignac, J. pr. 83. 202.)

100 pts. dissolve in 71.4 pts. H₂O at 17.5°, and sp. gr. of solution = 1.44825. Much more sol. in hot H₂O, and less sol. in alcohol, the stronger the alcohol. (Stolba, C. C. 1883. 292.)

Mercurous fluosilicate, Hg₂SiF₆.

Sl. sol. in H₂O without decomp. (Lemaire, C. C. 1897, I. 1046.)

+2H₂O. Sl. sol. in H₂O. More easily sol. in acidified H₂O, but precipitated by HCl + Aq. (Berzelius.)

Mercuric fluosilicate, basic, HgSiF₆, HgO + 3H₂O.

Decomp. by H₂O, but sol. in weakest acids. (Berzelius, Pogg. 1. 200.)

Mercuric fluosilicate, HgSiF₆ + 6H₂O.

Deliquescent, and easily sol. in H₂O. (Finkener, Pogg. 111. 246.)

Nickel fluosilicate, NiSiF₆ + 6H₂O.

Easily sol. in H₂O. (Marignac, Ann. Min. (5) 15. 262.)

Potassium fluosilicate, K₂SiF₆.

Sol. in 833.1 pts. H₂O at 17.5°, and 104.8 pts. at 100°. (Stolba, J. pr. 103. 396.) Sol. in 3800 pts. cold, and more easily sol. in hot H₂O. (Fresenius.)

More sol. in HCl + Aq than in H₂O.

Sol. in 337 pts. HCl + Aq of 26.5% at 14°; in 307 pts. of 25.7% at 15°; in 340 pts. of 14.1% at 14°; in 303 pts. of 13.6% at 15°; in 327 pts. of 9.6% at 14°; in 313 pts. of 9.2% at 15°; in 376 pts. of 2.7% at 14°; in 319 pts. of 2.4% at 15°; in 409 pts. of 1.8% at 14°. (Stolba, l. c.)

Sol. in 428 pts. sat., and 589 pts. dil. NH₄Cl + Aq. (Mallet.)

Much less sol. in K₂SO₄, KNO₃, or KCl + Aq, but more sol. in NH₄Cl + Aq than in H₂O. (Stolba.)

Sol. in 24,066 pts. K₂SO₄ + Aq containing 9.92% K₂SO₄ at 17°; in 17,858 pts. containing 6% at 18°; in 19,530 pts. containing 5% at 17°; in 10,721 pts. containing 1% at 17°.

Sol. in 125,000 pts. KNO₃ + Aq containing 18.4% KNO₃ at 15°; in 43,478 pts. containing 8.7% at 15°; in 1735 pts. containing 8.8% at 100°; in 35,814 pts. containing 4.3% at 15°; in 10,203 pts. containing 1.00% at 15°.

Sol. in 40,070 pts. KCl + Aq containing 25% KCl at 17°; in 38,352 pts. containing 18.4% at 17°; in 41,254 pts. containing 13.4% at 14°; in 24,032 pts. containing 6.7% at 12°; in 1200 pts. containing 0.65% at 17°; in 1095 pts. containing 0.45% at 18°.

Sol. in 358 pts. NH₄Cl + Aq containing 26.3% NH₄Cl at 17°; in 306 pts. containing 15% at 15°; in 339 pts. containing 10% at 15°; in 436 pts. containing 5% at 15°. (Stolba, J. pr. 103. 306.)

Insol. in liquid CO₂. (Büchner, Z. phys. Ch. 1906, 54. 674.)

Insol. in liquid NH₃. (Gore, Am. ch. J. 1898, 20. 829.)

Completely pptd. from aqueous solution by an equal vol. of alcohol.

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Rubidium fluosilicate, Rb_2SiF_6 .

Sol. in 625 pts. H_2O at 20° , and 73.05–74.5 pts. at 100° . More sol. in acidified water. Insol. in alcohol. (Stolba, J. pr. 101. 1.)

Insol. in H_2O . (Eggeling, Z. anorg. 1905, 46. 175.)

Less sol. in H_2O than K_2SiF_6 . (Gossner, Zeit. Kryst. 1904, 38. 149.)

Silver fluosilicate, $\text{Ag}_2\text{SiF}_6 + 4\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O . (Marignac, Ann. Min. (5) 15. 221.)

Sodium fluosilicate, Na_2SiF_6 .

Much more sol. in H_2O than K_2SiF_6 , especially in hot H_2O . Addition of acid does not increase solubility. (Berzelius.)

Sol. in 153.3 pts. H_2O at 17.5° , and 40.66 pts. at 100° . Easily forms supersaturated solutions. (Stolba, Z. anal. 11. 199.)

Much less sol. in $\text{NaCl} + \text{Aq}$ than in H_2O . (Stolba, J. pr. 1865 (1) 96. 26.)

Precipitated completely from aqueous solution by alcohol. (Rose.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Strontium fluosilicate, $\text{SrSiF}_6 + 2\text{H}_2\text{O}$.

Sol. in cold H_2O , but decomp. somewhat on heating. Sol. in 31.06 pts. H_2O . (Fresenius.)

Easily sol. in acidified H_2O without decomp. Sol. in alcohol.

Solubility in a mixture of H_2O , alcohol (96%), $\text{HCl} + \text{Aq}$ (20%), $\text{H}_2\text{SiF}_6 + \text{Aq}$ (3.7%). 1 pt. SrSiF_6 is sol. in pts. of solutions of given composition.

H_2O	Alcohol	$\text{HCl} + \text{Aq}$	$\text{H}_2\text{SiF}_6 + \text{Aq}$	SrSiF_6
50	50	0	0	15.29
74.1	25	0	0	82.93
70.8	25	4.2	0	50.9
77.95	20	0.9	1.15	55.0
73	25	0.9	1.1	82.97
75	25	0	0	147.4
95.24	0	2.04	2.72	7.3

(Fresenius, Z. anal. 29. 143.)

Thallous fluosilicate, $\text{Tl}_2\text{SiF}_6 + 2\text{H}_2\text{O}$.

Very easily sol. in H_2O . (Kuhlmann.)

Thorium fluosilicate, $\text{Th}(\text{OH})_2\text{SiF}_6$ (?).

(Cleve.)

Tin stannic fluosilicate, $\text{SnF}_4 \cdot \text{SiF}_4$.

Very easily sol. in H_2O . (Berzelius.)

Uranyl fluosilicate.

Very sl. sol. in acids. (Berzelius.)

Sol. in alcohol. (Stolba, Z. anal. 3. 71.)

Vanadium fluosilicate.

Deliquescent. Sol. in H_2O . (Guyard, Bull. Soc. (2) 25. 352.)

Yttrium fluosilicate.

Insol. in pure, sol. in acidified H_2O . (Berzelius.)

Zinc fluosilicate, $\text{ZnSiF}_6 + 6\text{H}_2\text{O}$.

Very easily sol. in H_2O . (Berzelius.)

Zirconium fluosilicate.

Sol. in H_2O . Solution clouds up on boiling. (Berzelius.)

Fluostannic acid.

Ammonium fluostannate, $(\text{NH}_4)_2\text{SnF}_6$.

Sol. in H_2O . (Marignac, Ann. Min. (5) 15. 224.)

$4\text{NH}_4\text{F}, \text{SnF}_4$. Sol. in H_2O . (Marignac.)

Barium fluostannate, BaSnF_6 .

Slowly sol. in H_2O .

$+3\text{H}_2\text{O}$. Sol. in 18 pts. H_2O at 15° . (Marignac, Ann. Min. (5) 15. 246.)

Decomp. by warming with H_2SO_4 with evolution of HF . (Emich, M. 1904, 25. 1912.)

Calcium fluostannate, $\text{CaSnF}_6 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac, Ann. Min. (5) 15. 250.)

Cadmium fluostannate, $\text{CdSnF}_6 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac.)

Cobaltous fluostannate, $\text{CoSnF}_6 + 6\text{H}_2\text{O}$.

(Gossner, Zeit. Kryst. 1907, 42. 482.)

Cupric fluostannate, $\text{CuSnF}_6 + 4\text{H}_2\text{O}$.

Not deliquescent. (Marignac, Ann. Min. (5) 15. 291.)

Lithium fluostannate, $\text{Li}_2\text{SnF}_6 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac, Ann. Min. (5) 15. 242.)

Magnesium fluostannate, $\text{MgSnF}_6 + 6\text{H}_2\text{O}$.

Not deliquescent. Sol. in H_2O . (Marignac, Ann. Min. (5) 15. 256.)

Manganous fluostannate, $\text{MnSnF}_6 + 6\text{H}_2\text{O}$.

Slowly efflorescent. (Marignac.)

Nickel fluostannate, $\text{NiSnF}_6 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac, Ann. Min. (5) 15. 262.)

Potassium fluostannate, $K_2SnF_6 + H_2O$.

Two modifications—(a) *Thin plates*. Sol. in 2.3 pts. H_2O at 100° , and in 15–16 pts. at 18° . (Marignac.)

(b) *Octahedra*. Sol. in 3 pts. H_2O at 100° , and 27 pts. at 18° . (Marignac.)

Sol. in hot H_2O . Can be crvst. from hot H_2O . With conc. H_2SO_4 , HF is evolved. (Emich, M. 1904, 25. 911.)

Potassium hydrogen fluostannate, $3KF, HF, SnF_4$.

Sol. in H_2O . (Marignac.)

Silver fluostannate, $Ag_2SnF_6 + 4H_2O$.

Sl. deliquescent. Easily sol. in H_2O . (Marignac.)

Sodium fluostannate, Na_4SnF_6 .

Sol. in 18–19 pts. H_2O at 20° . (Marignac.)

Strontium fluostannate, $SrSnF_6 + 2H_2O$.

Sol. in 5.5 pts. H_2O at 18° . (Marignac.)

Zinc fluostannate, $ZnSnF_6 + 6H_2O$.

Sol. in H_2O . (Marignac.)

Fluosulphonic acid, HSO_3F .

See Sulphuryl hydroxyl fluoride.

Ammonium fluosulphonate, FSO_3NH_4 .

Easily sol. in H_2O from which it can be crvst.

Sol. in ethyl alcohol, more sol. in methyl alcohol. Can be crvst. from abs. alcohol. (Traube, B. 1913, 46. 2528.)

Sodium fluosulphonate, FSO_3Na .

Hydroscopic.

Sol. in alcohol and acetone. (Traube.)

Fluosulphuric acid.

Tricassium difluodisulphate, $S_2O_7F_2Cs_3H + H_2O$.

As the K salt. (Weinland, Z. anorg. 1899, 21. 53.)

Tripotassium difluodisulphate,

$S_2O_7F_2K_3H + H_2O$.

Sol. in HF ; quite stable in air; sol. in H_2O with decomp. (Weinland, Z. anorg. 1899, 21. 51.)

Trirubidium difluodisulphate, $S_2O_7R_3Tb_3H + H_2O$.

Sol. in HF . (Weinland, Z. anorg. 1899, 21. 53.)

Fluotantallic acid.

Ammonium fluotantalate, $(NH_4)_2TaF_7$.

Very sol. in H_2O . (Marignac, A. ch. (4) 9. 272.)

$(NH_4)_2TaF_7$. (Balke, J. Am. Chem. Soc. 1905, 27. 1151.)

Cæsium fluotantalate, $CsTaF_6$.

Can be recryst. from $HF + Aq$. (Balke.)

Cs_2TaF_7 . Can not be recryst. from H_2O as it tends to go into CsF, TaF_5 . (Balke, J. Am. Chem. Soc. 1905, 27. 1151.)

$15CsF, TaF_5$. Sl. sol. in H_2O . (Pennington, J. Am. Chem. Soc. 1896, 18. 59.)

Calcium fluotantalate.

Difficulty sol. in H_2O (Berzelius.)

Cupric fluotantalate, $CuTaF_7 + 4H_2O$.

Deliquescent. Easily sol. in H_2O . (Marignac, A. ch. (4) 9. 294.)

Lead fluotantalate.

Difficulty sol. in H_2O . (Berzelius.)

Lithium fluotantalate, $LiF, TaF_5 + 2H_2O$.

Can be recryst. from conc. HF . (Balke, J. Am. Chem. Soc. 1905, 27. 1143.)

Potassium fluotantalate, K_2TaF_7 .

Sl. sol. in cold, much more easily in hot H_2O . Decomposes, with formation of a white precipitate on boiling. (Berzelius.)

Much more sol. in $HF + Aq$. 1 pt. of the salt is sol. in 200 pts. H_2O containing a trace of HF , and in 150–160 pts. of H_2O containing a little more HF . (Marignac, A. ch. (4) 9. 267.)

Potassium hydrogen fluotantalate, $KF, HF, TaF_5 (?)$.

Sol. in H_2O . (Berzelius.)

Rubidium fluotantalate, Rb_2TaF_7 .

Sol. in $HF + Aq$. (Pennington, J. Am. Ch. Soc. 1896, 18. 58.)

$3RbF, 2TaF_5$. (Balke, J. Am. Chem. Soc. 1905, 27. 1151.)

Sodium fluotantalate, $3NaF, TaF_5$.

Easily sol. in H_2O .

$Na_2TaF_7 + H_2O$. Sol. in H_2O . (Marignac.)

Thallous fluotantalate, Tl_2TaF_7 .

Sol. in H_2O . On boiling the aqueous solution tantalic acid separates.

Decomp. by conc. H_2SO_4 . Difficultly sol. in cold, easily sol. in hot HF . (Ephraim. B. 1909, 42. 4461.)

Zinc fluotantalate, $ZnTaF_7 + 7H_2O$.

Very deliquescent. Sol. in H_2O . (Marignac, A. ch. (4) 9. 249.)

Fluotelluric acid.

Ammonium fluotellurate, $NH_4TeF_6 + H_2O$.

Decomp. by H_2O . (Högbom, Bull. Soc. (2) 35. 60.)

Barium fluotellurate, $\text{Ba}(\text{TeF}_6)_2 + \text{H}_2\text{O}$.

As above.

Potassium fluotellurate, KTeF_6 .

As above.

$\text{TeO}_2\text{F}_2\text{K}_2 + 3\text{H}_2\text{O}$. Stable in dry air; only sl. sol. in H_2O with decomp.; sol. in HF . (Weinland, Z. anorg. 1899, 21. 61.)

Rubidium difluotellurate, $\text{TeO}_2\text{F}_2\text{Rb}_2 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O with decomp. Sol. in HF . (Weinland, Z. anorg. 1899, 21. 62.)

Fluotitanic acid.

Known only in solution as titanium hydrogen fluoride.

Ammonium fluotitanate, $(\text{NH}_4)_2\text{TiF}_6$.

Sol. in H_2O . (Marignac.)

$3\text{NH}_4\text{F}$, TiF_4 . Sol. in H_2O . (Marignac.)

Ammonium fluosequititanate, $6\text{NH}_4\text{F}$, Ti_2F_6 .

Easily sol. in H_2O . Sl. sol. in $\text{NH}_4\text{F} + \text{Aq}$. (Petersen, J. pr. (2) 40. 54.)

Insol. in $\text{NH}_4\text{F} + \text{Aq}$. (Piccini, C. R. 97. 1064.)

$4\text{NH}_4\text{F}$, Ti_2F_6 . Properties as the corresponding K salt. (Piccini, B. 18. 257 R.)

Barium fluotitanate, BaTiF_6 .

Very sl. sol. in H_2O . More easily sol. in dil. HNO_3 or HCl . (Engelskirchen, Dissert. 1903.)

$+ \frac{1}{2}\text{H}_2\text{O}$. (Emich, M. 1904, 25. 907.)

Cadmium fluotitanate, $\text{CdTiF}_6 + 6\text{H}_2\text{O}$.

Extremely sol. in H_2O . Easily sol. in 50% alcohol. (Engelskirchen, Dissert. 1903.)

Cæsium fluotitanate, Cs_2TiF_6 .

More sol. in hot than cold H_2O and much more sol. than the Rb comp. (Engelskirchen, Dissert. 1903.)

4CsF , TiF_4 . More sol. in H_2O than Cs_2TaF_7 and is not decomp. by pure H_2O . (Pennington, J. Am. Chem. Soc. 1896, 18. 60.)

Calcium fluotitanate, $\text{CaTiF}_6 + 3\text{H}_2\text{O}$.

Decomp. by pure H_2O . Sol. without decomp. in acidified H_2O . (Berzelius.)

Separates a precipitate with cold H_2O , which dissolves on heating. (Marignac, Ann. Min. (5) 15. 250.)

Cupric fluotitanate, $\text{CuTiF}_6 + 4\text{H}_2\text{O}$.

Sol. in pure H_2O with partial decomp.; easily and completely sol. in acidified H_2O . (Berzelius.)

Cupric fluotitanate ammonium fluoride, CuTiF_6 , $\text{NH}_4\text{F} + 4\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O . (Marignac, Ann. Min. (5) 15. 267.)

Cupric fluotitanate potassium fluoride, CuTiF_6 , $\text{KF} + 4\text{H}_2\text{O}$.

As the above salt. (Marignac.)

Ferrous fluotitanate, $\text{FeTiF}_6 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Weber, Pogg. 120. 2)

Ferric fluotitanate.

Decomp. by H_2O . (Berzelius.)

Lead fluotitanate.

Easily sol. in H_2O . (Berzelius.)

Lithium fluotitanate, $\text{Li}_2\text{TiF}_6 + 2\text{H}_2\text{O}$.

Very sol. in H_2O . (Engelskirchen, Dissert. 1903.)

Magnesium fluotitanate, $\text{MgTiF}_6 + 6\text{H}_2\text{O}$.

Easily sol. in cold H_2O . (Marignac, Min. (5) 15. 257.)

Nickel fluotitanate, $\text{NiTiF}_6 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Weber, Pogg. 282.)

Potassium fluotitanate, K_2TiF_6 .

Difficultly sol. in cold, much more eas hot H_2O .

100 pts. H_2O dissolve at:

0° 3° 6° 10° 14° 20°

0.556 0.667 0.775 0.909 1.042 1.28 pts. K, (Marignac, A. ch. (4) 8. 65.)

Sol. in 78.6 pts. H_2O at 21°. Sol. in : (Piccini, Gazz. ch. it. 1886, 16. 104.)

Sol. in 78 pts. H_2O at 20°; 9.4 pts. at By addition of small amount of HF solubility is increased. (Weiss and K Z. anorg. 1910, 65. 354.)

Sol. in HF . (Marchetti, Z. anorg. 10. 66.)

$+ \text{H}_2\text{O}$. Much less sol. in H_2O in pre of KBr or KI . (Hall, J. Am. Chem. 1904, 26. 1246.)

Sol. in H_2O or HF with decomp. (Marchetti, Z. anorg. 1895, 10. 66.)

Potassium fluosesquititanate, 4KF , Ti_2F_7 .

Scarcely sol. in H_2O ; sol. in dil. : (Piccini, B. 18. 257 R.)

Rubidium fluotitanate, Rb_2TiF_6 .

Very sl. sol. in cold, somewhat more s hot H_2O . (Engelskirchen, Dissert. 1903.)

Silver fluotitanate.

Very deliquescent. (Marignac.)

Sodium fluotitanate, Na_2TiF_6 .

Much more sol. in H_2O than the s sponding potassium salt. (Marignac, Min. (5) 15. 238.)

hydrogen fluotitanate, Na_2TiF_6 , TiF_3 .

H_2O . (Marignac.)

strontium fluotitanate, $\text{SrTiF}_6 + 2\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . Solution clouds up on heating. (Marignac.)

zinc fluotitanate, $\text{ZnTiF}_6 + 6\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Marignac, A. ch. (3) 60.)

Fluotitanic acid.

ammonium fluovanadate, $3\text{NH}_4\text{F}$, VF_3 .

Sl. sol. in H_2O . More easily sol. in acids. Nearly insol. in alcohol or MF. (Petersen, J. pr. (2) 40. 52.)

F_3 , $\text{VF}_3 + \text{H}_2\text{O}$. Easily sol. in H_2O . Sl. sol. in alcohol. (Petersen.)

F_3 , $\text{VF}_3 + 2\text{H}_2\text{O}$. As above. (Petersen.)

cadmium fluovanadate, CdF_2 , $\text{VF}_3 + 7\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Piccini and Giorgis, Ann. Chem. (3) 22, 1. 89.)

cobalt fluovanadate, CoF_2 , $\text{VF}_3 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O without decomp. (Petersen, J. pr. (2) 40. 51.)

nickel fluovanadate, NiF_2 , $\text{VF}_3 + 2\text{H}_2\text{O}$.

Co salt. (Petersen.)

potassium fluovanadate, 2KF , $\text{VF}_3 + \text{H}_2\text{O}$.

Sl. sol. in H_2O ; easily sol. in acids. Insol. in Aq. (Petersen, J. pr. (2) 40. 51.)

ammonium fluovanadate fluoxyvanadate, F_3 , VF_3 , VOF_2 .

Sl. sol. in H_2O , and still more easily in Aq. Sl. sol. in $\text{KF} + \text{Aq}$. (Petersen, J. pr. (2) 40. 274.)

sodium fluovanadate, 5NaF , $2\text{VF}_3 + \text{H}_2\text{O}$.

potassium salt. (Petersen.)

thallium fluovanadate, TlF , $\text{VF}_3 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O .

Decomp. in conc. H_2SO_4 , dil. H_2SO_4 , or cold dil. HCl .

Sl. sol. in $\text{NaOH} + \text{Aq}$. (Ephraim, B. 1909, 42. 4461.)

$\text{VF}_3 + \text{H}_2\text{O}$. Easily sol. in H_2O .

Sl. sol. in conc. H_2SO_4 , dil. HNO_3 , or cold dil. H_2SO_4 .

Sl. sol. in cold or hot $\text{NaOH} + \text{Aq}$.

(Ephraim, B. 1909, 42. 4461.)

zinc fluovanadate, ZnF_2 , $\text{VF}_3 + 7\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . Decomp. on heating. (Piccini and Giorgis, Ann. Chem. (3) 22, 1. 89.)

Fluoxycolumbic acid.

Ammonium fluoxycolumbate, $3\text{NH}_4\text{F}$, CbOF_3 .

Cubic salt. Sol. in H_2O . (Marignac, A. ch. (4) 8. 38.)

$2\text{NH}_4\text{F}$, CbOF_3 . Lamellar salt. Much more sol. in H_2O than 2KF , CbOF_3 . (M.)

$5\text{NH}_4\text{F}$, $3\text{CbOF}_3 + \text{H}_2\text{O}$. Hexagonal salt. (M.)

NH_4F , CbOF_3 . Rectangular salt. (M.)

Ammonium fluoxycolumbate columbium fluoride, $3\text{NH}_4\text{F}$, CbOF_3 , CbF_3 .

(Marignac.)

Cupric fluoxycolumbate, CuF_2 , $\text{CbOF}_3 + 4\text{H}_2\text{O}$.

Sl. deliquescent. Sol. in H_2O . (Marignac, A. ch. (4) 8. 42.)

Potassium fluoxycolumbate, 2KF , $\text{CbOF}_3 + \text{H}_2\text{O}$.

Sol. in 12.5-13 pts. H_2O at 17-21°. Much more sol. in hot H_2O , or H_2O containing HF . (Marignac.)

3KF , CbOF_3 . Decomp. by H_2O into above salt. (M.)

5KF , $3\text{CbOF}_3 + \text{H}_2\text{O}$. Sol. in H_2O . (M.)

4KF , $3\text{CbOF}_3 + \text{H}_2\text{O}$. Sol. in H_2O . (M.)

3KF , $2\text{Cb}_2\text{O}_3 + 5\text{H}_2\text{O}$. Sl. sol. in H_2O . (Petersen, J. pr. (2) 40. 287.)

KF , $\text{Cb}_2\text{O}_3 + 3\text{H}_2\text{O}$. Sl. sol. in H_2O . (Petersen.)

2KF , $3\text{Cb}_2\text{O}_3\text{F}$. Insol. in H_2O . Sol. in HF . (Krüss and Nilson, B. 20. 1689.)

See also Fluoxypercolumbate, potassium.

Potassium hydrogen fluoxycolumbate, 3KF , HF , CbOF_3 .

Sol. in H_2O . (Marignac.)

Sodium fluoxycolumbate, 2NaF , $\text{CbOF}_3 + 2\text{H}_2\text{O}$.

Sol. in H_2O .

NaF , $\text{CbOF}_3 + \text{H}_2\text{O}$. (Marignac.)

Zinc fluoxycolumbate, ZnF_2 , $\text{CbOF}_3 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac, A. ch. (4) 8. 41.)

Fluoxyhypomolybdic acid.

Ammonium fluoxyhypomolybdate, MoOF_3 , $2\text{NH}_4\text{F}$.

Decomp. by H_2O . (Mauro, Gazz. ch. it. 19. 179.)

3MoOF_3 , $5\text{NH}_4\text{F} + \text{H}_2\text{O}$. Decomp. by H_2O . (Mauro.)

Cupric fluoxyhypomolybdate, CuF_2 , $\text{MoOF}_3 + 4\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . (Mauro, Real. Ac. Linc. 1892, 1. 194.)

Potassium fluoxyhypomolybdate, MoOF_3 , $2\text{KF} + \text{H}_2\text{O}$.

Sol. in H_2O with decomp.

Sol. in HF or $\text{HCl} + \text{Aq.}$ (Mauro and Panabianco, Gazz. ch. it. 12. 80.)

3MoOF_3 , $5\text{KF} + \text{H}_2\text{O}$. Sol. in H_2O with decomp. (Mauro, Gazz. ch. it. 19. 179.)

Thallous fluoxyhypomolybdate, 2TlF , MoOF_3 .

(Mauro, B. 1894, 27R. 109.)

Zinc fluoxyhypomolybdate, ZnF_2 , $\text{MoOF}_3 + 6\text{H}_2\text{O}$.

Rapidly deliquescent. Sol. in H_2O . (Mauro, Real. Ac. Linc. 1892. 1. 194.)

Fluoxyhypovanadic acid.

See Fluoxyvanadic acid.

Fluoxymanganic acid.

Ammonium fluoxymanganate, $(\text{NH}_4)_2\text{MnOF}_4$.
Precipitate. (Nicklès.)

Potassium fluoxymanganate, K_2MnOF_4 .
Precipitate. (Nicklès, C. R. 65. 107.)

Sesquifluoxymanganic acid.

Potassium sesquifluoxymanganate,
 $\text{K}_4\text{Mn}_2\text{OF}_8 = 4\text{KF}$, Mn_2OF_4 .
Precipitate. (Nicklès.)

Fluoxymolybdic acid.

See also Fluoxyhypomolybdic, and fluoxypermolybdic acids.

Ammonium fluoxymolybdate, NH_4F , MoO_3F_2 .

Sol. in H_2O . (Mauro, Gazz. ch. it. 20. 109.)

$+ \text{H}_2\text{O}$. More sol. in H_2O than $2\text{NH}_4\text{F}$, MoO_3F_2 . (Delafontaine, N. Arch. Sci. ph. nat. 30. 250.)

Correct formula is $3\text{NH}_4\text{F}$, MoO_3F_2 . (Mauro, Gazz. ch. it. 18. 120.)

$2\text{NH}_4\text{F}$, MoO_3F_2 . Much more sol. than 2KF , MoO_3F_2 . (Delafontaine.)

$3\text{NH}_4\text{F}$, MoO_3F_2 . Sol. in H_2O . (Mauro.)

$5\text{NH}_4\text{F}$, $3\text{MoO}_3\text{F}_2 + \text{H}_2\text{O}$. Sol. in H_2O . (Mauro, Gazz. ch. it. 20. 109.)

Ammonium fluoxymolybdate molybdate,
 MoO_3F_2 , $4\text{NH}_4\text{F}$, $(\text{NH}_4)_2\text{MoO}_4$.

Sol. in H_2O , but with decomp. (Mauro, Gazz. ch. it. 18. 120.)

Cadmium fluoxymolybdate, CdF_2 , $\text{MoO}_3\text{F}_2 + 6\text{H}_2\text{O}$.

Sl. efflorescent. (Delafontaine, J. B. 1867. 236.)

Cobaltous fluoxymolybdate, CoF_2 , M , $6\text{H}_2\text{O}$.

Sol. in H_2O . (Delafontaine, J. 236.)

Cupric fluoxymolybdate, CuF_2 , M , $4\text{H}_2\text{O}$.

Deliquescent. (Mauro, Real. 1892, 1. 194.)

Nickel fluoxymolybdate, NiF_2 , M , $6\text{H}_2\text{O}$.

Sol. in H_2O . (Delafontaine, J. 236.)

Potassium fluoxymolybdate, 2KF , M , H_2O .

Easily sol. in boiling H_2O .

Sol. in H_2O with decomp. Sol. (Marchetti, Z. anorg. 1895, 10. 68.)

KF , $\text{MoO}_3\text{F}_2 + \text{H}_2\text{O}$. Gradually cent. (Delafontaine.)

Rubidium fluoxymolybdate, 2RbF , $+ 2\text{H}_2\text{O}$.

Sol. in cold, more sol. in hot H_2O (fontaine.)

Sodium fluoxymolybdate, NaF , M , $\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . (Delafontaine.)

Thallous fluoxymolybdate, 2TlF , M , H_2O .

Sol. in hot H_2O . (Delafontaine.)

Zinc fluoxymolybdate, ZnF_2 , MoO_3F_2 .
Sol. in H_2O . (Delafontaine.)

Fluoxypercolumbic acid.

Potassium fluoxypercolumbate, 2KF , $+ \text{H}_2\text{O}$.

(Piccini, Z. anorg. 2. 21.)

Sol. in H_2O with decomp.

Sol. in HF . (Marchetti, Z. anorg. 10. 67.)

Fluoxypermolybdic acid.

Ammonium fluoxypermolybdate,
 $3\text{NH}_4\text{F}$.

Sol. in H_2O . (Piccini, Z. anorg. 1.

Cæsium fluoxypermolybdate, MoO_3 , $+ \text{H}_2\text{O}$.

(Piccini.)

Potassium fluoxypermolybdate, M , $2\text{KF} + \text{H}_2\text{O}$.

Not very sol. in H_2O ; more sol. in Aq without decomp. (Piccini.)

n fluoxypermolybdate, MoO_3F_2 , $\text{F} + \text{H}_2\text{O}$.

What more sol. in H_2O than K salt.
Sol. in $\text{HF} + \text{Aq}$. (Piccini.)

ertantalic acid.

m fluoxypertantalate, 2KF , TaO_3F_2 , H_2O .

Insol. in H_2O . (Piccini, Z. anorg. 2. 21.)

ertitanic acid, TiO_3F_2 , HF .

Sol. only in solution. (Piccini, B. 18.)

um fluoxypertitanate, TiO_3F_2 , BaF_2 .

Insol. in H_2O . (Piccini, Gazz. ch. it. 17.)

um fluoxypertitanate, $3\text{NH}_4\text{F}$. Sol. in H_2O .

Insol. in H_2O_2 . (Piccini, Z. anorg. 1895, 10.)

um fluoxypertitanate, $3\text{NH}_4\text{F}$. Sol. in H_2O . (Piccini, 8 R.)

um fluoxypertitanate, TiO_3F_2 , BaF_2 .
Insol. in H_2O . Easily sol. in acids. (Piccini, 8 R.)

um fluoxypertitanate, 3BaF_2 . Insol. in H_2O ; sol. in dil. acids. (Piccini, Gazz. ch. it. 17. 479.)

n fluoxypertitanate, TiO_3F_2 , 2KF .
Insol. in H_2O . (Piccini, B. 21. 1391.)

Insol. in H_2O_2 . (Piccini, Z. anorg. 1895, 10.)

ertungstic acid.

n fluoxypertungstate, 2KF , WO_3F_2 +

H_2O . (Piccini, Z. anorg. 2. 11.)

intalic acid.

o Fluoxypertantalic acid.

um fluoxytantalate, $3\text{NH}_4\text{F}$, TaOF_3 .
Sol. in H_2O . The solution clouds up on warming. (Joly, C. R. 81.)

tanic acid.

o Fluoxypertitanic acid.

uoxytitanate, TiOF_2 , BaF_2 .

Insol. in H_2O ; sol. in dil. acids. (Piccini, it. 17. 479.)

ingstic acid.

m fluoxytungstate, $2\text{NH}_4\text{F}$, WO_3F_2 .
Sol. in H_2O . (Marignac, A. ch. (3) 69.)

NH_4F , $\text{WO}_3\text{F}_2 + \text{H}_2\text{O}$. Decomp. by H_2O .
Crystallises unchanged from H_2O containing HF . (Marignac.)

Ammonium fluoxytungstate tungstate,
 $4\text{NH}_4\text{F}$, WO_3F_2 , $(\text{NH}_4)_2\text{WO}_4$.

Incompletely sol. in H_2O . Residue dissolves in $\text{NH}_4\text{OH} + \text{Aq}$. (Marignac.)

Cadmium fluoxytungstate.

Very sol. in H_2O . (Marignac.)

Cupric fluoxytungstate, CuF_2 , $\text{WO}_3\text{F}_2 + 4\text{H}_2\text{O}$.

Very sol. in H_2O . (Marignac, C. R. 55. 888.)

Cupric fluoxytungstate ammonium fluoride,
 CuF_2 , WO_3F_2 , $\text{NH}_4\text{F} + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Marignac.)

Manganese fluoxytungstate.

Very sol. in H_2O . (Marignac.)

Nickel fluoxytungstate, NiF_2 , $\text{WO}_3\text{F}_2 + 10\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O . (Marignac.)

Potassium fluoxytungstate, KF , $\text{WO}_3\text{F}_2 + \text{H}_2\text{O}$.

Can be recrystallised without decomp. only from H_2O containing HF . (Marignac, A. ch. (3) 69. 70.)

2KF , $\text{WO}_3\text{F}_2 + \text{H}_2\text{O}$. Difficultly sol. in cold, more easily in hot H_2O . (Berzelius.)

Sol. in 17 pts. H_2O at 15° . (Marignac.)

Can be recrystallised without decomp. from H_2O , or H_2O containing HF . (Marignac.)

Sol. in H_2O with decomp. Sol. in HF . (Marchetti, Z. anorg. 1895, 10. 71.)

See also Fluoxypertungstate, potassium.

Silver fluoxytungstate.

Very easily sol. in H_2O . (Marignac.)

Sodium fluoxytungstate, 2NaF , WO_3F_2 .

More sol. in H_2O than the corresponding K compound. (Berzelius.)

Thallous fluoxytungstate, TlF , WO_3F_2 .

Insol. in H_2O . Decomp. by H_2O . (Ephraim and Heymann, B. 1909, 42. 4463.)

2TlF , WO_3F_2 . Insol. in H_2O but decomp. thereby. (Ephraim and Heymann, B. 1909, 42. 4462.)

3TlF , $2\text{WO}_3\text{F}_2$. Insol. in H_2O . Decomp. by acids. (Ephraim and Heymann, B. 1909, 42. 4462.)

Zinc fluoxytungstate, ZnF_2 , $\text{WO}_3\text{F}_2 + 10\text{H}_2\text{O}$.

Very sol. in H_2O . (Marignac.)

Fluoxyuranic acid.**Ammonium fluoxyuranate**, $3\text{NH}_4\text{F}$, UO_2F_2 .Easily sol. in H_2O , less in HF . Insol. in alcohol. (Bolton.)100 g. solution sat. at 27° contain 10.11 g. salt.100 g. solution sat. at 81.3° contain 20.71 g. salt.

(Bürger, Dissert. 1904.)

Barium fluoxyuranate, 3BaF_2 , $2\text{UO}_2\text{F}_2 + 2\text{H}_2\text{O}$.Traces dissolve in hot H_2O . Easily sol. in dil. acids. (Bolton.)**Potassium fluoxyuranate**, 3KF , UO_2F_2 .Sol. in 8 pts. H_2O at 21° . Insol. in alcohol and ether. (Bolton, J. pr. 99. 269.)

Does not exist. (Smithells, Chem. Soc. 43. 125.)

 4KF , UO_2F_2 . Insol. in H_2O . Easily sol. in dil. acids. (Ditte, C. R. 91. 115.) 5KF , $2\text{UO}_2\text{F}_2$. (Baker, Chem. Soc. 35. 760.) 3KF , $2\text{UO}_2\text{F}_2 + 2\text{H}_2\text{O}$. (Baker.)**Sodium fluoxyuranate**, NaF , UO_2F_2 . $+2\text{H}_2\text{O}$. Not efflorescent. $+4\text{H}_2\text{O}$. Insol. in H_2O and dil. acids. Sl. sol. in conc. $\text{HCl} + \text{Aq}$. Sol. in conc. H_2SO_4 . (Bolton, J. B. 1866. 212.) 4NaF , UO_2F_2 . (Ditte.)

Does not exist. (Smithells, Chem. Soc. 43. 125.)

Fluoxyvanadic acid.**Ammonium fluoxyvanadate**, $12\text{NH}_4\text{F}$, V_2O_5 , 2VOF_3 .Easily sol. in H_2O , and not attacked by cold conc. H_2SO_4 . (Baker, Chem. Soc. 33. 388.)Formula is $3\text{NH}_4\text{F}$, VO_2F . (Petersen, J. pr. (2) 40. 289.) $3\text{NH}_4\text{F}$, VO_2F . Sol. in H_2O . (Petersen, l. c.)Much less sol. in H_2O in presence of NH_4F . (Piccini and Giorgis, Gazz. ch. it. 27. 1. 65.) $+ \text{H}_2\text{O}$. (Piccini and Giorgis, Gazz. ch. it. 1892, 22. 55.) $3\text{NH}_4\text{F}$, VOF_3 . "Hypovanadate." Quite sol. in H_2O . Very sl. sol. in $\text{MF} + \text{Aq}$. Less sol. in alcohol than in H_2O . (Petersen, J. pr. (2) 40. 195.) $2\text{NH}_4\text{F}$, VOF_3 . Sol. in H_2O . (Petersen.) $+ \text{H}_2\text{O}$. (Piccini and Giorgis.) $7\text{NH}_4\text{F}$, $4\text{VOF}_3 + 5\text{H}_2\text{O}$. Very sol. in H_2O . (Petersen.) $3\text{NH}_4\text{F}$, $2\text{VO}_2\text{F}$. Sol. in H_2O without decomp. Sol. in conc. $\text{HF} + \text{Aq}$. (Piccini and Giorgis, Gazz. ch. it. 24. 1. 68.) $3\text{NH}_4\text{F}$, $2\text{VOF}_3 + \text{H}_2\text{O}$. Sol. in H_2O with decomp. V_2O_5 , $2\text{NH}_4\text{F}$. (Ditte, C. R. 103. 270.) V_2O_5 , $8\text{NH}_4\text{F} + 4\text{H}_2\text{O}$. As above.
 V_2O_5 , $4\text{NH}_4\text{F} + 4\text{H}_2\text{O}$. As above.
 H_2O .**Ammonium hydrogen fluodioxyvanadate**, $7\text{NH}_4\text{F}$, HF , $4\text{VO}_2\text{F}$.Very sol. in H_2O . (Petersen, J. p. 284.)**Ammonium hydrogen trifluoxyvanadate**, 3HF , $9\text{NH}_4\text{F}$, 5VOF_3 .Easily sol. in H_2O . Sl. sol. in MF . (Petersen, J. pr. (2) 40. 280.) $3\text{NH}_4\text{F}$, 3HF , 2VOF_3 . Sol. in (Baker, Chem. Soc. 33. 388.)Identical with 3HF , $9\text{NH}_4\text{F}$, 5 (Petersen.)**Barium fluoxyvanadate**, BaF_2 , VO_2F .

Ppt. (Ephraim, Z. anorg. 1903)

Cadmium fluoxyvanadate, CdF_2 , $7\text{H}_2\text{O}$.

"Hypovanadate." As the Co salt (Piccini and Giorgis.)

Cobalt fluoxyvanadate, CoF_2 , VOF_3 ."Hypovanadate." Sol. in H_2O . (Piccini and Giorgis.)**Nickel fluoxyvanadate**, NiF_2 , VOF_3 .

"Hypovanadate." As the Co salt (Piccini and Giorgis.)

Potassium fluoxyvanadate, 7KF , 3Very sl. sol. in H_2O and $\text{MF} + \text{A}$ sol. in dil. acids. (Petersen, J. p. 199.) 2KF , VOF_3 . As above. (Petersen, J. pr. (2) 40. 289.) 2KF , $2\text{V}_2\text{O}_5 + 8\text{H}_2\text{O}$. Sol. in H_2SO_4 . (Ditte, C. R. 103. 1067.) 2KF , $3\text{V}_2\text{O}_5 + 5\text{H}_2\text{O}$. As above. 2KF , $4\text{V}_2\text{O}_5 + 8\text{H}_2\text{O}$. As above. 4KF , V_2O_5 . Less sol. than $4\text{KF} + 2\text{H}_2\text{O}$, and $+3\text{H}_2\text{O}$. Sol. in H_2O . (Ditte, C. R. 103. 1067.) 8KF , $\text{V}_2\text{O}_5 + 2\text{H}_2\text{O}$, and $+3\text{H}_2\text{O}$. Sol. in H_2O .**Potassium trifluoxyvanadate**, 2KF , V_2O_5 , $2\text{VOF}_3 + 2\text{H}_2\text{O}$.

Ppt. (Petersen, J. pr. (2) 40. 270.)

 6KF , V_2O_5 , $2\text{VOF}_3 + 2\text{H}_2\text{O}$. Sol. in cold conc. H_2SO_4 . (Baker, Chem. Soc. 33. 300.)Formula is 3KF , $2\text{VO}_2\text{F}$. (Piccini and Giorgis.)

See also Fluovanadate fluoxy potassium.

Potassium fluodioxyvanadate, 2KF , VO_2F .Easily sol. in H_2O . (Petersen, J. p. 278.)

O_2F . As above. (Petersen.)
 VO_2F . Sol. in H_2O ; scarcely at-
 H_2SO_4 . (Piccini and Giorgis.)

hydrogen fluoxyvanadate, $3KF$,
 VOF_3 .

H_2O . (Petersen.)

fluoxyvanadate, $8NaF$, $3VOF_3$ +

H_2O . (Petersen, J. pr. (2) 40. 200.)
 VO_2F , VOF_3 (?). Very easily de-
 iccini and Giorgis.)
 $2V_2O_5 + 10H_2O$. Sol. in H_2O .
 R. 106. 270.)

V_2O_5 . As above.

$V_2O_5 + 18H_2O$. As above.

$V_2O_5 + 5H_2O$. As above.

$V_2O_5 + 3H_2O$. (As above.)

fluoxyvanadate, $2TlF$, VOF_3 .

at sol. in cold H_2O without decomp.
 boiling H_2O with exception of a
 due, which is easily sol. in dil.
 Ephraim, B. 1909, 42. 4460.)

VO_2F . Insol. in H_2O . Sol. in H_2O
 H_2SO_4 . (Ephraim and Hey-
 1909, 42. 4459.)

yanadate, ZnF_2 , ZnO , $2VOF_3$ +

on air; sol. in H_2O . (Baker,
 33. 388.)

position is represented by the for-
 $VO_2F + 7H_2O$. (Petersen.)

$VO_2F + 7H_2O$. Very sol. in H_2O .
 id Giorgis.)

$VOF_3 + 7H_2O$. "Hypovanadate."
 ld H_2O , but decomp. by boiling;

$HF + Aq$. (Piccini and Giorgis.)

nic acid.

a fluozirconate, $(NH_4)_2ZrF_6$.

H_2O .
 ZrF_4 . Sol. in H_2O . (Marignac.)

fluozirconate, $2CdF_2$, $ZrF_4 + 6H_2O$.

H_2O ; can be recrystallised therefrom.
 A. ch. (3) 60. 257.)

$+ 6H_2O$. Sol. in H_2O . (Marignac.)

uozirconate, CsF , $ZrF_4 + H_2O$.

H_2O without decomp. (Wells,
 1895, 10. 434.)

F_4 . Sol. in H_2O without decomp.
 anorg. 1895, 10. 434.)

$ZrF_4 + 2H_2O$. Only sl. sol. in H_2O .
 anorg. 1895, 10. 434.)

zirconate, $2CuF_2$, $ZrF_4 + 12H_2O$.

l. in cold H_2O . (Marignac, A. ch.
 .)

$2ZrF_4 + 16H_2O$. Sol. in H_2O .
 .)

Lithium fluozirconate, $2LiF$, ZrF_4 .

Ppt. (Wells, Am. J. Sci. 1897, (4) 3.
 468.)

$4LiF$, $ZrF_4 + \frac{1}{2}H_2O$. Sol. in H_2O with
 decomp. (Wells, Am. J. Sci. 1897, (4) 3.
 469.)

Magnesium fluozirconate, $MgZrF_6 + 5H_2O$.

Sol. in H_2O . (Marignac.)

Manganous fluozirconate, $MnZrF_6 + 5H_2O$.

Sol. in H_2O . (Marignac, J. pr. 83. 202.)

Nickel fluozirconate, $2NiF_2$, $ZrF_4 + 12H_2O$.

Sol. in H_2O . (Marignac, A. ch. (3) 60.
 291.)

$NiZrF_6 + 6H_2O$. Sol. in H_2O . (Marig-
 nac.)

Nickel potassium fluozirconate, K_2ZrF_6 ,

$NiZrF_6 + 8H_2O$.

Sol. in H_2O . (Marignac.)

Potassium fluozirconate, KF , $ZrF_4 + H_2O$.

Much more sol. in hot, than cold H_2O .
 (Marignac.)

$2KF$, $ZrF_4 = K_2ZrF_6$. 100 pts. H_2O dis-
 solve at 2° , 0.781 pt.; at 15° , 1.41 pts.; at 19°
 1.69 pts.; at 100° , 25.0 pts. K_2ZrF_6 . (Marig-
 nac.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J.
 1898, 20. 829.)

$3KF$, ZrF_4 .

Sodium fluozirconate, $5NaF$, ZrF_4 .

100 pts. H_2O dissolve 0.387 pt. at 18° , and
 1.67 pts. at 100° . (Marignac.)

$2NaF$, ZrF_4 . (Wells, Am. J. Sci. 1897, (4)
 3. 469.)

$5NaF$, $2ZrF_4$. Sol. in H_2O with decomp.
 (Wells.)

Tellurium fluozirconate, TeF , ZrF_4 .

Sol. in H_2O without decomp. (Wells,
 Am. J. Sci. 1897, (4) 3. 470.)

$+ H_2O$. Sol. in H_2O without decomp.
 (Wells.)

$3TeF$, ZrF_4 . Sol. in H_2O without decomp.
 (Wells.)

$5TeF$, $3ZrF_4$. Sol. in H_2O without decomp.
 (Wells.)

Zinc fluozirconate, $ZnZrF_6 + 6H_2O$.

Sol. in H_2O . (Marignac.)

$2ZnF_2$, $ZrF_4 + 12H_2O$. Sol. in H_2O . (Mar-
 ignac, A. ch. (3) 60. 257.)

Fulminating gold.

See Auroamidoimide.

Fulminating platinum.

See Fulminoplatinum.

Fulminating silver.

See Silver nitride.

Fulminoplatinum compounds.

See—

Dichlorofulminoplatinum.

Trichlorofulminoplatinum.

Tetrachlorofulminoplatinum.

Chloroxyfulminoplatinum.

Fuscocobaltic chloride, $\text{Co}(\text{NH}_3)_4(\text{OH})\text{Cl}$, $+\text{H}_2\text{O}$.

Sol. in H_2O , from which it is precipitated by $\text{NH}_4\text{Cl}+\text{Aq}$; decomp. by boiling H_2O ; pptd. from aqueous solution by alcohol. (Fremy, C. R. 32. 501.)

— nitrate, $\text{Co}(\text{NH}_3)_4(\text{OH})(\text{NO}_3)_2+\text{H}_2\text{O}$.

Sol. in H_2O . Properties as the chloride. (Fremy.)

— sulphate, $\text{Co}(\text{NH}_3)_4(\text{OH})\text{SO}_4+1\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . Insol. in $\text{NH}_4\text{OH}+\text{Aq}$. (Fremy, C. R. 32. 501.)

Insol. in H_2O . Sol. in conc. $\text{HCl}+\text{Aq}$, or H_2SO_4 , from which it is precipitated by H_2O . (Vortmann, N. 6. 412.)

Fusible white precipitate.

See Mercuridiammonium chloride.

Gadolinium, Gd.

(Marignac, C. R. 102. 92.)

Gadolinium bromide, $\text{GdBr}_3+6\text{H}_2\text{O}$.

Sol. in HBr . (Benedicks, Z. anorg. 1900, 22. 403.)

Gadolinium chloride, $\text{GdCl}_3+6\text{H}_2\text{O}$.

Somewhat deliquescent. Sol. in H_2O . (Benedicks.)

Gadolinium platinum chloride.

See Chloroplatinate, gadolinium.

Gadolinium fluoride, GdF_3 .

Insol. in H_2O ; sl. sol. in hot HF . (Popovici, B. 1908, 41. 635.)

Gadolinium hydroxide, $\text{Gd}(\text{OH})_3$.

Ppt. (Benedicks, Z. anorg. 1900, 22. 402.)

Gadolinium oxide, Gd_2O_3 .

Sol. in acids. (de Boisbaudran, C. R. 111. 394.)

Somewhat hygroscopic; easily sol. in acids. (Benedicks.)

Gallium, Ga.

Not decomp. by H_2O ; easily sol. in cold $\text{HCl}+\text{Aq}$. Slowly sol. in warm dil. HNO_3+Aq . Not attacked by conc. HNO_3 free from

N_2O_5 below $40-50^\circ$, and only slowly presence of N_2O_5 . (Dupré, C. R. 83. 72.) Easily sol. in cold or warm $\text{KOH}+$ (de Boisbaudran, A. ch. (5) 10. 100.)

Gallium bromide, GaBr_3 .

Deliquescent, and sol. in H_2O .

Gallium dichloride, GaCl_2 .

Deliquescent, and decomp. by H_2O . (son and Petersen, C. R. 107. 527.)

Gallium chloride, GaCl_3 .

Deliquescent, and very sol. in little H_2O . Decomp. by much H_2O , with formation of basic salt, which is slowly sol. in dil. $+\text{Aq}$.

Gallium hydroxide.

Sol. in acids; sol. in KOH or NaOH less easily in $\text{NH}_4\text{OH}+\text{Aq}$, even in presence of ammonium salts.

Gallium iodide, GaI_3 .

Deliquescent, and sol. in H_2O . (de Boisbaudran and Jungfleisch, C. R. 86. 578.)

Gallium suboxide, Ga_2O_3 (?).

Sol. in HNO_3+Aq . (Dupré.)

Sol. in dil. $\text{H}_2\text{SO}_4+\text{Aq}$.

Gallium oxide, Ga_2O_3 .

Sol. in acids.

Germanium, Ge.

Insol. in $\text{HCl}+\text{Aq}$. Easily sol. in regia. Decomp. by HNO_3+Aq to Conc. H_2SO_4 decomp. to sulphate. In boiling $\text{KOH}+\text{Aq}$. (Winkler, J. pr. (1) 177; 36. 177.)

Germanium tetrabromide, GeBr_4 .

Decomp. by H_2O . (Winkler.)

Germanium dichloride, GeCl_2 .

Decompl by H_2O . (Winkler.)

Germanium tetrachloride, GeCl_4 .

Sinks in H_2O , and is gradually decomposed thereby. (Winkler, J. pr. 34. 177.)

Insol. in and not attacked by hot H_2SO_4 . (Friedrich, W. A. B. 102, 2b.)

Germanium chloroform, GeHCl_3 .

Decomp. by H_2O . Sol. in HCl . (Winkler.)

Germanium tetrafluoride, GeF_4 .

Deliquescent, and sol. in $\text{H}_2\text{O}+3\text{H}_2\text{O}$. Deliquescent. Melts in crystal H_2O when warmed. (Winkler.)

ium potassium fluoride.
luogermanate, potassium.

ium hydride, GeH_4 .

l. in hot HCl . Sol. in $\text{NaOCl} + \text{Aq.}$
n, Z. anorg. 1902, 30. 327.)

ium tetraiodide, GeI_4 .

escent, and sol. in H_2O with decomp.
.)

um monoxide, GeO .

ppreciably sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$
l. in $\text{HCl} + \text{Aq.}$ Insol. in alkalies.
J. pr. (2) 34. 177.)

hat sol. in H_2O ; insol. in $\text{H}_2\text{SO}_4 + \text{Aq.}$
n hot and conc. (van Bemmelen, R.
15.)

um dioxide, GeO_2 .

ry difficultly sol. in H_2O .

247.1 pts. H_2O at 20° ; in 93.3 pts.
(Winkler.)

sol. in alkali carbonates or hydrates
sol. in acids.

um oxychloride, GeOCl_2 .

in H_2O ; sol. in acids. (Winkler, J.
177.)

um monosulphide, GeS .

402.9 pts. H_2O . Sol. in conc. hot
Sol. in $\text{KOH} + \text{Aq.}$ Sol. in
+ Aq when precipitated. Insol. in
+ Aq if crystalline. Also exists in a
state. (Winkler.)

um disulphide, GeS_2 .

221.9 pts. H_2O . Easily sol. in
q, or $\text{NH}_4\text{OH} + \text{Aq.}$ Insol. in acids.
o in a colloidal state. (Winkler.)

um and extensive researches have
e on the action of H_2O and various
on glass. The older work has a cer-
rical interest, but only a brief state-
ome of the more important results
iven here. For a very thorough
f the work before the year 1861,
Dictionary, p. 555, should be con-

s is more or less attacked by H_2O ,
easily the greater the amount of
sent, the finer it is powdered, and
the temperature.

hat of a flask, is decomposed to a consider-
y several days' boiling with H_2O , a portion
alkali being dissolved, but when powdered
ed with distilled H_2O in a mortar, the H_2O
and exhibits no alkalinity. (Scheele.)
lumbics is partially dissolved by long boil-
(Lavoisier.)

etc potash or soda from glass together with

a portion of the silica, the decomposition taking place
the more easily in proportion as the glass is richer in
alkalies, more minutely divided, or the temperature of
the water higher. (Bischof, Kasn. Arch. 1. 442.)

Powdered crown glass and some varieties of window
glass render cold H_2O alkaline when in contact there-
with. (Dumas.)

100 pts. finely divided flint glass lose 7 pts. potash
when boiled one week with H_2O . (Griffiths, Q. J. Sci.
30. 258.)

Retorts of ordinary or flint glass are partially dis-
solved by H_2O when it is evaporated therein. (Chev-
reuil, 1811.)

Finely powdered plate-glass (Faraday, Pogg. 18. 509),
and Thuringian potash glass (Ludwig, Arch. Pharm. 91.
47) reddens moistened turmeric paper

The alkaline reaction disappears by continued wash-
ing, but reappears when the glass is freshly rubbed.
(Griffiths.)

Cold H_2O takes up SiO_2 as well as alkali from glass
powder. (Fuchs.)

Powdered lead glass gives up appreciable amounts of
 PbO to weakly acidified H_2O . (Pelouze.)

When powdered white glass, containing 12.4% Na_2O ,
15.5% CaO , and 72.1% SiO_2 , is treated repeatedly with
 H_2O , more than 8% of the glass is dissolved, and the
undissolved part gives up 1.5% CaO to $\text{HCl} + \text{Aq}$ with
effervescence. A glass containing more alkali, i. e.
16.3% Na_2O , 6.4% CaO , 77.3% SiO_2 , lost with the same
treatment 18.2%, and the residue gave up 2% CaO to
 $\text{HCl} + \text{Aq}$. (Pelouze, C. R. 43. 117.)

In the above case the fineness of the glass has an
influence as well as its composition. When the same
sample of glass was boiled 1 hour with H_2O , amounts
were dissolved in the proportion 1 : 4 : 28, according as
the glass was in form of a coarse, fine, or very fine
powder. Glass of the composition of the above samples,
as given by Pelouze, lost 10 and 32% respectively.

If powdered glass is boiled with H_2O and CO_2 con-
ducted into the solution, it is absorbed; if boiled with
 K_2SO_4 , Na_2SO_4 is dissolved. (Pelouze.)

Glass tubes are converted into a white crystalline
mass by heating with H_2O several months to $75-150^\circ$;
lead glass and Bohemian glass most easily, English
crown glass least. A little H_2O attacks glass more
than much H_2O .

The action of H_2O is greatly increased by
finely pulverizing the glass.

H_2O dissolved 10% of a glass containing
12% Na_2O , 15.5% CaO , and 72.5% SiO_2 , and
32% of another glass containing 16.3% Na_2O ,
6.4% CaO , and 77.3% SiO_2 . (Vogel, B. A.
Munchen, 1867. 437.)

Action of H_2O on a glass containing 74%
 SiO_2 , 8.6% CaO , 14% Na_2O , 0.6% K_2O , with
traces of Al_2O_3 , Fe_2O_3 , MnO , and MgO .

By boiling with H_2O a decrease of 3.9 mg.
was observed for the first hour, which soon
became constant at 2.2 mg. per hour. The
action was then proportional to the time, and
also to the surface in contact with the liquid,
but independent of the amount of liquid
evaporating.

The action decreases rapidly with the tem-
perature, so that at $90-100^\circ$ only $\frac{1}{4}$ as much
glass is dissolved as by boiling H_2O . (Emmer-
ling, A. 180. 257.)

When steam condenses in tubes of Na glass,
they are so strongly attacked that the H_2O
has an alkaline reaction, but tubes of hard or
Bohemian K glass are not so strongly at-
tacked. (Tollens, B. 9. 1540.)

The effect of H_2O is so great as to impart a
distinctly alkaline reaction to water condens-
ing in a tube of ordinary glass. By condensing
water in long tubes of various kinds of glass
the following results were obtained.

I. Easily fusible Thuringian glass. Surface exposed = 324 sq. cm.

After 2 hours, 62.0 mg. KOH were dissolved.

After 3 hours more, 36.0 mg. KOH were dissolved.

After 3 hours more, 33.2 mg. KOH were dissolved.

After 3 hours more, 20.8 mg. KOH were dissolved.

After 3 hours more, 20.8 mg. KOH were dissolved.

Or, in 14 hours, 172.8 mg. KOH were dissolved.

II. Less easily fusible Thuringian glass. Surface exposed = 499 sq. cm.

After 3 hours, 19.2 mg. KOH were dissolved.

After 3 hours more, 15.2 mg. KOH were dissolved.

After 3 hours more, 12.4 mg. KOH were dissolved.

After 3 hours more, 11.2 mg. KOH were dissolved.

Or, after 12 hours, 58.0 mg. KOH were dissolved.

III. Combustion tubing of very difficultly fusible Bohemian glass. Surface exposed = 1130 sq. cm.

After 3 hours 4.16 mg. KOH were dissolved.

After 3 hours more 4.16 mg. KOH were dissolved.

After 3 hours more 4.16 mg. KOH were dissolved.

After 3 hours more 4.16 mg. KOH were dissolved.

Or, after 12 hours, 16.64 mg. KOH were dissolved.

IV. Easily fusible Bohemian glass. Surface exposed = 1394 sq. cm.

After 3 hours, 7.88 mg. KOH were dissolved.

After 3 hours more, 8.56 mg. KOH were dissolved.

After 3 hours more, 1.97 mg. KOH were dissolved.

Or, after 9 hours, 24.32 mg. KOH were dissolved. (Kreusler and Henzold, B. 17. 34.)

From the above the following table has been calculated.

50 ccm. H₂O dissolves from a surface of 1000 sq. m. in 1 hour:—

96.0 mg. from easily fusible Thuringian glass.

12.8 mg. from less fusible Thuringian glass.

1.2 mg. from combustion tube of Bohemian glass.

2.0 mg. from harder tube of Bohemian glass.

(Kreusler and Henzold, B. 17. 34.)

100 ccm. H₂O dissolves so much glass from a flask every 2 seconds when in contact therewith that 0.1 ccm. $\frac{1}{2}$ normal oxalic acid is neutralised thereby. (Bohlig, Z. anal. 23. 518.)

Action of H₂O on various kinds of Na glass
1 g. of finely powdered glass was boiled 10 minutes in a silver dish with 100 ccm. H₂O, and the per cent of Na₂O (or K₂O) in the solution was determined.

	% Na ₂ O (K ₂ O)
Orthoclase feldspar	0.17
Glass of a Bohemian combustion tube	0.85
“ flask (German manuf.)	0.89
“ champagne bottle	1.7
Natrolite	1.82
Glass of a wine bottle (Hungarian)	2.5
Glass which was attacked by H ₂ O under pressure	3.1
Lead glass	3.8
Glass that broke easily	4.8
Glass tubing that became rough when fused	6.1
Glass tubing that became opaque by fusing	14.36
Solid water glass	26.97

(Wartha, Z. anal. 24. 220.)

The relative ease by which various kinds of glass are attacked by H₂O is shown by the following table. The glass was powdered and heated on a water bath with exclusion of atmospheric CO₂.

Potassium water glass	291
Sodium water glass	196
Yellow glass rich in alkali	34
Thuringian glass	19
Ditto from Tittel and Co.	8
Window glass	8
Lead glass from Jena	6
Bohemian glass from Kavalier	2.4
Lead crystal glass	1.4
Thermometer glass, 16IV, from Jena	1.0
Zinc glass, 362, from Jena	0.8
Lead glass, 434, from Jena	0.6
Lead glass, 483, from Jena	0.2
Heaviest lead silicate, from Jena	0.0

(Mylius, C. C. 1888. 1313.)

Solubility of various kinds of glass in H₂O.

The amounts dissolved from various kinds of glass by heating 5 hours with H₂O were as follows.

Yellow glass rich in alkali (13% K ₂ O, 15% Na ₂ O)	249 mg.
Poor Thuringian glass (6.6% K ₂ O, 16.5% Na ₂ O)	91.4 “
Glass from Tittel and Co. (7.1% K ₂ O, 14.3% Na ₂ O)	30.4 “
Bottle glass from Schilling (4.2% K ₂ O, 11.9% Na ₂ O)	13.0 “
Bohemian glass from Kavalier (13.3% K ₂ O, 11.4% Na ₂ O)	10.1 “
Rhenish window glass (13.5% Na ₂ O)	8.4 “
Lead crystal glass from Ehrenfeld (12.1% K ₂ O)	8.5 “
Green bottle glass (1.3% K ₂ O, 9.5% Na ₂ O)	6.5 “

Stability of various kinds of glass in H₂O—
Continued.

thermometer glass 16III from Jena (14.0% Na ₂ O, 7% ZnO)	6.4 mg.
and glass, No. 483, from Jena (47% PbO, 7.3% K ₂ O)	3 3 "
and silicate	0.6 "

(Mylius and Forster, B. 22. 1100.)

By calculation from the electrical conductivity of the solutions formed, various data were obtained by Kohlrausch (B. 24. 3565), which showed that different varieties of glass were attacked in very different degree by cold H₂O, and, moreover, the amount dissolved was proportionately much greater during the first few minutes of treatment with H₂O than afterwards, and, furthermore, the rate of decrease was much faster for good glass than poor. Increase of temperature increased the rate of solubility to a very great degree, the increase for 1° C. being about 17%. In 7 hours at 80° half as much was dissolved as in 6 months at 18°. Extensive tables are given. (Kohlrausch, B. 2. 3651.) See also Kohlrausch (W. Ann. 44. 77).

A very extensive research on the action of H₂O on glass, with a historical review of the work previously done on the subject, has been published by Mylius and Forster. (Z. anal. L 241.) The general results may be summarized as follows—

1. The solution of glass in H₂O is caused by decomposition, by which free alkali is freed.
2. The silicic acid of the glass is brought to solution by a secondary reaction of the free alkali in the solution.
3. The constituents of the solution change according to the conditions of the digestion.
4. The amount of alkali going into solution on a given surface under certain conditions is a measure for the resistance of a glass under one condition.
5. The rate of attack of glass surfaces by cold H₂O decreases rapidly with the length of time of digestion, and finally approaches a constant value.
6. The solubility increases very rapidly with increase of temperature.
7. The ratio of the solubility of several kinds of glass is dependent on the temperature.
8. From glasses which show the same ease of attack unequal amounts of substance may be dissolved.
9. The solubility of a glass is influenced by the condition of the surface from "weathering" by prolonged exposure to the CO₂ and H₂O of the air.
10. The poorer a glass is the less will its solubility decrease by prolonged treatment with H₂O.
11. A good glass is essentially less easily

attacked after having been previously treated with H₂O.

12. After treatment with H₂O, glass surfaces have the property of fixing alkali from the solutions formed, and giving it up again by a subsequent treatment with H₂O.

13. Potassium glass is much more sol. than sodium glass (contrary to previous researches), but the difference decreases as the glass becomes richer in CaO.

14. In glass flasks which are to be only slightly attacked by cold or hot H₂O, the CaO, alkalis, and SiO₂ must stand in a fixed relation to each other.

15. Of the more common varieties of glass, lead flint glass is least sol. in H₂O, but its surface is corroded, and it is easily decomposed by acids.

(Mylius and Forster, Z. anal. 31. 241.)

Bottle glass containing much Al₂O₃ is easily attacked by acids.

From powdered flint glass, boiling HCl + Aq extracts K, but no Pb. (Griffiths.)

Bottles of flint glass with (NH₄)₂CO₃ + Aq became so fragile that on shaking pieces of glass were detached. (Griffiths.)

All glass is decomp. by HF.

Conc. H₃PO₄ also attacks all glass.

Glass containing small amounts of BiO₂ are attacked by H₂SO₄, poorer glass by boiling HCl, HNO₃, and aqua regia. (Berzelius.)

Conc. HNO₃ does not act on flint glass at 145–150°. (Sorby, C. R. 50. 990.)

Glass of ordinary chemical apparatus gives up traces of metals to HCl and HNO₃ + Aq, but hard Bohemian glass consisting of 75% SiO₂, 15% K₂O, 10% CaO, resists the action of warm conc. acids; also an easily fusible Na K glass with 77% SiO₂, 7.7% K₂O, 5% Na₂O, 10.3% CaO, is not easily attacked. (Stas.)

KOH, and NaOH + Aq dissolve SiO₂ from glass the more easily the hotter and the more conc. the solutions are. (Müller.) NH₄OH, and (NH₄)₂CO₃ + Aq attack many kinds of glass, especially flint glass. CaO₂H₂ attacks glass appreciably at 45° and lower; still more strongly on boiling. (Lamy, A. ch. (5) 14. 155.)

The action of various solvents on the glass mentioned on page 359 in Emmerling's experiments is as follows:

The action of HCl + Aq containing 0.2 to 3% HCl is practically null, but is increased either by dilution or concentration. A very small quantity (0.02%) HCl added to H₂O almost wholly prevents its action on glass. With HCl + Aq (11% HCl) a decrease of 4.2 mg. was noticed in the first hour, and only 3–4 mg. afterwards. The same is the case for HNO₃ + Aq in still greater degree, 0.008% HNO₃ sufficing to nearly counteract the solvent action of H₂O.

H₂SO₄ + Aq has about double the solvent effect possessed by H₂O.

Oxalic and acetic acids both diminish the solvent action of H₂O.

The addition of even traces (0.04%) of

Na_2CO_3 increases the solvent action, and this is further rapidly increased by an increase in the amount of Na_2CO_3 . $\text{Na}_2\text{CO}_3 + \text{Aq}$ containing 1% Na_2CO_3 dissolves about 10 times as much as pure H_2O , i. e. about 35 mg. per hour.

The above is also the case with $\text{KOH} + \text{Aq}$, but in even greater degree. $\text{KOH} + \text{Aq}$ containing 0.025% KOH dissolved three times as much as pure H_2O .

$(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ has about the same action as H_2O .

With $\text{NH}_4\text{OH} + \text{Aq}$ (9% NH_3) 7 mg. decrease for the first hour, and 3 mg. afterwards was noticed. The concentration of the $\text{NH}_4\text{OH} + \text{Aq}$ was apparently without effect.

The addition of NH_4Cl decreases the solvent action of H_2O proportionately to the amount added, but with new flasks large amounts are dissolved.

With $\text{NH}_4\text{Cl} + \text{Aq}$ (7% NH_4Cl) 4.2 mg. were dissolved in the first hour, and the amount dissolved gradually decreased to null after 24 hours on account of the liberation of HCl by the decomp. of NH_4Cl .

NaCl , KCl , KNO_3 , and Na_2SO_4 show a similar behaviour to that of NH_4Cl .

$\text{Na}_2\text{HPO}_4 + \text{Aq}$ containing 0.4% Na_2HPO_4

has six times the solvent action of pure H_2O , but the action is not increased by further concentration.

In general, those salts the acids of which form insol. Ca salts, as Na_2CO_3 , Na_2HPO_4 , $(\text{NH}_4)_2\text{C}_2\text{O}_4$, increase the solvent action of H_2O , and this effect is greater the more concentrated the solution. KNO_3 , NH_4Cl , and CaCl_2 decrease the effect, and the stronger the solution the less the effect.

All Na glass with approximately the above composition has the same power of resistance against H_2O ; Bohemian K glass shows greater resistance, especially against acids (Emmerling, A. 150. 257.)

Action of various reagents on hard Bohemian glass. 100 ccm. substance dissolved in glass in 6 days at 100° .

H_2O	10.0
$\text{H}_2\text{S} + \text{Aq}$	8.7
Dil. $(\text{NH}_4)_2\text{S} + \text{Aq}$	52.5
Conc. $(\text{NH}_4)_2\text{S} + \text{Aq}$	47.2
Conc. $\text{NH}_4\text{OH} + \text{Aq}$	42.5
Dil. $\text{NH}_4\text{OH} + \text{Aq}$	7.7
$\text{NH}_4\text{SH} + \text{Aq}$	51.2

(Cowper, Chem. Soc. 41. 254.)

Action of various solutions on glass of different composition. (The figures denote decrease in weight in mg. of a 100 ccm. flask.)

	Time	1	2	3	4	5	6	7	8	9	10
H_2O	5 hrs.	62	31	29	17	13	9	7	7	5	4
$\text{H}_2\text{SO}_4 + \text{Aq}$ (25% H_2SO_4)	3 "	..	43	35	8	7	6	5	5	5	5
$\text{HCl} + \text{Aq}$ (12% HCl)	3 "	85	..	27	4	2	1	1	1	0	0
$\text{NH}_4\text{OH} + \text{Aq}$ (10% NH_3)	3 "	62	11	8	7	7	6	5	4
$\text{Na}_2\text{HPO}_4 + \text{Aq}$ (12% Na_2HPO_4)	3 "	81	64	40	35	34	30	15	13
$\text{Na}_2\text{CO}_3 + \text{Aq}$ (2% Na_2CO_3)	3 "	283	160	130	124	50	45	42	42	26	22

Composition of above varieties of glass.

	1	2	3	4	5	6	7	8	9	10
SiO_2	76.22	74.09	76.39	68.56	74.48	74.69	66.75	74.12	77.07	74.40
Al_2O_3	..	0.40	0.50	1.85	0.50	0.45	1.31	0.50	0.30	0.70
CaO	4.27	5.85	5.50	7.60	7.15	7.85	13.37	8.55	8.10	8.85
K_2O	..	7.32	4.94	2.24	6.64	8.64	05.50	4.85	3.75	4.40
Na_2O	19.51	12.34	12.67	19.75	11.23	8.37	3.07	11.97	10.78	11.65

It is seen that glass which resists the attack of H_2O also resists acids and alkalis, and that the relative resistance of all varieties to any of the solutions is the same. Therefore the action of H_2O may be accepted as a criterion for judging of the resistance of a glass to all solvents. Glass No. 10, in which the molecular ratio of $\text{SiO}_2 : \text{CaO} : \text{K}_2\text{O} : \text{Na}_2\text{O}$ is 8.1:1.5, is recommended as best suited for chemical uses. (Weber and Sauer, B. 25. 70.)

Mylus and Forster (B. 25. 97) recommend a glass in which the molecular ratio of $\text{SiO}_2 : \text{CaO} : \text{K}_2\text{O} : \text{Na}_2\text{O}$ is 7.2:1:1.1 as the best suited for chemical apparatus.

In an exhaustive research on the action of aqueous solutions on glass, which cannot be given in full on account of its great length, the following conclusions are reached:—

1. Solutions of caustic alkalis act on glass much more strongly than H_2O , dissolving all the constituents of the glass—that is, the glass as such. Very dilute solutions form an exception.

2. Of the caustic alkalis, $\text{NaOH} + \text{Aq}$ has the strongest action, then come KOH , NH_4OH , and $\text{BaO} \cdot \text{H}_2\text{O} + \text{Aq}$ in the order named.

3. Increase in temperature increases the

length of the attack of alkalis very considerably.

4. At high temperatures, the ease with which glass is attacked increases at first rapidly with the concentration of the alkali, but afterwards more slowly.

5. At ordinary temperatures very concentrated alkali solutions have less action on glass than dil. solutions.

6. Solutions of pure alkalis, if not too concentrated, act less on glass than when contaminated with small amounts of SiO_2 .

7. Alkali carbonates + Aq. attack glass much more than H_2O , even when they are very dilute. The action corresponds less to that of the caustic alkalis than to that of other salts. With equivalent concentration, Na_2CO_3 + Aq. has a stronger action than K_2CO_3 + Aq.

8. The action of salt solutions on glass is a compound one, depending both on the concentration and the kind of salt dissolved, and is made up of the action of the H_2O and the salt in solution.

9. Each kind of attack is differently influenced by the composition of the glass.

10. Solutions of those salts, the acids of which form insol. Ca salts, have a stronger action than H_2O , and the action increases with the concentration.

11. Solutions of those salts, the acids of which form sol. Ca salts, have less action than H_2O , and the action decreases with the concentration. (Förster, B. 25. 2494.)

Data on this subject published since the first edition of this work have not been considered.

Glucinic acid.

Potassium glucinate, K_2GluO_2 .

Very deliquescent. Sol. in H_2O and acids. (Krüss and Moraht, B. 23. 733.)

Glucinum (Beryllium), Gl.

Not attacked by hot or cold H_2O . Sol. in cold dil. HNO_3 + Aq. (Wöhler, Pogg. 13. 577.)

Sol. only in boiling conc. HNO_3 + Aq. (Debray, A. ch. (3) 44. 5.)

Sol. in dil. HCl + Aq. dil. and conc. H_2SO_4 + Aq. and KOH + Aq. but insol. in NH_4OH + Aq. (Wöhler, Debray.)

Sol. in hot HCl , hot conc. HNO_3 , and hot conc. H_2SO_4 . (Lebeau, A. ch. 1899, (7), 16. 474.)

Glucinium azoimide, GluN_3 .

Decomp. by hot H_2O . (Curtius, J. pr. 1898, (2), 58. 292.)

Glucinum borocarbide, $3\text{Glu}_2\text{C}_3\text{B}_2\text{C}$.

Insol. in H_2O . Easily sol. in mineral acids especially HNO_3 . (Lebeau, A. ch. 1899, (7) 16. 481.)

$\text{C}_4\text{B}_5\text{Glu}_6$. Stable in air.

Easily sol. in mineral acids, conc. and dil. (Lebeau, C. R. 1898, 126. 1349.)

Glucinum bromide, GluBr_3 .

Sol. in H_2O with evolution of much heat. (Wöhler.)

Sol. in abs. alcohol. (Lebeau, A. ch. 1899, (7) 16. 484.)

Glucinum carbide, Glu_2C .

Decomp. by H_2O . Slowly attacked by cold or hot conc. HCl and HNO_3 . Gradually but completely sol. in dil. acids. (Lebeau, A. ch. 1899, (7), 16. 476.)

GluC_2 . Not easily decomp. by strong acids. Easily sol. with decomp. in dil acids. (Lebeau, C. R. 1895, 121. 497.)

Glucinum chloride, GluCl_2 .

Anhydrous. Fumes and deliquesces in air. Sol. in H_2O with hissing and evolution of much heat. Easily sol. in alcohol.

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 828.)

Sol. in alcohol and ether. Sl. sol. in C_6H_6 , CHCl_3 , Cl_4 and CS_2 . (Lebeau, A. ch. 1899, (7) 16. 493.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+4 H_2O . Very hygroscopic.

Easily sol. in H_2O and in alcohol. (Mieleitner, Z. anorg. 1913, 80. 73.)

Glucinum chloride ammonia, $\text{GluCl}_2 \cdot 4\text{NH}_3$.

(Mieleitner, Z. anorg. 1913, 80. 73.)

Glucinum chloride iodine trichloride, $2\text{GluCl}_2 \cdot \text{GlCl}_2 + 8\text{H}_2\text{O}$.

Hygroscopic. (Weinland, Z. anorg. 1902, 30. 140.)

Glucinum ferric chloride, $\text{GluCl}_2 \cdot \text{FeCl}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O . (Neumann, A. 244. 329.)

Glucinum mercuric chloride, $\text{GluCl}_2 \cdot 3\text{HgCl}_2 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Atterberg, B. 6. 1288.)

Glucinum thallic chloride, $3\text{GluCl}_2 \cdot 2\text{TlCl}_3$.

Cryst. from HCl solution. (Neumann, A. 244. 348.)

Glucinum stannic chloride.

See Chlorostannate, glucinum.

Glucinum fluoride, basic, $2\text{G}\text{I}\text{O}$, $5\text{G}\text{I}\text{F}_2$.

Sol. in H_2O . (Lebeau, A. ch. 1899, (7) 16. 484.)

Glucinum fluoride, $\text{G}\text{I}\text{F}_2$.

Deliquescent. Sol. in H_2O . Insol. in anhydrous HF . Sol. in alcohol. (Lebeau, C. R. 1898, 126. 1421.)

Sol. in H_2O in all proportions. Somewhat sol. in abs. alcohol. Easily sol. in 90% alcohol, also in a mixture of alcohol and ether. Insol. in anhydrous HF . (Lebeau, A. ch. 1899, (7) 16. 484.)

Glucinum potassium fluoride, $\text{G}\text{I}\text{F}_2$, KF .

Sl. sol. in H_2O . (Awdejew.) Much more sol. in hot than cold H_2O . (Berzelius.)

$\text{G}\text{I}\text{F}_2$, $2\text{K}\text{F}$. Sol. in about 50 pts. H_2O at 20° , and 19 pts. boiling H_2O . (Marignac.)

Glucinum sodium fluoride, $\text{G}\text{I}\text{F}_2$, $2\text{Na}\text{F}$.

Sol. in 34 pts. H_2O at 100° , and 68 pts. at 18° . (Marignac.)

Glucinum hydroxide, $\text{G}\text{I}\text{O}_2\text{H}_2$.

Easily sol. in acids. Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in $\text{CO}_2 + \text{Aq}$; 100 ccm. sat. $\text{CO}_2 + \text{Aq}$ dissolve 0.0185 g. GIO . (Sestini, Gazz. ch. it. 20. 313.)

Also sol. in KOH , NaOH , NH_4OH , or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$, especially when freshly precipitated; also in Na_2CO_3 , or $\text{K}_2\text{CO}_3 + \text{Aq}$. (Debray.)

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$ containing $\text{NH}_4\text{Cl} + \text{Aq}$.

Very sl. sol. in $\text{Li}_2\text{CO}_3 + \text{Aq}$. (Gmelin.)

Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Berthier.)

Sol. in $\text{BaO}_2\text{H}_2 + \text{Aq}$, from which it is pptd. by NH_4 salts, but not by boiling. Sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$ when freshly pptd.

Sol. in $\text{NH}_4\text{F} + \text{Aq}$. (Helmholtz, Z. anorg. 3. 130.)

Solubility of $\text{G}\text{I}\text{O}_2\text{H}_2$ in $\text{NaOH} + \text{Aq}$.

G. Na in 20 ccm.	G. Gl. in 20 ccm.
0.3358	0.0358
0.6717	0.0882
0.8725	0.1175
1.7346	0.2847

(Rubenbauer, Z. anorg. 1902, 30. 334.)

When glucinum hydroxide is treated with alkali, more dissolves at first than corresponds with the true equilibrium under the prevailing conditions, for such solutions spontaneously deposit more or less glucinum hydroxide according to the concentration. (Rubenbauer.)

Solubility of freshly pptd. $\text{G}\text{I}\text{O}_2\text{H}_2$ in NaOH Aq at room temperature.

millimols Na per l.	G. GIO dissolved in l
649	3.6
540	2.92
540	2.53
483	1.69
383	1.64
388	1.53
386	1.45
390	1.24

In the first two cases the values were obtained by adding to $\text{G}\text{I}\text{Cl}_2 + \text{Aq}$ at 0° , ice NH_4OH and treating the ppt. with NaOH Aq. In the remaining cases by dissolving basic $\text{G}\text{I}\text{CO}_3$ in HCl and pouring into NaOH Aq. (Haber, Z. anorg. 1904, 38. 386.)

Solubility of $\text{G}\text{I}\text{O}_2\text{H}_2$, which is one week in $\text{NaOH} + \text{Aq}$ at t° .

NaOH	t°	G. GIO in
$\frac{1}{2}\text{-N}$	$20\text{-}23^\circ$	0.06
1-N	$20\text{-}23^\circ$	0.17
2-N	$20\text{-}23^\circ$	0.57
$\frac{1}{2}\text{-N}$	$50\text{-}53^\circ$	0.08
1-N	$50\text{-}53^\circ$	0.23
2-N	$50\text{-}53^\circ$	0.90
$\frac{1}{2}\text{-N}$	100°	0.08
1-N	100°	0.29
2-N	100°	1.02

(Haber.)

Solubility of $\text{G}\text{I}\text{O}_2\text{H}_2$ in $\text{NaOH} + \text{Aq}$ at

G. mols.	
Na	Gl
0.268	0.0330
0.318	0.0492
0.446	0.0841
0.526	0.089
0.563	0.101
0.801	0.143
0.854	0.202

(Wood, Chem. Soc. 1910, 97. 884.)

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$ and in alkyl alcohols. (Renz, B. 1903, 36. 2753.)

Sol. in $\text{G}\text{I}\text{SO}_4 + \text{Aq}$. (Parsons, J. Chem. 1907, 11. 658.)

A form insol. in acids and alkalis is obtained by sufficiently long heating in boiling H_2O , $\text{Na}_2\text{CO}_3 + \text{Aq}$, $\text{NH}_4\text{OH} + \text{Aq}$, solutions of NaOH or KOH so dil. that $\text{G}\text{I}\text{O}_2\text{H}_2$ is either insol. or very sl. sol. (van Oordt, C. C. 1906, 1. 108.)

100 ccm. of glycerine + Aq containing 60% by vol. of glycerine dissolve 0.1 g. (Müller, Z. anorg. 1905, 43. 322.)

Contains $\frac{1}{2}\text{H}_2\text{O}$ (Schaffgotzsch); $\frac{1}{2}\text{H}_2\text{O}$ (Terberg).

Solubility of GlO_2H_2 , which has been boiled with alkali, in various solvents.

Alkali used	Time	Solvent	Solubility
10-N NaOH	2½ hrs.	0.106-N NaOH 0.39-N NaOH 0.97-N NaOH 2.0-N NaOH 2.0-N NaOH	1 mol. GlO_2H_2 : 331 mole NaOH 1 mol. GlO_2H_2 : 183 mole NaOH 1 mol. GlO_2H_2 : 91.8 mole NaOH 1 mol. GlO_2H_2 : 49 mole NaOH 1 mol. GlO_2H_2 : 49 mole NaOH
NaOH NaOH	Long time 2 hrs.	10-N K_2CO_3 Dil. HCl Hot conc. HCl Dil. acetic acid	Insol. Sl. sol. Slowly sol. Almost insol.
KOH	Till flocculent ppt. appeared	1-N KOH 1-N NaOH Warm dil. HCl	Insol. " Sol.
½-N Na_2CO_3 { ½-N Na_2CO_3 for then 1/10-N Na_2CO_3 for	3 hrs. 3 hrs. and 4 hrs.	1-N NaOH Dil. HCl 1-N NaOH Dil. HCl	Easily sol. " Sl. sol. Slowly sol.
½-N K_2CO_3 10-N K_2CO_3	6 hrs. Short time	10-N K_2CO_3 1-N NaOH Cold 10-N K_2CO_3	Insol. Easily sol. Very slowly sol.
NH_4OH	5 hrs.	½-N NaOH	Very sl. sol.

(Haber.)

Glucinum iodide, GII_2 .

Sol. in H_2O with evolution of much heat. (Wöhler.)

Decomp. in moist air and by H_2O .

Sol. in abs. alcohol. Insol. in most neutral organic solvents, as benzene, toluene, etc. (Lebeau, C. R. 1898, 126. 1273.)

Insol. in C_6H_6 , C_7H_8 , and oil of turpentine. Easily sol. in CS_2 . Sol. in anhydrous alcohols without evolution of heat. (Lebeau, A. ch. 1899, (7) 16. 490.)

Glucinum iodide ammonia, $2\text{GII}_2, 3\text{NH}_3$.

Ppt. (Lebeau, A. ch. 1899, (7) 16. 492.)

Glucinum oxide, GlO .

Crystalline. Insol. in acids except conc. H_2SO_4 . (Ebelmen, C. R. 32. 710.)

Amorphous. Absolutely insol. in H_2O . The higher the temp. to which the substance has been heated the more insol. is it in acids. Insol. in $\text{NH}_4\text{OH} + \text{Aq}$ or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Insol. in conc. $\text{NH}_4\text{Cl} + \text{Aq}$, or KOH , and $\text{NaOH} + \text{Aq}$. (Rose.)

When obtained by ignition of GlSO_4 , it is very slowly but completely sol. in HCl , and $\text{H}_2\text{SO}_4 + \text{Aq}$. (Rose.)

Insol. in hydracids. Sol. in conc. H_2SO_4 . (Lebeau, C. R. 1896, 123. 819.)

Insol. in liquid NH_3 . (Gore, Am. ch. J. 1898, 20. 828.)

Glucinum peroxide basic, $2\text{GlO}_2, 3\text{GlO}$.

+ $8\frac{1}{2}\text{H}_2\text{O}$. (Komarovski, Chem. Soc. 1913, 104 (2) 707.)

Glucinum oxybromides.

Sol. in H_2O if three or less equivalents of base are present to one of acid; insol. if more of the base is present. (Ordway, Am. J. Sci. (2) 26. 207.)

Glucinum oxychloride, $\text{Gl}_2\text{OCl}_2 = \text{GlO}, \text{GlCl}_2$.

Insol. in H_2O .

$3\text{GlCl}_2, 2\text{GlO} + 2\text{H}_2\text{O}$ (?). Sol. in H_2O . (Atterberg.)

$\text{GlCl}_2, 3\text{GlO} + 3\text{H}_2\text{O}$ (?). Sol. in H_2O , but solution soon becomes cloudy and deposits a fine ppt. By boiling the solution it is decomp. into above salt, and $\text{GlCl}_2, 12\text{GlO}_2\text{H}_2 + 10\text{H}_2\text{O}$, which is insol. in H_2O ; decomp. into GlO_2H_2 by washing. Sol. in acids. (Atterberg.)

Glucinum oxyfluoride, $5\text{GlF}_2, 2\text{GlO}$.

Readily sol. in H_2O . (Lebeau, C. R. 1898, 126. 1419.)

Glucinum phosphide.

Decomp. by H_2O . (Wöhler.)

Glucinum selenide.

Sl. sol. in H_2O . (Berzelius.)

Glucinum sulphide.

Slowly sol. without decomp. in H_2O , but easily decomp. by acids. (Wöhler.)

Gold, Au.

Gold which has been pptd. from $AuCl_3 + Aq$ by $FeSO_4$ is al. sol. in HCl . (Auerkiew, Z. anorg. 1909, 81, 10.)

Not attacked by H_2O . Insol. in HNO_3 or $HCl + Aq$. Easily sol. in aqua regia or any mixture evolving Cl or Br . Sol. in selenic acid, or antimonie acid + Aq ; less easily in arsenic acid + Aq . Sol. in mixtures of HCl and nitrates, or HNO_3 and chlorides; also in $(NaCl + KNO_3 + K_2Al_2(SO_4)_4) + Aq$ (?). Insol. in H_2SO_4 , except in presence of $KMnO_4$, HNO_3 , or HIO_4 . Sol. in a solution of I in ether in direct sunlight.

Sol. in solutions of ferric, and cupric salts.

Sol. in $HCl + Aq$ containing H_2CrO_4 , H_2MnO_4 , H_2SeO_4 , H_2AsO_4 , or $FeCl_3$. (Wurtz.)

Attacked by fuming HCl (sp. gr. 1.178) at ord. temp. in direct light, especially in the presence of a trace of $MnCl_2$, but not attacked in the dark even in the presence of this salt. (Berthelot, C. R. 1904, 138, 1298.)

100 ccm. hot conc. HCl dissolve 0.008 g. yellow Au powder in 4 hours. (Hanriot and Raoult, C. R. 1912, 155, 1086.)

Upon boiling 25 and 50 cc. $HCl + Aq$ (sp. gr. 1.178), dil. to 125 cc. with 250 mg. sheet Au $\frac{1}{2}$ in. square, 0.009 in. thick, weighing 250 mg. for several hours, there was no loss of weight of Au . (McCaughy, J. Am. Chem. Soc. 1909, 31, 1263.)

From 5 g. finely divided ordinary yellow gold; 100 cc. HNO_3 of 22° B. dissolve 0.002 g. in 2 hrs.

100 cc. HNO_3 of 32° B. dissolve 0.0119 g. in 2 hrs.

100 cc. HNO_3 of 36° B. dissolve 0.028 g. in 2 hrs.

100 cc. HNO_3 monohydrate dissolve 0.076 g. in 2 hrs. (Hanriot and Raoult, C. R. 1912, 155, 1085.)

From 5 g. brown gold:

100 ccm. HNO_3 of 22° Baume dissolve 0.006 g. in 2 hrs.

100 ccm. HNO_3 of 32° Baume dissolve 0.039 g. in 2 hrs.

100 ccm. HNO_3 of 36° Baume dissolve 0.078 g. in 2 hrs.

100 ccm. HNO_3 monohydrate dissolve 1.540 g. in 2 hrs. (Hanriot and Raoult.)

Sl. sol. in boiling HNO_3 (sp. gr. = 1.42). The solution deposits Au by standing several days. (Dewey, J. Am. Chem. Soc. 1910, 32, 320.)

Best composition of aqua regia for dissolving Au is 200 cc. HCl (sp. gr. 1.1946) 45 cc. HNO_3 (sp. gr. 1.4) and 245 cc. H_2O . 1 pt. Au is sol. in 4.3 pts. of such a mixture. (Priwoznik, C. C. 1910, II, 1743.)

Sol. in 1 pt. $HNO_3 + 4$ pts. HCl as representing the most economical mixture. (Priwoznik, Chem. Soc. 1911, 100, (2), 484.)

Easily sol. in nitrosulphonic acid from phuric acid manufacture, when mixed with equal parts conc. $HCl + Aq$. (Berthelot, Rep. anal. Ch. 1887, 741.)

Sol. in hot conc. H_2SO_4 in the presence of MnO_2 , Mn_2O_3 , Mn_2O_4 , PbO_2 , Pb_2O_3 , PbO , CrO_3 , CrO_4 and Ni_2O_3 . Solution also takes place slowly in the cold. Sol. in hot $H_2SO_4 + KMnO_4$. Slowly sol. in cold, more rapidly in hot $H_2SO_4 + HNO_3$. (Lenher, J. Am. Chem. Soc. 1904, 26, 550.)

Sol. in a hot solution of crystalline telluric acid in H_2SO_4 or H_3PO_4 .

Sol. in hot H_3PO_4 in the presence of MnO_2 , Mn_2O_3 , Mn_2O_4 , the higher oxides of lead, CrO_3 , chromium tetroxide and nickel oxide. Solution takes place more slowly in the cold. Sol. in hot H_3PO_4 in the presence of $KMnO_4$. Slowly sol. in a cold, more rapidly sol. in a hot mixture of H_3PO_4 and HNO_3 .

Sol. in hot arsenic acid in the presence of MnO_2 , Mn_2O_3 and Mn_2O_4 . (Lenher, J. Am. Chem. Soc. 1904, 26, 550.)

Gold leaf is not attacked by cold conc. H_2SeO_4 , when completely free from halogens, but is dissolved by conc. H_2SeO_4 at 300°. (Lenher, J. Am. Chem. Soc. 1902, 24, 354.)

Solubility of thin sheet Au in HCl solution of iron alum.

Time, hours	Mg. Au dissolved			
	1 g. Fe 25 cc. HCl	1 g. Fe 50 cc. HCl	2 g. Fe 25 cc. HCl	2 g. Fe 50 cc. HCl

Temp. 38–43°.

16	1.00	1.30	1.08	1.47
22	1.12	1.55	1.20	1.81
40	1.52	2.15	1.82	2.75
46	1.71	2.34	2.02	2.95
64	1.96	3.10	2.60	3.79
72	2.12	3.30	2.83	4.05
89	2.32	3.65	3.22	4.65
100	2.40	3.76	3.38	4.81
113	2.45	3.95	3.51	5.12
124	2.60	4.09	3.63	5.30
161	2.78	4.36	3.95	5.60
185	2.90	4.49	4.11	6.22

Temp. 98–100°.

1	1.13	0.78	1.15	1.27
2	1.99	1.74	2.56	2.66
4	3.48	3.31	4.55	5.06
16	10.09	11.37	13.15	15.56
20	12.20	13.72	15.59	19.41
24	14.37	16.49	17.96	23.20
36	17.38	23.27	22.07	31.73
42	18.79	26.30	24.62	35.29
54	20.91	31.39	29.49	42.11
50½	21.64	33.12	30.64	44.43

duction contained the given amounts of iron alum, the sp. gr. of the HCl 8, and the solutions diluted to 125 cc. ghey, J. Am. Chem. Soc. 1909, 31.

solvent action of ferric salt occurs presence of a ferrous salt, but de- with increase of concentration of alt. (McCaughy.)

r of thin sheet Au in HCl solution of CuCl_2 .

Mg. Au dissolved

1 g. Cu 5 cc. HCl	1 g. Cu 50 cc. HCl	2 g. Cu 25 cc. HCl	2 g. Cu 50 cc. HCl
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Temp. 38-43°.

0.03	0.26	0.05	0.39
0.09	0.38	0.05	0.54
0.10	0.54	0.14	0.94
0.12	0.61	0.15	1.07
0.11	0.76	0.17	1.40
0.14	0.87	0.22	1.58
0.14	0.92	0.24	1.75
0.16	1.02	0.27	1.90
0.18	1.34	0.32	2.45
0.23	1.60	0.39	2.84

Temp. 98-100°.

0.15	0.34	0.17	0.46
0.55	1.23	0.55	1.35
1.34	5.00	2.12	8.80
1.63	6.52	2.78	11.86
2.17	9.13	3.59	15.70
3.13	13.98	5.07	23.14
3.61	16.54	5.77	26.62
4.07	19.26	6.26	30.80
4.82	26.37	7.47	39.09

ions the same as above for HCl +
t. (McCaughy.)

powdered gold is sol. in conc HCl
sence of alcohol, etc.

g. Au. is sol. in 100 cc. HCl + 100 cc.

g. Au. is sol. in 100 cc. HCl + 100 cc.

g. Au. is sol. in 100 cc. HCl + 100 cc.

g. Au. is sol. in 100 cc. HCl + 100 cc.

g. Au. is sol. in 100 cc. HCl + 50 cc.
 $(\text{OH})_2$.

verkiev, C. C. 1908, II. 1566.)

Solubility of Au which has been pptd. from
 $\text{AuCl}_3 + \text{Aq}$ by various precipitants in
HCl + formaldehyde.

HCHO (40%) ccm.	HCl (1.19) ccm.	Au dis- solved g.	Au used was pptd. from $\text{AuCl}_3 + \text{Aq}$ by
150	150	0.0007	FeSO_4
100	100	0.0006	"
125	25	0.0006	"
100	200	0.0009	sugar
150	150	0.0003	oxalic acid
250	250	0.0002	"
500	500	0.0008	FeSO_4
100	500	0	oxalic acid
100	200	0.0004	"
100	200	0	"
150	300	0.0006	sugar
250	125	0.001	"
150	75	0.0013	FeSO_4
100	200	0.0008	"
100	200	0.0006	HCHO
100	100	0.0005	"
180	540	0	
360	720	0.001	FeSO_4
360	720	0.0019	sugar
209	400	0.0013	FeSO_4

In the last four cases the solubility was
determined at the ordinary temp.; in the
first sixteen the gold was boiled with the
mixture of HCl and HCHO.

(Averkiew, Z. anorg. 1909, 61. 3.)

Solubility of Au in boiling HCl +
paraformaldehyde.

$(\text{CH}_2\text{O})_n$ g.	HCl (1.19) g.	Dissolved Au g.	Au used was pptd. from $\text{AuCl}_3 + \text{Aq}$ by
5	25	0.0005	FeSO_4
5	25	0.0004	oxalic acid
25	125	0.006	sugar
20	400	0.0034	CHOH
20	400	0.003	CH_2OH
20	400	0.0065	"
20	400	0.0044	sugar
20	400	0.0005	formic acid
20	400	0.0005	"
40	400	0.001	CH_2OH
20	300	0.0024	FeSO_4
20	300	0.003	"
10	200	0.0008	CH_2OH
10	200	0.0008	oxalic acid
60	120	0.0015	FeSO_4

(Averkiew.)

Solubility of Au in boiling HCl+methyl alcohol.

CH ₃ OH (99%) ccm.	HCl (1.19) ccm.	Dissolved Au g.	Au used was pptd. from AuCl ₃ +Aq. by
100	100	0.0302	FeSO ₄
150	150	0.0043	oxalic acid
150	150	0.028	sugar
25	25	0.001	CH ₃ OH
50	50	0.0002	oxalic acid
50	50	0.0005	CH ₃ OH
50	50	0.0002	oxalic acid
50	50	0.0018	FeSO ₄
50	50	0.002	"
100	100	0.0009	oxalic acid
500	500	0.0128	FeSO ₄
1000	1000	0.0281	"
50	100	0.0084	"
100	200	0.0006	"
100	300	0	"
75	25	0.005	HCOH
90	30	0.005	"
75	25	0.0014	"
80	20	0.0005	"
100	50	0.0018	FeSO ₄
100	50	0.0008	CH ₃ OH
50	200	0.001	"

(Averkiew.)

Solubility of Au in boiling HCl+ethyl alcohol.

C ₂ H ₅ OH (95%) ccm.	HCl (1.19) ccm.	Dissolved Au g.	Au used was pptd. from AuCl ₃ +Aq. by
25	50	0.0006	FeSO ₄
100	200	0.0111	"
200	400	0.0017	"
150	50	0.0003	sugar
100	300	0.0004	"
100	100	0.0015	"
200	200	0.0055	C ₂ H ₅ OH
250	250	0.0021	sugar
300	300	0.0197	FeSO ₄
1000	1000	0.007	CH ₃ OH
150	150	0.008	"

(Averkiew.)

Solubility of Au in boiling HCl+amyl alcohol.

C ₅ H ₁₁ OH g.	HCl (1.19) g.	Dissolved Au g.	Au used was pptd. from AuCl ₃ +Aq. by
100	100	0.019	FeSO ₄
100	200	0.0048	"
100	100	0.0024	sugar
100	100	0.0027	"
150	100	0.0032	"
300	100	0.0023	FeSO ₄
200	200	0.0067	C ₅ H ₁₁ OH
500	500	0.028	FeSO ₄

(Averkiew.)

Solubility of Au in boiling HCl+

C ₂ H ₅ OH g.	HCl (1.19) g.	Dissolved Au g.	Au pptd. from AuCl ₃
10+25	100	0.001	C
10	40	0.0004	ox
20	50	0.0003	
25	100	0.0005	
25	150	0	F
50	200	0.0005	
25	250	0.0005	
25	250	0.0012	

(Averkiew.)

Solubility of Au in boiling HCl+cl

CHCl ₃ g.	HCl (1.19) g.	Dissolved Au g.	Au pptd. from AuCl ₃
50	100	0.0009	
100	100	0.023	
100	100	0.0017	
100	50	0.0012	
200	400	0.0024	
250	250	0.002	
300	100	0.0106	

(Averkiew.)

Colloidal gold is sol. in dil. alkali B. 1902, 35. 2236.)

Rather quickly sol. in 10-15% I in KI+Aq.

Very slowly sol. in 5% solution of Aq.

Scarcely sol. in more dil. solution KI+Aq.

Easily sol. in 10% NH₄I+I. sol in 5% NH₄I+I. (Düring.)Sol. in cold Na₂S+Aq when Na₂S in proportion of 843 pts. Na₂S to (Becker, Sill, Am. J. (3) 33. 199.)

In finely divided state Au is sol. KCN+Aq. Not attacked by boil +Aq. (Vogel, J. pr. 20. 355.)

Solubility of Au (diaks) in KCN-

(A) oxygen passed through sol (B) agitated with oxygen.

% KCN	g Au dissolved in 24 h	
	A	
1	0.00845	0
5	0.01355	0
20	0.0115	0
50	0.00505	0

(MacLaurin, Chem. Soc. 1893, 61)

The solution of Au in KCN+essentially hastened by sunlight. (Proc. Chem. Soc. 1904, 20. 199.)

sence of oxygen is necessary for n of Au in KCN+Aq. The rate of Au in KCN+Aq varies with h of the solution, being small for tions, increasing as the solution ore dilute, reaching a maximum at N, and then again diminishing. (Chem. Soc. 1895, 67. 211.)

of Au (strips) in dil. KCN+Aq.

CN	Mg. Au dissolved in 24 hours
	0.010
5	0.043-0.07
	0.10-0.23
6	0.16
	0.44
25	1.77
	4.29
	48.43
	74.96
5	150.54
	168.12

, Elektrochem. Z. 1901, 7. 205.)

divided metallic gold is completely ord. temp. in solutions of potas- cyanide. Solution takes place, even when the potassium ferro- solution is boiling. (Beutel, Z. 2, 78. 158.)

RbCl.I+Aq. (Erdmann, Arch. 4, 232, 30.)

acked by FeCl₃+Aq when air is ut very energetically attacked in f HCl and oxygen. (McIlhiney, 1896, (4), 2. 293.)

acked by several days heating, at 150°. At 200° there is sl. 10 days. (North, J. Am. Chem. 34. 892.)

n excess dissolves pulverulent Au hours heating at 160°. (North, 4), 9. 647.)

Cl₂. (Baudrimont, A. ch. (4) 2.

ol. in acid solutions of thiocar- pecially in presence of suitable ompounds. (Moir, Chem. Soc 45.)

ide, AuAs.

leohol slowly extracts As; HNO₃+ ts into Au and H₃AsO₄. Sol. in

Not attacked by cold, decomp. . H₂SO₄. (Tivoli, C. C. 1887. 778; 610.)

thide, Au₂Bi.

ldonite. Sol. in aqua regia.

mide, AuBr.

H₂O. (Thomsen, C. C. 1860.

Insol. in H₂O, HNO₃, H₂SO₄.

Sol. in NH₄OH+Aq. with decomp.

Decomp. by HBr and KBr+Aq. Sol. in KCN+Aq without decomp. Slowly decomp. by alcohol, ether, acetone and moist CHCl₃. (Lengfeld, Am. Ch. J. 1901, 26. 325.)

Gold (auroauric) bromide, Au₂Br₄.

Not deliquescent. H₂O or ether dissolves out AuBr₃. (Thomsen, C. C. 1860. 606.)

Does not exist. (Krüss, B. 20. 640.)

Existence is maintained by Petersen. (J. pr. (2) 46. 334.)

Auric bromide, AuBr₃.

Not deliquescent. Slowly sol. in H₂O, more readily in ether.

Can be recryst. from AsBr₃, SbBr₃, TiBr₃ or SnBr₃. (Lindet, Bull. Soc. 1886, (2) 45. 149.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Aurous phosphorus tribromide, AuBr, PBr₃.

Decomp. by H₂O. (Lindet, J. pr. (2) 32. 494.)

Auric phosphorus pentabromide, AuBr₃, PBr₅.

Decomp. by H₂O. (Lindet.)

Aurous bromide phosphorus trichloride, AuBr, PCl₃.

Decomp. by H₂O. (Lindet.)

Auric praseodymium bromide, PrBr₃, AuBr₃+10H₂O.

Very sol. in H₂O; sol. in conc HBr. (Von Schule, Z. anorg. 1898, 18. 355.)

Aurous bromide ammonia, AuBr, 2NH₃.

Decomp. by H₂O and dil. HCl.

Sol. in aqua regia. (Meyer, C. R. 1906, 143. 281.)

Gold carbide, Au₂C₂.

Ppt. Decomp. by boiling H₂O without evolution of C₂H₂. Decomp. by HCl with evolution of C₂H₂. (Mathews, J. Am. Chem. Soc. 1900, 22. 110.)

Aurous chloride, AuCl.

Insol. in H₂O, but gradually decomp. thereby into Au and AuCl₃. (Thomsen, J. pr. (2) 13. 341.)

Insol. in H₂O and dil. HNO₃.

Decomp. by conc. HNO₃ to Au and AuCl₃.

Sol. in HCl, HBr and in sol. of alkali chloride and bromides, with decomp.

Decomp. by alcohol, ether and acetone. (Lengfeld, Am. Ch. J. 1901, 26. 324.)

Gold (auric) chloride, AuCl_3 .

Deliquescent. Very sol. in H_2O . Sol. in 1.47 pts. H_2O . (Abl.) Sol. in conc. HCl , or $\text{HNO}_3 + \text{Aq}$ without decomp.

AsCl_3 dissolves about 22% at 160° and 2.5% at 15° . Solubility in SbCl_3 is about the same. Much less sol. in SnCl_4 or TiCl_4 , SnCl_4 dissolving 4% at 160° and hardly a trace at 0° . Very sl. sol. in hot or cold SiCl_4 . (Lindet, Bull. Soc. (2) 45. 149.)

Sl. sol. in liquid NH_3 . (Franklin and Kraus, Am. Ch. J. 1898, 20. 829.)

Sol. in alcohol with gradual decomp. (Gmelin.) Sol. in ether with decomp. in light or on long standing. Ether extracts AuCl_3 from $\text{AuCl}_3 + \text{Aq}$ (Proust). Sol. in volatile oils with gradual decomp.

Sol. in ether. (Mylius, Z. anorg. 1911, 70. 207.)

Very sol. in ether. (Willstätter, B. 1903, 36. 1830.)

Completely sol. in ether. (Frank, C. C. 1913, II. 541.)

Insol. in or decomp. by alcohol, ether, CS_2 , C_6H_6 , oil of turpentine, pentane, hexane, CHCl_3 , CCl_4 , ethyl nitrate, nitrobenzol, ethyl acetate, ethyl propionate and pyridine. (Lenher, J. Am. Chem. Soc. 1903, 25. 1138.)

+ $2\text{H}_2\text{O}$. (Thomsen.)

Auroauric chloride, Au_2Cl_4 .

Decomp. by H_2O into AuCl_3 and AuCl . (Thomsen, J. pr. (2) 13. 357.)

Does not exist. (Krüss and Schmidt, J. pr. (2) 38. 77.)

Existence is maintained by Christensen. (J. pr. (2) 46. 328.)

Auric chloride with MCl .

See Chloraurate, M.

Auric nitrosyl chloride, AuCl_2 , NOCl .

Sol. in H_2O with decomp. (Sudborough, Chem. Soc. 59. 662.)

Aurous phosphorus trichloride, AuCl , PCl_3 .

Decomp. by H_2O . Sol. in about 100 pts. PCl_3 at 15° , and about 8 pts. at 120° . Sol. in AsCl_3 . (Lindet, C. R. 101. 1492.)

Auric phosphorus pentachloride, AuCl_3 , PCl_5 .

Decomp. by H_2O . Nearly insol. in PCl_5 . Sol. in AsCl_3 . (Lindet.)

Aurous potassium chloride, AuCl , KCl .

Decomp. by H_2O or $\text{HCl} + \text{Aq}$ into KCl , KAuCl_4 , and Au . (Berzelius.)

Auric potassium chloride.

See Chloraurate, potassium.

Auric selenium chloride, AuCl_3 , SeCl_4 .

Decomp. by H_2O . Sol. in AsCl_3 . (Lindet, C. R. 101. 1492.)

Gold (aurous) sodium chloride, AuCl

Insol. in H_2O . Sol. in alcohol. (J. Pharm. 3. 447.)

Formula is 4NaCl , AuCl , AuCl_3 . (sen.)

Auric sodium chloride.

See Chloraurate, sodium.

Auric sulphur chloride, AuCl_3 , SCl_4 .

Easily decomp. by H_2O . (Lindet, 101. 1492.)

Aurous chloride ammonia, AuCl , NH_3 .

Ppt. Unstable. (Diemer, J. Am. Soc. 1913, 35. 554.)

AuCl , 3NH_3 . Decomp. by H_2O acids.

Sol. in aqua regia. Sol. in conc with decomp. (Meyer, C. R. 1906, 14.

AuCl , 12NH_3 . (Meyer.)

Auric fluoride, AuF_3 .

Very unstable.

Is incapable of existence not only in presence of H_2O but under the ordinary conditions met with in the laboratory and in nature. (Lenher, J. Am. Chem. Soc. 1903, 25. 1138.)

Auric hydroxide, AuO_2H_3 .

Nearly insol. in most acids. Easily dissolved in very conc. $\text{HNO}_3 + \text{Aq}$ (Proust), from all AuO_2H_3 is separated by dilution. Extremely sl. sol. in fuming HNO_3 , dil. $\text{HNO}_3 + \text{Aq}$ when pure (Krüss, 281). Not attacked by H_3PO_4 . Insol. in HCl , or $\text{HBr} + \text{Aq}$ (Fremy).

Sol. in $\text{H}_2\text{SeO}_4 + \text{Aq}$. (Mitscherlich.) Sl. sol. in conc. H_2SO_4 ; somewhat in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Rose.)

Nearly insol. in cold $\text{KOH} + \text{Aq}$, dissolved on boiling. Insol. in NH_4OH or alkali carbonates + Aq (Rose). Slightly sol. in boiling $\text{CaCl}_2 + \text{Aq}$, $\text{NaCl} + \text{Aq}$, $\text{BaCl}_2 + \text{Aq}$ (Pelletier). Sol. in NH_4CN , and KCN (Himly).

Sl. sol. in KCl , or $\text{NaCl} + \text{Aq}$. (Fremy.) Sol. in $\text{K}_4\text{Fe}(\text{CN})_6 + \text{Aq}$ at ordinary temperature, rapidly on boiling. (Beutel, Z. anorg. 78. 154.)

AuO , $\text{OH} = \text{Au}_2\text{O}_3$, H_2O . (Krüss.)

Auroauric hydroxide, $\text{Au}_2\text{O}_3(\text{OH})_2$, + $2\text{H}_2\text{O}$.

Insol. in boiling conc. $\text{KOH} + \text{Aq}$. Slightly sol. by conc. HCl or $\text{HNO}_3 + \text{Aq}$ into Au_2O_3 , which dissolves. (Schöttgen, Z. anorg. 217. 336.)

Aurous iodide, AuI .

Insol. in cold, decomp. by hot H_2O , HCl , or $\text{HNO}_3 + \text{Aq}$, with separation of Au . Decomp. immediately by ether, not by alcohol.

ly sol. in KI, FeI_3 , or $\text{HI} + \text{Aq}$. Sl. attacked by NH_4OH , or aq at 35° (Fordos). Instantly de-
 $\cdot \text{KOH} + \text{Aq}$.

ric) iodide, AuI_3 .

in H_2O . Sol. in alkali iodides, and
 Decomp. on air or by alkalies.
 n, Phil. Mag. J. 9. 266.)

xide ammonia, AuI , NH_3 .

p. by H_2O or dil. acids. (Meyer,
 18, 143. 281.)

aqua regia.

NH_3 . (Meyer.)

xide, Au_2O .

n H_2O or alcohol. Decomp. by boil-
 $\cdot \text{HCl} + \text{Aq}$ into Au and AuCl_3 .
 HNO_3 , or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ do not
 Sol. in cold aqua regia. Sol. in HI
 sol. in KOH , or $\text{NaOH} + \text{Aq}$ when
 precipitated. (Berzelius.)

ing to Krüss (A. 237. 281) all hitherto
 Au_2O is impure. Pure Au_2O is sol.
 I_2O when freshly precipitated, from
 xeroxide is precipitated by boiling.
 ol. in HCl , or $\text{HBr} + \text{Aq}$. Sol. in
 $\cdot \text{NaOH} + \text{Aq}$ when freshly precipi-
 Not affected by any other acid or
 (Krüss.)

-called solution of Au_2O in H_2O is
 a colloidal suspension. (Vanino, B.
 462.)

de, Au_2O_3 .

ric hydroxide.

c oxide, Au_2O_3 .

cold $\text{HCl} + \text{Aq}$; forms insol. comp.
 (Prat, C. R. 70. 842.)

ed pure by Krüss (A. 237. 296.)

ephide, Au_4P_4 .

acked by $\text{HCl} + \text{Aq}$. HNO_3 forms
 and leaves undissolved Au . (Schröt-
 1849. 247.)

Decomp. on air or with H_2O . (Ca-
 uss. ch. it. 15. 40.)

Readily attacked by aqua regia
 aq . (Granger, C. R. 1897, 124. 498.)

ple (mixture of Au and SnO_2).

in H_2O . Easily sol. in aqua regia.
 Aq dissolves all Sn and leaves Au .
 $\cdot \text{HNO}_3 + \text{Aq}$ dissolves a little Sn .

in boiling $\text{KOH} + \text{Aq}$ (Berzelius).
 $\cdot \text{aq}$ extracts excess of SnO_2 , and the
 comes sol. in H_2O , from which it is
 $\text{NH}_4\text{Cl} + \text{Aq}$. (Figuier, A. ch. (3)

ben still moist, in $\text{NH}_4\text{OH} + \text{Aq}$, but
 has been dried.

ed in colloidal state in aqueous solu-
 aining 0.58 g. Au . and 5.41 g. SnO_2

in a litre. This solution may be concentrated
 without coagulation. The solution is coagul-
 ated by dil. HNO_3 , or $\text{HCl} + \text{Aq}$, more easily
 by dil. $\text{H}_2\text{SO}_4 + \text{Aq}$; also by KCl , HgCl_2 ,
 $\text{FeSO}_4 + \text{Aq}$, and many other salts. Not
 coagulated by alcohol, but easily when ether
 is added to the alcohol. (Schneider, Z. anorg.
 5. 80.)

Gold (auric) selenide, Au_2Se_3 .

$\text{HNO}_3 + \text{Aq}$ dissolves out Se . Sol. in aqua
 regia or alkali sulphides + Aq . (Uelsmann,
 J. B. 1860. 90.)

Aurous sulphide, Au_2S .

Easily sol. in H_2O when freshly prepared,
 but precipitated from aqueous solution by
 HCl , KCl , or $\text{NaCl} + \text{Aq}$. When dried is
 insol. in H_2O .

Insol. in boiling dil. or conc. HCl , or H_2SO_4
 $+ \text{Aq}$. Easily sol. in aqua regia, $\text{HCl} + \text{Aq}$
 with KClO_4 , etc. Slowly sol. in alkali mono-
 sulphides + Aq . Easily sol. in polysulphides
 $+ \text{Aq}$.

Insol. in $\text{KOH} + \text{Aq}$. Sol. in $\text{KCN} + \text{Aq}$.
 (Krüss, B. 20. 2369.)

Known also in colloidal state in aqueous
 solution containing 1.74 g. Au_2S per l.
 (Schneider, B. 24. 2241.)

Auric sulphide, Au_2S_3 .

Insol. in H_2O and acids except aqua regia;
 sol. in alkali sulphides, or $\text{KOH} + \text{Aq}$. (Ber-
 zelius.)

Does not exist (Krüss, B. 22. 2369), but
 has since been made by Antony and Lucchesi
 (Gazz. ch. it. 20. 601). Insol. in HCl , or dil.
 $\text{HNO}_3 + \text{Aq}$. Decomp. by conc. HNO_3 , KOH ,
 or $\text{NaOH} + \text{Aq}$ with separation of Au . Sl.
 decomp. by $\text{NH}_4\text{OH} + \text{Aq}$. Easily sol. in
 $\text{KCN} + \text{Aq}$; decomp. by $(\text{NH}_4)_2\text{S} + \text{Aq}$. Sol.
 in cold Na_2S or $\text{K}_2\text{S} + \text{Aq}$; decomp. on boil-
 ing. (Antony and Lucchesi, Gazz. ch. it. 21,
 2. 209.)

Insol. in ether. (Hofmann, B. 1904, 37.
 246.)

Auroauric sulphide, Au_2S_7 .

Insol. in H_2O or acids except aqua regia.
 Sl. sol. in cold alkali monosulphides + Aq ,
 but easily sol. on warming. Sol. in cold
 polysulphides + Aq , but less in ammonium
 polysulphide than the other alkali poly-
 sulphides.

Not attacked by cold, but easily sol. in hot
 $\text{KOH} + \text{Aq}$. Sol. in $\text{KCN} + \text{Aq}$. (Hoffmann
 and Krüss, B. 20. 2704.)

Obtained also in colloidal state in aqueous
 solution containing 0.8 g. per l. (Schneider.)

Insol. in $\text{Na}_2\text{S} + \text{Aq}$. sat. with S . (Ditte,
 A. ch. 1907, (8) 12. 273.)

Aurous potassium sulphide, Au_2S , $3\text{K}_2\text{S}$.

(Antony and Lucchesi, Gazz. ch. it. 1896,
 26. (2) 350.)

Au_2S , $4\text{K}_2\text{S}+12\text{H}_2\text{O}$. Very sol. in H_2O . (Ditte, C. R. 1895, 120. 322.)

Gold silver sulphide, Ag_3AuS_2 .

Ppt. (Lucchesi, Gazz. ch. it. 1896, 26. 350-53.)

Aurous sodium sulphide, $\text{NaAuS}+4\text{H}_2\text{O}$.

Sol. in H_2O and alcohol. (Yorke, Chem. Soc. Q. J. 1. 236.)

$+5\text{H}_2\text{O}$. Sol. in H_2O . (Ditte, C. R. 1895, 120. 321.)

Na_3AuS_2 . Sol. in H_2O . (Lucchesi, Gazz. ch. it. 1896, 26. 350-53.)

Au_2S , $2\text{Na}_2\text{S}+20\text{H}_2\text{O}$. Very sol. in H_2O . (Ditte, C. R. 1895, 120. 321.)

Gold telluride.

Ppt. (Berzelius, Pogg. 8. 178.)

Gold silver telluride, Au_2Te_2 , Ag_2Te .

Min. *Sylvanite*. Sol. in HNO_3 + Aq with separation of Au, in aqua regia with separation of AgCl.

$3\text{Ag}_2\text{Te}$, Au_2Te . Min. *Petzite*.

Hartshorn, salts of.

See Carbonate carbamate, ammonium hydrogen.

Helium, He.

Coefficient of absorption for H_2O at $18.2^\circ = 0.0073$. (Ramsay, Z. phys. Ch. 1906, 55. 347.)

Absorption of helium by H_2O at t° and 760 mm. pressure.

t°	Coefficient of absorption
0	0.01500
0.5	0.01487
5	0.01460
10	0.01442
15	0.01396
20	0.01386
25	0.01371
30	0.01382
35	0.01380
40	0.01387
45	0.01403
50	0.01404

(Estreicher, Z. phys. Ch. 1899, 31. 184.)

Absorption by H_2O at t° .

t°	Coefficient of absorption
0	0.0134
10	0.0100
20	0.0138
30	0.0161
40	0.0191
50	0.0226

(Antropoff, Roy. Soc. Proc. 1910, 83. A 480.)

Completely insol. in benzene and in alk. (Ramsay, Chem. Soc. 1895, 67. 684.)

Hexamine chromium compounds.

See Luteochromium compounds.

Hexamine cobaltic compounds,

$\text{Co}_2(\text{NH}_3)_6\text{X}_4$.

See Dichrocobaltic compounds.

$\text{Co}(\text{NH}_3)_6\text{X}_3$.

See Luteocobaltic compounds.

Hexamine iridium chloride, $\text{Ir}_3(\text{NH}_3)_6$

See Iridotriamine chloride.

Hexathionic acid, $\text{H}_2\text{S}_6\text{O}_4$.

Known only in aqueous solution, decomposes rapidly, even in presence of sulphuric acid. (Debus, A. 244. 76.)

Potassium hexathionate, $\text{K}_2\text{S}_6\text{O}_4$.

Sol. in H_2O , with rapid decomp. obtained in pure state. (Debus, A. 244.

Holmium, Ho.

Holmium oxide, Ho_2O_3 .

(Cleve, C. R. 89. 478; 91. 328.)

Consists of at least two elements. (de Boisbaudran, C. R. 102. 1005.)

Consists of seven elements. (Krtz Nilson.)

Sesqui-hydraurylamine, $(\text{AuOH})_2\text{N}$, $\text{Au}_2\text{N}_2+3\text{H}_2\text{O}$.

Decomp. by boiling with H_2O . (R. A. 235. 341.)

Hydrazidophosphoric acid.

Barium hydrazidophosphate,

$\text{OP}(\text{N}_2\text{H}_3)\text{O}_2\text{Ba}$.

(Ephraim, B. 1911, 44. 3420.)

Lead hydrazidophosphate, $\text{OP}(\text{N}_2\text{H}_3)\text{O}$

Easily sol. in HNO_3 . (Ephraim.)

Potassium hydrazidophosphate,

$\text{OP}(\text{N}_2\text{H}_3)(\text{OK})_2$. (Ephraim.)

Sodium hydrazidophosphate,

$\text{OP}(\text{N}_2\text{H}_3)(\text{ONa})_2$.

Can be cryst. from dil. alcohol. (Ephraim.)

Hydrazine, $\text{N}_2\text{H}_4 = \text{NH}_2 - \text{NH}_2$.

Very sol. in H_2O . (Curtius, B. 20.

Very hygroscopic; decomp. by H_2O ent for sulphur, KCl, KBr, KNO_3 Bruyn, R. t. c. 1894, 13. 433-40; Chem. Soc. 1895, 68 (2) 347.)

Mixes in all proportions with alk. sol. in organic solvents. (Lobry de Chem. Soc. 1897, 72 (2) 22.)

amidosulphonate,
 $\text{HSO}_3, \text{NH}_2$.
 in H_2O . (Sabanejeff, Z. anorg. 2.)

azoimide, $\text{N}_2\text{H}_4, \text{HN}_3$.
 cent. Easily sol. in H_2O . Sl. sol. and can be crystallised therefrom. 24. 2344.)

borate, $(\text{N}_2\text{H}_4)_2(\text{B}_2\text{O}_3)_6$.
 H_2O . (Dschawachow, C. C. 1902,

$\text{H}_2\text{B}_4\text{O}_7$). Sol. in H_2O . (Dschawachow, C. 1902, I. 1394.)
 Sol. in H_2O . (Dschawachow, I. 1394.)
 Ppt. (Dschawachow, C. C. 4.)

cuprous bromide chloride,
 $\text{N}_2\text{H}_4, \text{Br}, 3\text{CuBr}$.
 i, Real Ac. Linc. 1906 (5) 15, II,

mercuric bromide,
 $2(\text{N}_2\text{H}_4, \text{HBr}) + \text{H}_2\text{O}$.
 in H_2O , sol. in alcohol and acetone.
 cetic ether.
 ethyl ether. (Ferratini, C. A.)

zinc bromide, $2\text{N}_2\text{H}_4, \text{HBr}, \text{ZnBr}_2$.
 in H_2O ; sol. in alcohol and acetone.
 C. A. 1912. 1612.)

carbonate.
 quiescent, but only sl. sol. in H_2O .
 n alcohol. (Curtius and Jay, J.) 39. 41.)

zinc carbonate hydrazine,
 $\text{N}_2, \text{NH}_3, \text{NH}_2$, $2\text{N}_2\text{H}_4$.
 sl. in H_2O . (Ebler and Schott, J.) 79, 72.)

chlorate, $\text{N}_2\text{H}_4, \text{HClO}_3$.
 in H_2O ; sl. sol. in alcohol. Insol. CHCl_3 and benzol. (Salvadori, 1907, 37, (2) 32.)

dichlorate, $\text{N}_2\text{H}_4, 2\text{HClO}_3$.
 comp. in aq. solution. (Turrentine, m. Soc. 1915, 37. 1123.)

cupric chloride, $\text{N}_2\text{H}_4, \text{Cl}, \text{CuCl}_2 +$
 by H_2O . (Ranfaldi, Real. Ac. (5) 15, II. 95.)
 Decomp. by H_2O . (Ranfaldi.)

Hydrazine mercuric chloride, $\text{N}_2\text{H}_4, \text{Cl}, \text{HgCl}_2$.
 (Hoffmann and Marburg, A. 1899, 305. 221.)

Hydrazine dithionate, $\text{N}_2\text{H}_4, \text{H}_2\text{S}_2\text{O}_6$.
 Sol. in H_2O ; decomp. on standing in aq. solution. (Sabanejeff, Z. anorg. 1899, 20. 21.)
 $2\text{N}_2\text{H}_4, \text{H}_2\text{S}_2\text{O}_6$. Sol. in H_2O . (Sabanejeff.)

Hydrazine fluosilicate, $\text{N}_2\text{H}_4, \text{H}_2\text{SiF}_6$.
 Easily sol. in H_2O .
 Difficultly sol. in ethyl and methyl alcohol. (Ebler, J. pr. 1910, (2) 81. 552.)

Hydrazine fluotitanate, $(\text{N}_2\text{H}_4)_2, \text{H}_2\text{TiF}_6 + 2\text{H}_2\text{O}$.
 Ppt. Sol. in H_2O . (Ebler, J. p r. 1910, (2) 81. 555.)

Hydrazine monohydrobromide, $\text{N}_2\text{H}_4, \text{HBr}$.
 Very easily sol. in H_2O or hot alcohol. (Curtius and Schultz, J. pr. (2) 42. 537.)

Hydrazine dihydrobromide, $\text{N}_2\text{H}_4, 2\text{HBr}$.
 Easily sol. in H_2O . Sl. sol. in alcohol. (Curtius and Schulz, J. pr. (2) 42. 535.)

Hydrazine monohydrochloride, $\text{N}_2\text{H}_4, \text{HCl}$.
 Extremely sol. in H_2O . Sl. sol. in boiling absolute alcohol. (Curtius and Jay, J. pr. (2) 39. 38.)

Hydrazine dihydrochloride, $\text{N}_2\text{H}_4, 2\text{HCl}$.
 Easily sol. in cold H_2O ; sl. sol. in hot alcohol. (Curtius, l. c.)
 Sol. in 2.67 pts. H_2O at 23° . Sp. gr. of sat. solution at $20^\circ/4^\circ = 1.4226$. (Schiff, Z. phys. Ch. 1896, 21. 292.)

Sp. gr. of aqueous solution at 20° .

% salt	Sp. gr.
25	1.1183
20	1.0923
15	1.0675
10	1.0436
5	1.0206

(Schiff, Z. phys. Ch. 1896, 21. 292.)

Nearly insol. in hot absolute alcohol. (Curtius and Jay, J. pr. (2) 39. 37.)

Hydrazine dihydrofluoride, $\text{N}_2\text{H}_4, 2\text{HF}$.
 Easily sol. in H_2O . Nearly insol. in alcohol. (Curtius and Schulz, J. pr. (2) 42. 533.)

Hydrazine monohydroiodide, $\text{N}_2\text{H}_4, \text{HI}$.
 Easily sol. in H_2O . (Curtius and Schulz.)

Hydrazine dihydroiodide, $\text{N}_2\text{H}_4, 2\text{HI}$.
 Very deliquescent. Easily sol. in H_2O . Sl. sol. in alcohol. (Curtius and Schulz, J. pr. (2) 42. 536.)

Trihydrazine dihydroiodide, $3\text{N}_2\text{H}_4, 2\text{HI}$.

Easily sol. in H_2O and alcohol. (Curtius and Schulz, J. pr. (2) 42. 540.)

Hydrazine hydroxide, $\text{N}_2\text{H}_4, \text{H}_2\text{O}$.

Miscible with H_2O or alcohol, but not with ether, chloroform, or benzene. (Curtius and Schulz, J. pr. (2) 42. 530.)

Hydrazine hypophosphate, $\text{N}_2\text{H}_4, \text{H}_4\text{P}_2\text{O}_6$.

Sl. sol. in H_2O . 1.5 pts. dissolve in 100 pts. H_2O . (Sabanejeff, Z. anorg. 1898, 17. 490.)

Hydrazine hypophosphate ammonia,
 $\text{N}_2\text{H}_4, \text{H}_4\text{P}_2\text{O}_6, \text{NH}_3$.

Sol. in H_2O . (Sabanejeff, Z. anorg. 1899, 20. 23.)

Hydrazine mercuric iodide,
 $2\text{N}_2\text{H}_4\text{I}, \text{HgI}_2 + \text{H}_2\text{O}$.

Very sol. in H_2O , alcohol, acetone, acetic ether.

Insol. in ether. Excess of H_2O decomp. (Ferratini, Gazz. ch. it. 1912, 42. (1), 172.)

Hydrazine zinc iodide,
 $\text{ZnI}_2, 2\text{N}_2\text{H}_4, \text{HI} + \frac{1}{2}\text{H}_2\text{O}$.

Very sol. in H_2O ; sol. in alcohol and acetone. (Ferratini, C. A. 1912. 1612.)

Hydrazine nitrate, $\text{N}_2\text{H}_4, \text{HNO}_3$.

Very sol. in H_2O ; sl. sol. in abs. alcohol. (Sabanejeff, Z. anorg. 1899, 20. 24.)

Solubility in H_2O at t° .

t°	g. $\text{N}_2\text{H}_4, \text{HNO}_3$ per 100 g.	
	Sat. solution	H_2O
10	63.63	174.9
15	68.47	217.2
20	72.70	266.3
25	76.61	327.5
30	80.09	402.2
35	83.06	490.3
40	85.86	607.2
45	88.06	737.6
50	91.18	1034.
55	93.58	1458.
60	95.51	2127.

(Sommer, Z. anorg. 1914, 86. 85.)

$\text{N}_2\text{H}_4, 2\text{HNO}_3$. Very unstable.

Sol. in H_2O but solution cannot be concentrated beyond 30% without decomp.

Decomp. by abs. alcohol.

Very sol. in hydrazine hydroxide + Aq. (Sabanejeff, Z. anorg. 1898, 20. 25.)

Hydrazine nitrite, $\text{N}_2\text{H}_4, \text{HNO}_2$.

Very sol. in H_2O . [Pptd. by ether from solution in alcohol; insol. in ether.

Very hygroscopic. (Sommer, Z. an 1913, 83. 125.)

Hydrazine perchlorate, $\text{N}_2\text{H}_4, \text{HClO}_4$.

Sol. in 1.48 pts. H_2O at 30° and alcohol; crystallizes from boiling alcohol. (Salvadori, Ch. Z. 31. 680.)

+ $3\text{H}_2\text{O}$.

Solubility in H_2O at t° .

t°	% $\text{N}_2\text{H}_4, \text{HClO}_4$	Sp.
18	41.72	1.2
35	66.9	1.3

(Carlson, Dissert. 1910.)

Hydrazine diperchlorate, $\text{N}_2\text{H}_4, 2\text{H}_2\text{O}$.

Efflorescent.

Deliquescent.

102 pts. salt sol. in 100 pts. H_2O at

2.8 pts. salt sol. in 100 pts. alc

1.0 pt. " " " 100 " eth

(Turrentine, J. Am. Chem. Soc. 1123.)

Hydrazine phosphate, $\text{N}_2\text{H}_4, \text{H}_2\text{PO}_4$.

Very sol. in H_2O . (Sabanejeff, 1898, 17. 488.)

$\text{N}_2\text{H}_4, 2\text{H}_2\text{PO}_4$. Sol. in H_2O . (Sa

Hydrazine phosphite, $\text{N}_2\text{H}_4, \text{H}_2\text{PO}_3$.

Sol. in H_2O . (Sabanejeff, Z. anorg. 468.)

Hydrazine hydrogen phosphite,
 $\text{N}_2\text{H}_4, 2\text{H}_2\text{PO}_3$.

Less sol. in H_2O than the nor (Sabanejeff.)

Hydrazine selenate, $\text{N}_2\text{H}_4, \text{H}_2\text{SeO}_4$.

Sol. in H_2O with decomp. (Rimi 1907, I. 86.)

Hydrazine zinc selenate,
 $(\text{N}_2\text{H}_4)_2\text{SeO}_4, \text{ZnSeO}_4$.

Sl. sol. in H_2O but more sol. than (Rimini and Malagnini, Gazz. ch. 37. (1), 265.)

Hydrazine sulphocyanide, $\text{N}_2\text{H}_4, \text{H}$

Very deliquescent. Sol. in H_2O and (Curtius and Herdenreich, J. pr. 184 488.)

Hydrazine sulphate, $\text{N}_2\text{H}_4, \text{H}_2\text{SO}_4$.

Sol. with difficulty in cold, easily in H_2O . Insol. in alcohol. (Curtius, 1

100 pts. H_2O dissolve 8.055 pts. (Curtius and Jay, J. pr. (2) 30. 30.)

$2\text{N}_2\text{H}_4, \text{H}_2\text{SO}_4$. Very deliquescent

Insol. in alcohol. (Curtius, J. 11.)

ulphite, $(N_2H_4)_2H_2SO_3$.
f, Z. anorg. 1899, 20, 24.)

rosulphite, $2N_2H_4, H_2S_2O_3$.
with decomp.; insol. in alcohol.
Z. anorg. 1899, 20, 23.)

iosulphate, $(N_2H_4)_2H_2S_2O_3$.
t pure.) (Ferratini, Gazz. ch. 1) 138.)

ad thiosulphate,
 $2(N_2H_4)_2H_2S_2O_3 + H_2O$.
[H_2O and alcohol.
[Cl + Aq and in HNO_3 + Aq.
A. 1912. 1612.)

ver thiosulphate,
 $(N_2H_4)_2H_2S_2O_3$.
 H_2O ; sol. in NH_4OH + Aq and in
ratini.)

ulphinic acid.

zine disulphinate,
 $(N_2)_2$.
 H_2O ; sol. in acids. (Ephraim,
190.)

ulphonic acid, $N_2H_4SO_3OH$.
at 24 pts. H_2O at ord. temp. De-
mineral acids; nearly insol. in
ther organic solvents. (Traube,
1941.)

hydrazinesulphonate,
 NH_4 .
it; decomp. by acids. (Traube.)

azinesulphonate,
 $(N_2)_2Ba + 2H_2O$.
; pptd. by alcohol; decomp. by
be.)

azinesulphonate,
 $(N_2)_2Ca + H_2O$.
. Decomp. by acids. Insol. in
ube.)

irazinesulphonate, $N_2H_4SO_3K$.
r acids. (Traube.)

nesulphonate, $N_2H_4SO_3Ag$.

zinesulphonate,
 $Na + H_2O$.
r acids. (Traube.)

Strontium hydrazinesulphonate,
 $(N_2H_4SO_3)_2Sr + 2H_2O$.

Sol. in H_2O . Decomp. by acids. Insol. in
alcohol. (Traube.)

Hydriodic acid, HI.
See Iodhydric acid.

Hydrobromic acid, HBr.
See Bromhydric acid.

Hydrochloric acid, HCl.
See Chlorhydric acid.

Hydrofluorboric acid, HBF_4 .
See Fluoborhydric acid.

Hydrofluoric acid, HF.
See Fluorhydric acid.

Hydrogen, H_2 .
Sl. absorbed by H_2O .

Sol. in 150 pts. H_2O ; 1 vol. H_2O absorbs 0.018 vol. H .
Recently boiled H_2O absorbs 1.53% H . (Henry, 1893.)
100 vols. H_2O at 18° absorb 4.6 vols. H . (de Saussure,
1814.)

1 vol. H_2O absorbs 0.0193 vol. H at 760 mm.
and all temperatures between 0° and 23.6° .
(Bunsen.)

Later work does not confirm the above state-
ment.

Absorption of H by H_2O at t and 760 mm.
 β = coefficient of absorption; β_1 = "solu-
bility" (see under Oxygen).

t°	β	β_1
0	0.02153	0.02140
1	0.02134	0.02120
2	0.02115	0.02100
3	0.02097	0.02081
4	0.02079	0.02062
5	0.02061	0.02043
6	0.02044	0.02025
7	0.02027	0.02207
8	0.02010	0.01989
9	0.01994	0.01971
10	0.01978	0.01954
11	0.01962	0.01937
12	0.01947	0.01920
13	0.01932	0.01904
14	0.01918	0.01888
15	0.01903	0.01872
16	0.01889	0.01856
17	0.01876	0.01840
18	0.01863	0.01825
19	0.01850	0.01810
20	0.01837	0.01795
21	0.01825	0.01781
22	0.01813	0.01767
23	0.01802	0.01753
24	0.01791	0.01739
25	0.01780	0.01725
26	0.01770	0.01712

(Timofejeff, Z. phys. Ch. 6. 147.)

Absorption of H by H₂O at t° and 760 mm.
 β = coefficient of absorption.

t°	β	t°	β	t°	β
0	0.0203	16	0.0182	32	0.0161
1	0.0202	17	0.0180	33	0.0160
2	0.0200	18	0.0179	34	0.0159
3	0.0199	19	0.0178	35	0.0157
4	0.0198	20	0.0177	36	0.0156
5	0.0196	21	0.0175	37	0.0155
6	0.0195	22	0.0174	38	0.0154
7	0.0194	23	0.0172	39	0.0153
8	0.0192	24	0.0171	40	0.0152
9	0.0191	25	0.0170	45	0.0149
10	0.0190	26	0.0168	50	0.0146
11	0.0189	27	0.0167	60	0.0144
12	0.0187	28	0.0166	70	0.0146
13	0.0186	29	0.0164	80	0.0149
14	0.0184	30	0.0163	90	0.0155
15	0.0183	31	0.0162	100	0.0166

(Bohr and Bock, W. Ann. 44. 318.)

Absorption of hydrogen by H₂O at t° and 760 mm. pressure. β = coefficient of absorption. β_1 = "solubility" (see under Oxygen).

t°	β	β_1
0	0.02148	0.02135
1	0.02126	0.02112
2	0.02105	0.02090
3	0.02084	0.02068
4	0.02064	0.02047
5	0.02044	0.02026
6	0.02025	0.02006
7	0.02007	0.01987
8	0.01989	0.01968
9	0.01972	0.01950
10	0.01955	0.01932
11	0.01940	0.01915
12	0.01925	0.01899
13	0.01911	0.01883
14	0.01897	0.01867
15	0.01883	0.01851
16	0.01869	0.01836
17	0.01856	0.01821
18	0.01844	0.01706
19	0.01831	0.01792
20	0.01819	0.01777
21	0.01805	0.01761
22	0.01792	0.01746
23	0.01779	0.01730
24	0.01766	0.01715
25	0.01754	0.01700
26	0.01742	0.01685
27	0.01731	0.01670
28	0.01720	0.01656
29	0.01709	0.01642
30	0.01699	0.01630
31	0.01692	0.01618
32	0.01685	0.01606
33	0.01679	0.01596
34	0.01672	0.01585
35	0.01666	0.01574

Absorption of hydrogen by H₂O at t° and 760 mm. pressure.—Continued.

t°	β	β_1
36	0.01661	0.01564
37	0.01657	0.01554
38	0.01652	0.01544
39	0.01648	0.01535
40	0.01644	0.01525
41	0.01640	0.01515
42	0.01635	0.01504
43	0.01631	0.01494
44	0.01627	0.01482
45	0.01624	0.01473
46	0.01620	0.01460
47	0.01617	0.01449
48	0.01614	0.01437
49	0.01611	0.01425
50	0.01608	0.01413
52	0.01606	0.01392
54	0.01605	0.01369
56	0.01603	0.01343
58	0.01602	0.01311
60	0.01600	0.01287
62	0.01600	0.01256
64	0.01600	0.01223
66	0.01600	0.01189
68	0.01600	0.01150
70	0.01600	0.01109
72	0.01600	0.01070
74	0.01600	0.01017
76	0.01600	0.00966
78	0.01600	0.00912
80	0.01600	0.00853
82	0.01600	0.00790
84	0.01600	0.00723
86	0.01600	0.00652
88	0.01600	0.00575
90	0.01600	0.00494
92	0.01600	0.00407
94	0.01600	0.00313
96	0.01600	0.00216
98	0.01600	0.00111
100	0.01600	0.0000

(Winkler, B. 24. 99.)

Critical t, —232°. (Natanson, Z. phys. Ch. 1895, 17. 43–48.)

Coefficient of absorption for H₂O = 0.017 at 25°; 0.01905 at 20°; 0.02059 at 15°; 0.02213 at 10°; 0.02366 at 5°. (Braun, phys. Ch. 1900, 33. 734.)

Solubility in H₂O at 25° = 0.01926. (Gecken, Z. phys. Ch. 1904, 49. 267.)

Coefficient of absorption for H₂O = 0.018 at 20.11°. (Hufner, Z. phys. Ch. 1907, 13. 623.)

Solubility in H₂O at 25° = 0.01963. (Drucker and Moles, Z. phys. Ch. 1910, 14. 417.)

Coefficient of absorption for H₂O at 15° = 0.01892; at 20° = 0.01829. (Müller, Z. phys. Ch. 1912, 81. 493.)

Solubility of hydrogen in water at 25°.
Solubility calculated according to formula, for which see original article.
P=Pressure.

	S	P	S
6	0.0199	1095	0.0195
7	0.0198	1097	0.0197
0	0.0200	1244	0.0202
7	0.0193	1252	0.0196
6	0.0194	1380	0.0196
2	0.0198	1393	0.0198

(Landlay, Chem. Soc. 1912, 101. 1465.)

Absorption by H₂O at different pressures.
Hg-pressure in metres.
S=coefficient of solubility.
Table I. Volume of the absorbing liquid = 7 ccm. T = 19.5°.

	λ	P	λ
110	0.01798	3.3926	0.01789
167	0.01796	4.1405	0.01776
309	0.01799	4.6629	0.01761
133	0.01800	5.4705	0.01748
711	0.01794	5.9580	0.01725
148	0.01791	6.6507	0.01706
102	0.01793	7.4548	0.01674
208	0.01793	7.8783	0.01652
371	0.01795	8.2439	0.01632

Table II. Volume of the absorbing liquid = 7 ccm. T = 23°.

	λ	P	λ
154	0.01736	4.6220	0.01716
758	0.01739	5.1130	0.01702
416	0.01733	5.9702	0.01687
712	0.01731	7.1920	0.01649
724	0.01734	7.4493	0.01631
115	0.01732	7.8696	0.01618
229	0.01728	8.1913	0.01603

(Cassuto, Phys. Zeit. 1904, 5. 235.)

Absorption of H₂ by acids + Aq.
M = content in gram-equivalents per liter.
S = solubility. (See under Oxygen.)
Absorption of H₂ by HNO₃ + Aq.

M	S 25°
0.741	0.01851
0.753	0.01868
1.22	0.01812
1.45	0.01782
2.09	0.01739
2.96	0.01690
3.18	0.01667
3.22	0.01633
4.13	0.01611
4.23	0.01589

(Geffcken, Z. phys. Ch. 1904, 49. 267.)

Absorption of H₂ by HCl + Aq.

M	S 25°
0.426	0.01875
0.432	0.01868
1.063	0.01789
1.602	0.01732
1.802	0.01699
1.928	0.01688
2.338	0.01652
2.438	0.01627
2.836	0.01606

(Geffcken.)

Absorption of H₂ by $\frac{H_2SO_4}{2}$ + Aq.

M	S 25°
0.527	0.01869
0.562	0.01838
0.985	0.01780
1.122	0.01768
1.866	0.01642
1.905	0.01632
2.605	0.01575
3.045	0.01496
3.174	0.01456
3.962	0.01422
3.989	0.01402

(Geffcken.)

Solubility of H₂ in H₂SO₄ + Aq at 20°.

% H ₂ SO ₄	λ 20°
0	0.0208
35.82	0.00954
61.62	0.00708
95.6	0.01097

(Christoff, Z. phys. Ch. 1906, 55. 627.)

Solubility of H₂ in colloidal ferric hydroxide solution is practically the same as its solubility in pure H₂O.

Solubility of H₂ in a solution containing 18.11 g. Fe(OH)₃ per liter at 25° = 0.3085.

Solubility of H₂ in a solution containing 18.309 g. Fe (OH)₃ per liter at 25° = 0.3083.

(Geffcken, Z. phys. Ch. 1904, 49. 299.)

Absorption of H₂ by bases + Aq.

M = content in gram-equivalents per liter.
S = solubility. (See under Oxygen.)

Absorption of H₂ by KOH + Aq.

M	S 25°
0.536	0.01658
0.715	0.01539
1.059	0.01378
1.056	0.01389
1.480	0.01195

(Geffcken, Z. phys. Ch. 1904, 49. 267.)

Absorption of H ₂ by NaOH + Aq.		
M	S 25°	
0.543	0.01632	
0.571	0.01608	
0.692	0.01442	
0.974	0.01409	
1.059	0.01372	
1.137	0.01348	
1.850	0.01018	
3.400	0.00648	
3.430	0.00639	
4.687	0.00483	
(Geffcken.)		
Solubility of H ₂ in salts + Aq at 15°.		
Salt	% Salt	Coefficient of absorption
H ₂ O	0.00	0.01883
CaCl ₂	3.47	0.01619
	6.10	0.01450
	11.33	0.01138
	17.52	0.00839
	26.34	0.00519
MgSO ₄	4.94	0.01501
	10.19	0.01159
	23.76	0.00499
LiCl	3.48	0.01619
	7.34	0.01370
	14.63	0.0099
K ₂ CO ₃	2.82	0.01628
	8.83	0.01183
	16.47	0.00761
	24.13	0.00462
	41.81	0.00160
KCl	3.83	0.01667
	7.48	0.01489
	12.13	0.01279
	19.21	0.01012
	22.92	0.00892
KNO ₃	4.73	0.01683
	8.44	0.01559
	16.59	0.01311
	21.46	0.01180
NaNO ₃	5.57	0.01603
	11.16	0.01370
	19.77	0.01052
	37.43	0.00578
Na ₂ CO ₃	2.15	0.01639
	8.64	0.01385
	11.53	0.00839
Na ₂ SO ₄	4.58	0.01519
	8.42	0.0154
	16.60	0.00775

(Gordon, Z. phys. Ch. 1895, 18. 14.)

Absorption by salts + Aq.					
Salt	t,°	Sp. gr. salt + Aq. at t,°/4°	% salt	t,°	Absorption-coefficient at t,°
KCl	15°	1.1565	22.92	16.62	0.00886
		1.1294	19.21	18.52	0.0096
		1.0794	12.13	18.71	0.0124
		1.0480	7.48	19.00	0.0144
		1.0240	3.83	19.23	0.0161
KNO_3	15°	1.1460	21.46	17.81	0.0115
		1.1101	16.59	17.27	0.0129
		1.0936	14.26	16.81	0.0137
		1.0539	8.44	17.58	0.0153
		1.0295	4.73	17.82	0.0163
K_2CO_3	15°	1.4395	41.81	13.17	0.0016
		1.3112	30.99	12.77	0.0029
		1.2353	24.13	12.62	0.0047
		1.1555	16.47	12.51	0.0077
		1.0807	8.83	11.32	0.0121
		1.0405	4.53	12.29	0.0153
$NaCl$	15°	1.1817	23.84	13.48	0.0080
		1.1088	14.78	13.17	0.0093
		1.0607	11.09	13.80	0.0114
		1.0315	4.47	13.56	0.0156
$NaNO_3$	18°	1.2963	37.43	17.27	0.0066
		1.2099	27.95	17.36	0.0079
		1.1417	19.77	17.40	0.0102
		1.0765	11.16	17.65	0.0134
Na_2CO_3	15°	1.0367	5.57	17.80	0.0157
		1.1213	11.53	13.07	0.0065
		1.0835	8.02	12.08	0.0116
		1.0457	4.64	11.61	0.0142
Na_2SO_4	18°	1.0217	2.15	11.94	0.0167
		1.1008	16.69	18.41	0.0071
		1.0768	8.42	18.57	0.0122
		1.0412	4.58	18.51	0.0146
$LiCl$	15°	1.0643	14.63	12.77	0.0106
		1.0416	7.34	12.40	0.0136
		1.0192	3.48	10.47	0.0167
$MgSO_4$	15°	1.2679	23.76	18.26	0.0046
		1.1805	16.64	17.55	0.0078
		1.1064	10.19	17.30	0.0114
		1.0503	4.97	17.10	0.0147
$ZnSO_4$	18°	1.3265	26.51	18.56	0.0089
		1.1961	17.23	18.22	0.0094
		1.1304	12.73	17.95	0.0116
		1.0602	6.03	17.79	0.0149
$CaCl_2$	15°	1.2470	26.34	17.68	0.0051
		1.2160	23.33	17.84	0.0062
		1.1506	17.52	18.09	0.0082
		1.0981	11.23	18.40	0.0111
		1.0513	6.10	18.04	0.0141
$AlCl_3$	15°	1.0255	3.47	18.52	0.0157
		1.2647	32.30	17.14	0.0066
		1.1589	20.75	17.28	0.0099
		1.0914	12.46	17.30	0.0139
CuH_2O_{11}	17.5°	1.0488	6.88	17.46	0.0146
		1.2184	47.65	13.80	0.0090
		1.1286	30.06	14.19	0.0126
		1.0872	16.67	14.16	0.0167

(Steiner, W. Ann. 1894 (2), 53. 2)

Solubility in salt solutions.
C = concentration of the solution in terms normal.

α = coefficient of absorption.

Absorption of hydrogen by $\text{NH}_4\text{NO}_3 + \text{Aq}$ at 20° .

P	C	α
1.037	0.1308	0.01572
2.167	0.2765	0.01845
3.179	0.4363	0.01823
4.823	0.6933	0.01773
6.773	0.9069	0.01744
11.550	1.6308	0.01647

(Knopp, Z. phys. Ch. 1904, 40. 103.)

Absorption of hydrogen by $\text{KNO}_3 + \text{Aq}$ at 20° .

P	C	α
1.244	0.1245	0.01835
2.094	0.2114	0.01818
4.010	0.4127	0.01785
5.925	0.6225	0.01743
7.742	0.8293	0.01667
13.510	1.5436	0.01436

(Knopp.)

Absorption of hydrogen by $\text{NaNO}_3 + \text{Aq}$ at 20° .

P	C	α
1.041	0.1236	0.01839
2.192	0.2634	0.01774
4.405	0.5416	0.01694
6.702	0.8142	0.01518
12.637	1.7394	0.01300

(Knopp.)

Absorption of hydrogen by $\text{KCl} + \text{Aq}$ at 20° .

P	C	α
1.089	0.1475	0.01823
2.123	0.2907	0.01757
4.070	0.5687	0.01661
6.375	0.9127	0.01531
7.380	1.0682	0.01472
13.612	2.1222	0.01255

(Knopp.)

Absorption of H_2 by $\text{NaCl} + \text{Aq}$.
 αt° = coefficient of absorption at t° .

Percent of NaCl in solution	$\alpha 25^\circ$	$\alpha 20^\circ$	$\alpha 15^\circ$	$\alpha 10^\circ$	$\alpha 5^\circ$
5.999	0.01383	0.01528	0.01640	0.01749	0.01839
5.506	0.01395	0.01532	0.01655	0.01769	0.01861
4.496	0.01429	0.01587	0.01714	0.01817	0.01920
3.798	0.01476	0.01623	0.01760	0.01876	0.01977
1.523	0.01603	0.01754	0.01898	0.02030	0.02155
1.250	0.01621	0.01771	0.01914	0.02052	0.02180

(Braun, Z. phys. Ch. 1900, 33. 735.)

Absorption of H by $\text{BaCl}_2 + \text{Aq}$.
 αt° = coefficient of absorption at t° .

Percent of BaCl_2 in the solution	$\alpha 25^\circ$	$\alpha 20^\circ$	$\alpha 15^\circ$	$\alpha 10^\circ$	$\alpha 5^\circ$
7.002	0.01455	0.01591	0.01715	0.01833	0.01937
6.453	0.01474	0.01605	0.01734	0.01857	0.01957
3.800	0.01562	0.01700	0.01839	0.01971	0.02089
3.291	0.01570	0.01719	0.01847	0.01983	0.02110

(Braun.)

1 vol. alcohol at t° and 760 mm. absorbs V vols. H gas reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	0.06925	9	0.06799	18	0.06690
1	0.06910	10	0.06787	19	0.06679
2	0.06896	11	0.06774	20	0.06668
3	0.06881	12	0.06761	21	0.06657
4	0.06867	13	0.06749	22	0.06646
5	0.06853	14	0.06737	23	0.06636
6	0.06839	15	0.06725	24	0.06621
7	0.06826	16	0.06713
8	0.06813	17	0.06701

(Bunsen's Gasometry, p. 286.)

One vol. alcohol absorbs 0.06925—0.000148t + 0.000001t² vols. H at t° .
(Bunsen.)

Solubility in alcohol + Aq at 20° and 760 mm.

Wt. % alcohol	Vol. H_2 absorbed	Wt. % alcohol	Vol. H_2 absorbed
0	1.93	28.57	1.04
9.09	1.43	33.33	1.17
16.67	1.29	50	2.02
23.08	1.17	66.67	2.55

(Lubarsch, W. Ann. 1889, (2) 37. 525.)

Absorption of hydrogen by alcohol.

t°	Coeff. of absorption	t°	Coeff. of absorption
0	0.0676	13.4	0.0705
6.2	0.0693	18.8	0.0740

(Timofejeff.)

Solubility of H_2 in ethyl ether at t° .

t°	Solubility
0	0.1115
5	0.1150
10	0.1195
15	0.1257

(Christoff, Z. phys. Ch. 1912, 79. 459.)

Coefficient of absorption in petroleum = 0.0582 at 20° , and 0.0652 at 10° . (Griewass and Walfisz, Z. phys. Ch. 1. 70.)

Absorption of hydrogen by chloralhydrate + Aq at 20°.

P	C	a
4.911	0.310	0.01839
7.69	0.504	0.01802
14.56	1.030	0.01712
18.77	1.397	0.01653
29.50	2.530	0.01542
32.00	2.845	0.01518
38.42	3.770	0.01440
49.79	6.000	0.01353
60.12	9.120	0.01324
63.90	10.700	0.01307

(Knopp, Z. phys. Ch. 1904, 48, 103.)

Absorption of H₂ by chloralhydrate + Aq.
t° = temp. of the solution.

P = % chloralhydrate in the solution.

$\beta_{t^{\circ}}$ = coefficient of absorption at t°.

$\beta_{15^{\circ}}$ = coefficient of absorption at 15°.

t°	P	$\beta_{t^{\circ}}$	$\beta_{15^{\circ}}$
15.0	10.0	0.01740	0.01740
16.4	16.1	0.01719	0.01737
15.8	33.35	0.01475	0.01484
15.0	39.4	0.01470	0.01470
15.6	51.0	0.01300	0.01306
16.2	60.8	0.01281	0.01230
15.5	70.7	0.01282	0.01287
15.0	79.0	0.01320	0.01320
$\beta_{20^{\circ}}$			
19.4	15.5	0.01732	0.01724
17.4	28.3	0.01569	0.01540
18.7	46.56	0.01388	0.01375
16.5	52.0	0.01314	0.01280
17.0	63.0	0.01270	0.01243
17.2	66.0	0.01285	0.01260
17.9	68.0	0.01286	0.01270
18.3	78.4	0.01398	0.01380

(Müller, Z. phys. Ch. 1912, 81, 499.)

Solubility of H₂ in glycerol + Aq.

t°	% glycerol	Coefficient of absorption
14	0	0.0193
	2.29	0.0189
	5.32	0.0186
	8.57	0.0182
	10.83	0.01815
	15.31	0.01765
21	0	0.0184
	2.29	0.0181
	5.68	0.0177
	6.46	0.0176
	10.40	0.0171
	18.20	0.0160

(Henkel, in Landolt-Börnstein, Tab. 4th Ed. 602.)

Absorption of H₂ by glycerine + Aq.
t° = temp. of the solution.

P = % glycerine in the solution.

$\beta_{t^{\circ}}$ = coefficient of absorption at t°.

$\beta_{15^{\circ}}$ = coefficient of absorption at 15°.

t°	P	$\beta_{t^{\circ}}$	$\beta_{15^{\circ}}$
14.5	14.9	0.01654	0.01647
13.0	22.8	0.01532	0.01510
13.8	38.0	0.01226	0.01216
14.5	43.5	0.01117	0.01110
13.7	49.15	0.01019	0.01010
14.9	51.5	0.01028	0.01025
12.3	68.0	0.00822	0.00806
18.0	90.7	0.00853	0.00870

(Müller, Z. phys. Ch. 1912, 81, 496.)

Solubility of H₂ in glycerine + Aq at 25°.

G = % by wt. glycerine in the solvent.

S = Solubility of H₂.

P = corrected pressure at end of experiment in mm. Hg at 0°.

P	G	S
716.3	4.0	0.0186
736.1	10.5	0.0178
684.3	22.0	0.0154
709.9	49.8	0.0099
730.1	50.5	0.0097
672.2	52.6	0.0090
741.1	67.0	0.0067
708.0	80.0	0.0051
665.5	82.0	0.0051
662.3	88.0	0.0044
741.8	95.0	0.0034

(Drucker and Moles, Z. phys. 1910, 75, 417.)

Absorption of H₂ by glucose + Aq.

t° = temp. of the solution.

P = % glucose in the solution.

$\beta_{t^{\circ}}$ = coefficient of absorption at t°.

$\beta_{20^{\circ}}$ = coefficient of absorption at 20°.

t°	P	$\beta_{t^{\circ}}$	$\beta_{20^{\circ}}$
20.5	12.2	0.01595	0.01600
20.5	20.7	0.01445	0.01430
21.1	32.56	0.01243	0.01210
21.8	45.8	0.01000	0.01015
21.2	59.0	0.00775	0.00760

(Müller, Z. phys. Ch. 1912, 81, 494.)

Solubility in sugar + Aq at 15°.

% Sugar	Absorption Coefficient
16.67	0.01561
30.08	0.01284
47.65	0.01110

(Gordon, Z. phys. Ch. 1895, 12, 14.)

Absorption of H_2 by sucrose + Aq.
 = temp. of the solution in the absorption
 1.
 = % sucrose in the solution.
 ' = coefficient of absorption at t° .
 ° = coefficient of absorption at 15° .

	P	β_{t°	β_{15°
7		0.01862	0.01892
3		0.01840	$\beta_{20^\circ} =$ 0.01829
2	5.04	0.01723	0.01726
6	14.7	0.01547	0.01510
	20.26	0.01500	0.01462
7	29.86	0.01290	0.01257
8	31.74	0.01220	0.01185
3	39.65	0.01047	0.01033
6	42.94	0.00956	0.00939

(Müller.)

sorption of H_2 by propionic acid + Aq.
 at t° = coefficient of absorption at t° .

Content of ionic in solution	α_{25°	α_{20°	α_{15°	α_{10°	α_5°
910	0.01602	0.01782	0.01908	0.02029	0.02129
763	0.01638	0.01788	0.01929	0.02042	0.02120
500	0.01690	0.01829	0.01925	0.02093	0.02181
267	0.01705	0.01842	0.01983	0.02117	0.02239
373	0.01706	0.01866	0.01987	0.02120	0.02221
634	0.01722	0.01876	0.2003	0.02142	0.02245

(Braun, Z. phys. Ch. 1900, 33. 735.)

bsorption of H_2 by organic acids + Aq.
 [= content in gram-equivalents per liter.
 = solubility.

Absorption of H_2 by CH_3COOH + Aq.

M	S_{25°
0.517	0.01925
0.528	0.01923
1.160	0.01903
1.20	0.01895
1.963	0.01885
1.980	0.01882
3.178	0.01862
3.220	0.01858
4.157	0.01849

(Geffcken, Z. phys. Ch. 1904. 49. 267.)

sorption of H_2 by $CH_2ClCOOH$ + Aq.

M	S_{25°
0.527	0.01905
0.990	0.01852
1.773	0.01783

(Geffcken.)

Absorption of H by organic substances +
 Aq at t° .

V = absorbed volume reduced to 0° and
 760 mm.

α = coefficient of absorption.

Substance	Grams in 1 liter	Vol. of solution used in ccm.	t°	V ccm.	α
Glucose	174	409.94	20.28°	5.48	0.01516
Glucose	80.8	"	20.16°	6.12	0.01649
Glucose	41.45	"	20.00°	6.36	0.01759
Urea	60	"	20.17°	6.26	0.01703
Acetamide	59	"	20.11°	6.51	0.01795
Alanine	89	"	20.08°	5.57	0.01555
Glycocoll	75	"	20.16°	5.67	0.01577

(Hüfner, Z. phys. Ch. 1907, 57. 623-4.)

Solubility in organic solvents.

Solvent	Solubility at $25^\circ C$	Solu- bility at $20^\circ C$	$\frac{ds}{dt}$
Glycerine	Not measurable		
Water	0.01992	0.02000	-0.000016
Aniline	0.02849	0.03033	-0.000368
Amyl alcohol	0.03708	0.03533	+0.00035
Nitrobenzene	0.03708	0.03533	+0.00035
Carbon bisulphide	0.03753	0.03358	+0.00079
Glacial acetic	0.06330	0.06172	+0.000316
Benzene	0.07560	0.07071	+0.000978
Acetone	0.07641	0.07027	+0.001228
Amyl acetate	0.07738	0.07432	+0.00061
Xylene	0.08185	0.07834	+0.000702
Ethyl acetate	0.08516	0.07877	+0.001278
Toluene	0.08742	0.08384	+0.000716
Ethyl alcohol	0.08935	0.08620	+0.00063
(99.8%)			
Methyl alcohol	0.09449	0.09016	+0.000866
Isobutyl acetate	0.09758	0.09287	+0.000942
Chloroform	no constant value		
Carbon tetra- chloride			

(Just, Z. phys. Ch. 1901, 37. 359.)

Extended investigations have been made
 by Findlay and Shen. (Chem. Soc. 1912,
 101. 1465) on the effect of colloids on solu-
 bility of H_2 in H_2O . See original article.

Hydrogen arsenide.

See Arsenic hydride.

Hydrogen peroxide, H_2O_2 .

Miscible with H_2O . Not stable in conc.
 solution. Aqueous solution gives up its H_2O_2
 to ether. Ethereal solution is more stable
 than an aqueous solution of the same strength,
 and may be distilled without decomp. Mis-
 cible with alcohol.

Very stable in aq. solution of various
 conc. if perfectly free from impurity such as
 compds. of heavy metals, etc. (Woffenstein,
 B. 1894, 27. 3307.)

Coefficient of distribution between ether
 and H_2O determined at 3° , 7° , and 17.5°
 with varying quantities (1.7—5%) of H_2O_2 .
 (Osipoff, C. C. 1903, II. 1265.)

As sol. in ether as in H_2O . A 50% solution
 in H_2O still contained about 25% H_2O_2 after

being shaken 6 times with large quantities of fresh ether.

Insol. in petroleum ether. (Brühl, B. 1895, 28. 2855.)

+H₂O, and +2H₂O. Does not solidify at -20°. (Wolffenstein, B. 1894, 27. 3311).

Hydrogen phosphide, gaseous (Phosphine), PH₃.

Very slightly absorbed by H₂O.

Statements as to solubility in H₂O vary considerably.

(a) *Difficultly inflammable gas*—

1 vol. H₂O absorbs 0.1122 vol. PH₃. (Dyb-kowsky, J. B. 1866. 735.)

1 vol. H₂O absorbs 0.125 vol. PH₃. (H. Davy.)

(b) *Easily inflammable gas*—

1 vol. H₂O absorbs 0.018 vol. PH₃. (Gengembre, Crell. Ann. 1. 450.)

1 vol. H₂O absorbs 0.0214 vol. PH₃. (Henry.)

1 vol. H₂O absorbs 0.025 vol. PH₃. (Davy.)

1 vol. H₂O absorbs 0.125 vol. PH₃. (Dalton, Ann. Phil. 11. 7.)

1 vol. H₂O absorbs 0.255 vol. PH₃. (Raymond, Scher. J. 5. 389.)

1 vol. H₂O dissolves 0.26 vols. PH₃ at 17°. (Stock, Böttger and Lenger, B. 1909, 42. 2855.)

Sol. in conc. H₂SO₄ without immediate decomp. (Buff, Pogg. 16. 363.)

1 vol. 50% H₂SO₄ dissolves 0.05 vol. PH₃. (S. B. and L.)

Absorbed by CuSO₄+Aq and by Br. (Berthelot.)

Absorbed rapidly by Cu₂Cl₂+Aq with formation of Cu₂Cl₂, 2PH₃, and Cu₂Cl₂, 4PH₃. (Riban, C. R. 88. 581.)

1 vol. alcohol of 0.85 sp. gr. absorbs 0.5 vol.; 1 vol. ether absorbs 2 vols. (Graham.)

Sol. in volatile oils; 1 vol. oil of turpentine absorbs 3.25 vols. (Graham.)

Several varieties of blood absorb PH₃.

Hydrogen phosphide, liquid, P₂H₄.

Insol. in H₂O. Apparently sol. in alcohol and oil of turpentine, but solution is very quickly decomp. (Thénard, A. ch. (3) 145.)

Hydrogen phosphide, solid, P₄H₂.

Insol. in H₂O and alcohol. (Leverrier, A. ch. 60. 174.)

Insol. in all liquids except liquid PH₃. (Thénard, A. ch. (3) 14. 5.)

Instantly decomp. by HNO₃, or H₂SO₄+Aq. Sol. with decomp. in alcoholic solution of KOH. (Thénard.)

Somewhat sol. in liquid phosphorus. (Buck, Dissert. 1904.)

P₄H₂. Insol. in all solvents. (Stock, Böttger and Lenger, B. 1909, 42. 2851.)

P₁₂H₄. Sol. in liquid hydrogen phosphide and molten P. There are no other solvents which appreciably dissolve it. Insol. in liquid PH₃. (S. B. and L.)

Hydrogen selenide, H₂Se.

More sol. in H₂O than hydrogen sulphide (Berzelius.)

Solubility coefficient of H₂Se at t°.

t°	Sol. Coeff.
4	3.77 vols.
9.65	3.43
13.2	3.31
22.5	2.70

(Forcrand and Fonzes-Diacon, C. R. 1902, 134. 171.)

Sol. in cold COCl₂. (Besson, C. R. 1892, 122. 140.)

Hydrogen silicide.

See Silicon hydride.

Hydrogen sulphide, H₂S.

(a) *Liquid*. Dissolves S on warming, which separates on cooling.

(b) *Gas*.

1 vol. H₂O absorbs 1.08 vols. H₂S at 10°. (Henry, 1803.)

1 vol. H₂O absorbs 2.53 vols. H₂S at 15°. (de Saussure, Ann. Phil. 6. 340.)

1 vol. H₂O absorbs 3 vols. H₂S at 11°. (Gay-Lussac and Thénard.)

1 vol. H₂O absorbs 3.66 vols. H₂S at ord. temp. (Thompson.)

1 vol. H₂O absorbs 2.5 vols. H₂S at ord. temp. (Dalton.)

1 vol. H₂O absorbs 4.3706-0.083687t+0.0005213t² vols. H₂S at temperatures between 2 and 43.3°. (Bunsen and Schönfeld, A. 93. 26.)

At 0° and about 820 mm. pressure, 1 ccm. H₂O absorbs 100 ccm. H₂S, while only about 4 ccm. are absorbed at ord. pressure. (de Forcrand and Villard, C. R. 103. 1402.)

1 vol. H₂O at 760 mm. pressure and t° absorbs V vols. H₂S, reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	4.3706	14	3.3012	28	2.4357
1	4.2874	15	3.2326	29	2.3819
2	4.2053	16	3.1651	30	2.3290
3	4.1243	17	3.0986	31	2.2771
4	4.0442	18	3.0331	32	2.2262
5	3.9652	19	2.9687	33	2.1764
6	3.8872	20	2.9053	34	2.1277
7	3.8103	21	2.8430	35	2.0799
8	3.7345	22	1.7817	36	2.0332
9	3.6596	23	2.7215	37	1.9876
10	3.5858	24	2.6623	38	1.9430
11	3.5132	25	2.6091	39	1.8994
12	3.4415	26	2.5470	40	1.8569
13	3.3708	27	2.4909

(Schönfeld, A. 93. 26.)

Absorption coefficient of H_2S in H_2O at $0^\circ = 4.6796$. (Prytz and Holst, W. Ann. 1885, 54, 137.)
 1 l. H_2O dissolves 0.1004 mol. H_2S at 25° and 760 mm. (Pollitzer, Z. anorg. 1909, 64, 145.)

Absorption of hydrogen sulphide by H_2O at t° .

t°	Coefficient of absorption
0°	4.686
10°	3.520
20°	2.672

[Calc. fr. data of Fauser. (C. C. 1889, 1, 54.)]
 (Winkler, Z. phys. Ch. 1906, 55, 350.)

Absorption of hydrogen sulphide by H_2O at t° .

t°	Coefficient of absorption
0	4.621
10	3.362
20	2.554
30	2.014
40	1.642
50	1.376
60	1.176

(Winkler, Z. phys. Ch. 1906, 55, 350.)

Solubility of H_2S in H_2O .

Layer rich in H_2S

t°	Mols. H_2S per 100 mols. H_2O
0	0.4
6	0.5
17	0.8
26	1.2
29.5	1.6

Layer rich in H_2O

t°	Mols. H_2S per 100 mols. H_2O
29.4	96.3
28.5	96.9
26.9	97.3
26.3	97.5
23.8	98.1
23.3	98.1
22.9	98.2
17.2	98.8
13.7	99.1
11.4	99.25
5.3	99.5

Sheffer, Proc. K. Ak. Amsterdam, 1911, 14, 198.)

Difficultly sol. in conc. H_2SO_4 with decomp.
 Instantly decomp. by fuming HNO_3 .
 Solubility of H_2S in $\text{HI} + \text{Aq}$ at 25° and 760 mm.

HI Mol. per l.	H_2S Mol. per l.
0.00	0.1004
1.01	0.111
1.51	0.113
1.93	0.125
2.65	0.130
2.64	0.138
3.42	0.142
4.38	0.163
5.005	0.165
5.695	0.181
6.935	0.197
* (9.21)	0.267

* Not exact.

(Pollitzer, Z. anorg. 1909, 64, 145.)

Solubility in acids + Aq.

l = value of H_2S dissolved in acid + Aq as determined by titration.

l_0 = value of H_2S dissolved in H_2O as determined by titration.
 $t^\circ = 25^\circ$.

Acid	l/l_0
$\frac{1}{2}$ -N. HCl	0.975
$\frac{1}{2}$ -N. H_2SO_4	0.905

(McLauchlan, Z. phys. Ch. 1903, 44, 615.)

Less sol. in NaCl , or $\text{CaCl}_2 + \text{Aq}$ than in H_2O .

Sol. in $\text{CdCl}_2 + \text{NH}_4\text{OH} + \text{Aq}$. (Crobaugh, Z. anorg. 1894, 5, 321.)

Solubility in salts + Aq.

l = value of H_2S dissolved in salt + Aq as determined by titration.

l_0 = value of H_2S dissolved in H_2O as determined by titration.
 $t^\circ = 25^\circ$.

Salt + Aq.	l/l_0	Salt + Aq.	l/l_0
$\frac{1}{2}$ -N. Na_2SO_4	0.73	$\frac{1}{4}$ -N. Na_2SO_4	0.855
$\frac{1}{2}$ -N. K_2SO_4	0.78	$\frac{1}{4}$ -N. K_2SO_4	0.890
$\frac{1}{2}$ -N. $(\text{NH}_4)_2\text{SO}_4$	0.82	$\frac{1}{4}$ -N. $(\text{NH}_4)_2\text{SO}_4$	0.91
N. NaCl	0.847	$\frac{1}{2}$ -N. NaCl	0.930
N. KCl	0.853	N. NaBr	0.935
N. NH_4Cl	0.960	N. KBr	0.945
N. NaNO_3	0.893	N. NH_4Br	1.00
N. KNO_3	0.913	N. KI	0.98
N. NH_4NO_3	0.990		

(McLauchlan, Z. phys. Ch. 1903, 44, 615.)

Solubility of H ₂ S in NaSH + Aq. (g. mol. H ₂ S dissolved in 1 l.)			
t°	0.05 g. mol. NaSH per l.	0.1 g. mol. NaSH per l.	0.2 g. mol. NaSH per l.
15	...	0.132	0.129
25	...	0.104	0.1035
35	0.082	0.082	...
45	0.064

(Goldschmidt and Larsen, Z. phys. Ch. 1910, 71. 449.)

At 18° and ord. pressure, 100 vols. alcohol of 0.84 sp. gr. absorb 606 mols. H₂S. (de Saussure, 1814.)

1 vol. alcohol absorbs 17.891-0.65598t + 0.00661t² vols. H₂S between 0° and 22°. (Carius.)

1 vol. alcohol at t° and 760 mm. absorbs V vols. H₂S reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	17.891	9	12.523	18	8.225
1	17.242	10	11.992	19	7.814
2	16.606	11	11.475	20	7.415
3	15.983	12	10.971	21	7.030
4	15.373	13	10.480	22	6.659
5	14.776	14	10.003	23	6.300
6	14.193	15	9.539	24	5.955
7	13.623	16	9.088
8	13.066	17	8.650

(Carius, A. 94. 140.)

Solubility in alcohol + Aq at 25°.

Molecules of C ₂ H ₅ OH in 100 molecules C ₂ H ₅ OH + H ₂ O	Molecules H ₂ O in 100 molecules C ₂ H ₅ OH + H ₂ O	l/lo
0.00	100	1.00
1.60	98 (?)	0.96
5.18	94.82	0.933
9.25	90.75	0.91
23.60	76.40	1.28
47.75	52.25	1.95
* (100	0	2.16)

* Carius.

(McLauchlan.)

Sol. in methyl acetate (Marchand), ether (Higgins).

Insol. in caoutchin.
Sol. in glycerine in less amount than in H₂O. If a certain vol. of H₂O dissolves 100 pts. H₂S, the same vol. of glycerine (1 pt. glycerine + 1pt. H₂O) dissolves only 60 pts. H₂S, but the solution is very stable. After standing a year there is no appreciable decomp. (Lapage, J. Pharm. (4) 5. 256.)

According to Lindo (C. N. 57. 173), the solution in glycerine is no more stable than that in H₂O.

Sol. in CS₂.

Solubility in organic substances + Aq
l = value of H₂S dissolved in organic
stance + Aq as determined by titration.
lo = value of H₂S dissolved in H₂O
terminated by titration.
t° = 25°.

Solution	l/lo
N-NH ₄ C ₂ H ₃ O ₃	1.09
N-C ₄ H ₉ O ₆	0.944
3-N-C ₄ H ₉ O ₆	0.858
N-(NH ₂) ₂ CO	1.02
pure C ₂ H ₅ (OH) ₂	0.863

(McLauchlan, Z. phys. Ch. 1903, 44. 1)

Solubility in acetic acid + Aq at 25°

Molecules of CH ₃ COOH in 100 molecules CH ₃ COOH + H ₂ O	Molecules of H ₂ O in 100 molecules CH ₃ COOH + H ₂ O	l/lo
0	100	1.0
8.85	91.15	0.9
16.7	83.30	0.8
21.0	79	1.0
35.5	64.5	1.0
53.5	40.5	1.0
55.7	...	1.0
67.8	32.2	1.0
81.0	19.0	1.0
98.58	1.42	3.0

(McLauchlan.)

Hydrogen persulphide, H₂S₂ or H₂S₈.

Decomp. by contact with H₂O, in v is apparently insol. Sol. in ether with quent decomp. Sol. in CS₂. (Thén ch. 48. 79.)

H₂S₂. Quickly decomp. by ether, ether, ethyl, or amyl alcohol. H₂S action.

Conc. HCl, or HC₂H₃O₂ + Aq have tion. Sol. in a solution of S in CS₂, liquid hydrocarbons.

Chloroform dissolves without decomp (Sabatier, C. R. 100. 1346, 1585.)

Alkalies, and K₂S + Aq decomp. in Decomp. by H₂O, dil. and conc. HC H₂SO₄, alkali and alcohol. Sol. in containing HCl but soon decomp. solution. Miscible in all proportio without decomp. with benzene, ether a (Bloch, B. 1908, 41. 1977.)

Formula is H₂S₈. (Rebs, A. 244 + 7H₂O. Easily decomp. by heat Forcrand and Villard, C. R. 103. 140

Hydrogen trisulphide, H₂S₃.

Decomp. by H₂O, dil. and conc. HC H₂SO₄, alkali and alcohol. Somewhat alcohol containing HCl, but slowly d in this solution. Miscible with ethe zene and CS₂, and these solutions are re stable. (Bloch, B. 1908, 41. 1974.)

telluride, H_2Te .
in H_2O . Decomp. in the air.
anorg. 1900, 25. 313.)

phuric acid, H_2S .
rogen sulphide.

phurous acid, H_2SO_2 .
osulphurous acid.

amic acid.

ydroxylamate, $\text{Ca}(\text{ONH}_2)_2$.
plosive; decomp. by H_2O . (Ebler
, J. pr. 1908, (2) 78. 323.)

xylamate $\text{Zn}(\text{ONH}_2)_2$.
. by H_2O . (Ebler and Schott.)

xylamate, hydroxylamine,
 $(\text{NO})_2$, $3\text{NH}_2\text{O}$.
stable.
abs. alcohol. (Ebler and Schott.)

amine, $\text{NH}_2\text{O} = \text{NH}_2(\text{OH})$.
only in solution.
lcohol. (Lossen, J. pr. 96. 462.)
l in free state by de Bruyn.
liquescent, and sol. in H_2O and
sl. sol. or insol. in CHCl_3 , C_6H_6 ,
thyl acetate.
alcohol at 5° dissolves 35%; ethyl
 15° , 15%; boiling dry ether, 1.2%;
yl acetate, 1.6%. (de Bruyn, R.
)

mine arsenate, $\text{AsO}_4\text{H}_2(\text{NH}_2\text{O})_2$.
n cold H_2O ; sol. in hot H_2O from
an be cryst. (Hofmann, A. 1899,

mine azoimide.
mide, hydroxylamine.

mine bromide, NH_2OH , HBr .
in H_2O ; insol. in ether by which
from solution in alcohol. (Adams,
1902, 28. 205.)
 I , HBr . Easily sol. in H_2O ; insol.
and ligroin. Sl. sol. in alcohol.

mine mercuric bromide hydroxyla-
 $2\text{NH}_2\text{OH}$, 2HBr , HgBr_2 , $2\text{NH}_2\text{OH}$.
. by H_2O and methyl alcohol.
decomp. by alkalies. (Adams.)

nine calcium, HO.Ca.ONH_2 .
decomp. by H_2O at ordinary
Hofmann, Z. anorg. 1898, 16. 464.)

Hydroxylamine chloride, basic, $\text{NH}_2(\text{OH})\text{Cl}$,
 NH_2OH .

Sol. in H_2O . Alcohol precipitates from
aqueous solution. Insol. in ether. (Lossen.)
 $2\text{NH}_2(\text{OH})\text{Cl}$, NH_2OH . Deliquescent;
very sol. in H_2O , less in alcohol, and insol. in
ether. (Lossen.)

Hydroxylamine chloride, $\text{NH}_2(\text{OH})\text{Cl}$.

Not deliquescent. Very sol. in H_2O and
hot ordinary alcohol. Sl. sol. in absolute al-
cohol. Insol. in ether. (Lossen.)

Sol. in 1.2 pts. H_2O at 17° . (Schiff, Z. phys.
Ch. 1896, 21. 290.)

Sp. gr. of aqueous solution at 17° .

% salt	sp. gr.
40	1.1852
28	1.1260
20	1.0888
14	1.0616
10	1.0437
7	1.0303
5	1.0214
3.5	1.0147

(Schiff, Z. phys. Ch. 1896, 21. 290.)

100 pts. absolute methyl alcohol dissolve
16.4 pts. at 19.75° ; 100 pts. absolute ethyl
alcohol dissolve 4.43 pts. at 19.75° . (de
Bruyn, Z. phys. Ch. 10. 783.)

Somewhat sol. in alcohol. (Adams, Am.
Ch. J. 1902, 28. 204.)

Hydroxylamine mercuric chloride, NH_2OH ,
 HCl , HgCl_2 .

Very sol. in H_2O and alcohol. Less sol. in
ether. (Adams, Am. Ch. J. 1902, 28. 213.)

$5(\text{NH}_2\text{OH})_2$, HCl , 2HgCl_2 . Sol. in cold
 H_2O , alcohol and ether. More easily sol. in
methyl alcohol. Sol. in HCl . The slightest
trace of alkali causes decomp. (Adams.)

Hexahydroxylamine cobaltic bromide,
 $[\text{Co}(\text{NH}_2\text{OH})_6]\text{Br}_3$.

(Werner, B. 1905, 38. 897.)

Hexahydroxylamine cobaltic chloride,
 $[\text{Co}(\text{NH}_2\text{OH})_6]\text{Cl}_3$.

Very stable toward HCl . (Werner, B.
1905, 38. 895.)

Hexahydroxylamine cobaltic nitrate,
 $[\text{Co}(\text{NH}_2\text{OH})_6](\text{NO}_3)_3$.

(Werner.)

Hexahydroxylamine cobaltic sulphate,
 $[\text{Co}(\text{NH}_2\text{OH})_6]_2(\text{SO}_4)_3 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O . (Werner.)

Hydroxylamine columbate, $\text{CbO}_6\text{N}_6\text{H}_{12}$.

Explosive. Sl. sol. in H_2O . (Hofmann, Z.
anorg. 1898, 16. 473.)

Hydroxylamine dithionate, $(\text{NH}_2\text{OH})_2, \text{H}_2\text{S}_2\text{O}_6$.

Sol. in H_2O ; decomp. on heating the aq. solution. (Sabanejeff, Z. anorg. 1898, 17. 485.)

Hydroxylamine fluosilicate, $(\text{NH}_2\text{O})_2, \text{H}_2\text{SiF}_6 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O . Nearly insol. in methyl and abs. ethyl alcohol. (Ebler, J. pr. 1908, (2), 78. 338.)

Hydroxylamine fluotitanate, $(\text{NH}_2\text{O})_2, \text{H}_2\text{TiF}_6$.

Sol. in H_2O . Sl. sol. in methyl alcohol. (Ebler, J. pr. 1908, (2) 78. 340.)

Hydroxylamine hypophosphite, $(\text{NH}_2\text{OH})\text{H}_2\text{PO}_2$.

Very sol. in H_2O . (Sabanejeff, Z. anorg. 1898, 17. 483.)

Sol. in H_2O and absolute alcohol. Insol. in ether. (Hofmann and Kohlschütter, Z. anorg. 1898, 16. 469.)

Hydroxylamine potassium hypophosphite, $(\text{H}_2\text{PO}_2)_2(\text{NH}_2\text{O})_2\text{K}_2$.

Easily sol. in H_2O , decomp. on heating; sol. in hot abs. alcohol. (Hofmann and Kohlschütter, Z. anorg. 1898, 16. 468.)

Hydroxylamine hypophosphate, $(\text{NH}_2\text{OH})_2\text{H}_2\text{P}_2\text{O}_6$.

Easily sol. in H_2O . (Sabanejeff, Z. anorg. 1898, 17. 489.)

Hydroxylamine iodide, $\text{NH}_2\text{OH}, \text{HI}$.

Hygroscopic; sol. in methyl alcohol. Very explosive. (Wolffenstein and Groll, B. 1901, 34. 2419.)

Dihydroxylamine iodide, $(\text{NH}_2\text{OH})_2, \text{HI}$.

Deliquescent. More sol. in H_2O , methyl and ethyl alcohol than the tri compound. Is decomp. when recryst. from these solvents. Insol. in ether. (Dunstan, Chem. Soc. 1896, 69. 841.)

Trihydroxylamine iodide, $(\text{NH}_2\text{OH})_3, \text{HI}$.

Deliquescent in moist air. Sol. in H_2O , methyl and ethyl alcohol. Insol. in ether. (Dunstan.)

Hydroxylamine nitrate, $\text{NH}_2(\text{OH})\text{NO}_3$.

Very sol. in H_2O and absolute alcohol. (Lossen.)

Hydroxylamine orthophosphate, $(\text{NH}_2\text{OH})_3\text{PO}_4$.

Sl. sol. in cold H_2O . (Lossen.)

Only sl. sol. in H_2O . (Hofmann, A. 1899, 307. 330.)

Moderately sol. in H_2O . (Ross, Chem. Soc. 1906, 90, (2) 19.)

Solubility in H_2O .

1 g. of aqueous solution contain	
0°	10°
0.012	0.015 g. hydroxylamine
20°	30°
0.019	0.027 g. hydroxylamine
40°	50°
0.040	0.055 g. hydroxylamine
60°	70°
0.077	0.102 g. hydroxylamine
80°	90°
0.133	0.168 g. hydroxylamine

(Adams, Am. Ch. J. 1902, 28.

$(\text{NH}_2\text{OH})\text{H}_2\text{PO}_4$. Hygroscopic. tion is decomp. on heating. (Sat 1897, 30. 287.)

Hydroxylamine phosphite, $(\text{NH}_2\text{O})_2$.

Sol. in H_2O and absolute alcohol. (mann and Kohlschütter, Z. anorg 467.)

$(\text{NH}_2\text{OH})\text{H}_2\text{PO}_2$. Sol. in H_2O , alcohol. (Sabanejeff, Chem. Soc. (2), 14.)

Hydroxylamine phosphite ammoni, $(\text{NH}_2\text{OH})\text{H}_2\text{PO}_2, \text{NH}_3$.

Sol. in H_2O . (Sabanejeff, Chem 78, (2) 14.)

Hydroxylamine sodium, NaONH_2 .

Very hygroscopic. (de Bruyn 1892, 11. 18.)

Hydroxylamine sulphate, $(\text{NH}_2\text{OH})_2$.

Easily sol. in H_2O . Precipitates concentrated aqueous solution by alcohol. (sen.)

Sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. In alcohol and ether. (Preibisch, J. pr. 7. 480.)

Not deliquescent. Sol. in $\frac{3}{4}$ of H_2O at 20°. (Divers and Haga, 1896, 69. 1665.)

1 g. of aqueous solution contain		
—8°	0°	+10°
0.307	0.329	0.366 g. hydroxylamin
20°	30°	40°
0.413	0.441	0.482 g. hydroxylamin
50°	60°	90°
0.522	0.560	0.685 g. hydroxylamin

(Adams, Am. Ch. J. 1902, 28.

Dry hydroxylamine sulphate is abs. and almost insol. in 95% (Adams.)

For double salts, see under sulph. $\text{NH}_2\text{OH}, \text{H}_2\text{SO}_4$. Deliquescent in H_2O . (Divers, Chem. Soc. 1895.

Hydroxylamine tungstate, $4\text{NH}_2\text{OH}, \text{H}_2\text{WO}_6, 3\text{H}_2\text{O}$.

Moderately sol. in H_2O . (Allenschalk, Am. Ch. J. 1902, 27. 338.)

amine uranate, $\text{UO}_4(\text{NH}_3\text{O})_2 + \text{H}_2\text{O}$.
by heat. (Hofmann, Z. anorg. 1.)

amine uranate ammonia, $\text{H}_2\text{O})_2, 2\text{NH}_3$.
by H_2O . (Hofmann, Z. anorg. 1.)

amine metavanadate, $\text{VO}_4\text{N}_3\text{H}_{10}$.
by moisture. (Hofmann, Z. 16. 472.)

amine metavanadate ammonia, $(\text{NH}_3\text{O})_2, 2\text{NH}_3$.
decomp. by H_2O and HCl . (Hofmann, 1898, 16. 471.)

amine monosulphonic acid, $\text{I}(\text{SO}_3\text{H})$.
"hydric acid" of Fremy.
"hydroxylamic acid" of Claus.
 I_2O . Slowly decomp. on boiling. (A. 241. 161.)

amine hydroxylamine monosulphonate, $\text{IN}, \text{SO}_3\text{NH}_4$.
eff, Z. anorg. 1898, 17. 491.)

amine — — —, $(\text{HONHSO}_3)_2\text{Ba} +$
sol. in H_2O . (Divers and Haga, 55. 760.)

amine — — —, $\text{Ba}(\text{HONSO}_3)_2\text{Ba} +$
insol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$. (Divers and Haga, Chem. Soc. 55. 760.)

amine — — —, $\text{HONH}(\text{SO}_3\text{K})$.
"amine sulphhydroxylamate" of Claus.
"amine sulphazotate" of Fremy.
sol. in H_2O . Easily sol. in hot H_2O .
decomp. Insol. in alcohol. (Raschig, 55. 760.)
(Divers and Haga, Chem. Soc. 55. 760.)

amine disulphonic acid, $(\text{SO}_3\text{H})_2$.
"hydroazotic acid" of Claus.
"azotic acid" of Fremy.
known in free state. (Raschig, A. 241. 161.)

amine hydroxylamine disulphonate, $(\text{S}_2\text{O}_7)_2 + 4\text{H}_2\text{O}$ and $+ 8\text{H}_2\text{O}$.
sol. in H_2O . Sol. in $\text{NH}_4\text{Cl} +$ water. (Divers and Haga, Chem. Soc. 1894, 65. 559.)

amine potassium — — —.
 $(\text{NS}_2\text{O}_7)_2 + 9\text{H}_2\text{O}$.
 $\text{O}_7 + \text{H}_2\text{O}$.
 $3\text{K}_2\text{H}(\text{NS}_2\text{O}_7)_2 + \text{H}_2\text{O}$.

$\text{Ba}_2\text{K}_2(\text{NS}_2\text{O}_7)_7 + 14\text{H}_2\text{O}$.
 $3\text{Ba}(\text{OH})_2, \text{Ba}_2\text{K}_2(\text{NS}_2\text{O}_7)_7$.
Above salts are all ppts. (Divers, Chem. Soc. 1894, 65. 561.)

Barium sodium hydroxylamine disulphonate, $\text{Ba}_2\text{Na}_2(\text{NS}_2\text{O}_7)_8 + 7\text{H}_2\text{O}$.
 $\text{Ba}_2\text{Na}_2(\text{NS}_2\text{O}_7)_7 + 7\text{H}_2\text{O}$.
 $\text{Ba}_{12}\text{Na}_{12}(\text{NS}_2\text{O}_7)_{17} + 24\text{H}_2\text{O}$.
Above salts are ppts. (Divers.)

Potassium — — —, $\text{HON}(\text{SO}_3\text{K})_2 + 2\text{H}_2\text{O}$.
"Potassium disulphhydroxyazotate" of Claus (A. 158. 75). Insol. in cold H_2O .
Very unstable. Very difficultly sol. in H_2O , more easily in dil. $\text{KOH} + \text{Aq}$. (Raschig, A. 241. 161.)
 $\text{HON}(\text{SO}_3\text{K})_2, \text{KON}(\text{SO}_3\text{K})_2 + \text{H}_2\text{O}$. True composition of potassium sulphazotate of Fremy. (Divers and Haga, Chem. Soc. 1900, 77. 432.)

Potassium sodium — — —.
 $3\text{K}_2\text{NS}_2\text{O}_7, 2\text{Na}_2\text{NS}_2\text{O}_7 + 2\text{H}_2\text{O}$. Sol. in H_2O .
 $6\text{K}_2\text{NS}_2\text{O}_7, \text{Na}_2\text{NS}_2\text{O}_7, \text{H}_2\text{NS}_2\text{O}_7 + 20\text{H}_2\text{O}$. Sol. in H_2O .
 $\text{K}_2\text{Na}_{16}\text{H}_2(\text{NS}_2\text{O}_7)_7 + 5\text{H}_2\text{O}$. Less sol. in H_2O than the others.
 $\text{K}_{12}\text{Na}_4\text{H}_4(\text{NS}_2\text{O}_7)_8 + 9\text{H}_2\text{O}$. Sol. in H_2O .
 $\text{KNa}_4\text{H}(\text{NS}_2\text{O}_7)_2 + \text{H}_2\text{O}$. Readily sol. in H_2O .
 $\text{K}_2\text{NaH}_2(\text{NS}_2\text{O}_7)_3 + 2\text{H}_2\text{O}$. Moderately sol. in H_2O .
 $\text{KNaHNS}_2\text{O}_7 + 3\text{H}_2\text{O}$. Sol. in H_2O .
(Divers, Chem. Soc. 1894, 65. 552.)

Potassium strontium — — —,
 $(\text{HO}, \text{Sr})_2\text{NS}_2\text{O}_7, 8(\text{SrKNS}_2\text{O}_7, 2\text{H}_2\text{O})$.
Ppt. (Divers.)

Potassium hydroxylamine disulphonate nitrite, $\text{HON}(\text{SO}_3\text{K})_2, \text{KNO}_2$.
Very sl. in H_2O . (Divers and Haga, Chem. Soc. 1900, 77. 433.)
 $\text{K}_2\text{H}(\text{NS}_2\text{O}_7)_2, 3\text{KNO}_2 + \text{H}_2\text{O}$. Decomp. by H_2O . (Divers and Haga.)
 $2\text{KON}(\text{SO}_3\text{K})_2, \text{KNO}_2 + 4.4\text{H}_2\text{O}$. Very sol. in H_2O which decomp. it into its constituent salts. (Divers and Haga.)
 $+ 6\text{H}_2\text{O}$. Very sol. in H_2O which decomp. it into its constituent salts. (Divers and Haga.)
 $2\text{K}_2\text{H}(\text{NS}_2\text{O}_7)_2, 7\text{KNO}_2 + 3\text{H}_2\text{O}$. Decomp. by H_2O . (Divers and Haga.)
 $3\text{K}_2\text{H}(\text{NS}_2\text{O}_7)_2, 7\text{KNO}_2$. Decomp. by H_2O . (Divers and Haga.)

Potassium hydroxylamine disulphonate sodium chloride, $5\text{K}_2\text{HNS}_2\text{O}_7, 8\text{NaCl} + 3\text{H}_2\text{O}$.
Decomp. by H_2O . (Divers, Chem. Soc. 1894, 65. 551.)

Sodium hydroxylamine disulphonate,
 $\text{HON}(\text{SO}_3\text{Na})_2$.

Sol. in somewhat more than its own wt. of H_2O at 14° . (Divers, Chem. Soc. 1894, 65. 546.)

$\text{Na}_2\text{HNS}_2\text{O}_7$, $2\text{Na}_2\text{NS}_2\text{O}_7 + 3\text{H}_2\text{O}$. Sol. in less than 1.5 pts. H_2O at 14° . (Divers.)

Hydroxylamine' isomonosulphonic acid,
 NH_2 , O, SO_3 , OH.

Very hygroscopic. Sol. in water; sol. in alcohol. (Sommer, B. 1914, 47. 1226.)

[Compare Raschig, A. 1887, 241. 161.]

Hydroxylamine isodisulphonic acid.

Ammonium hydroxylamine isodisulphonate,
 $(\text{SO}_3\text{NH}_4)\text{ONH}(\text{SO}_3\text{NH}_4)$.

3 pts. are sol. in 2 pts. H_2O at 18° . Apt to form supersat. solutions. (Haga, Chem. Soc. 1906, 89. 246.)

Dipotassium — — — — —, $\text{K}_2\text{HS}_2\text{O}_7\text{N}$.

Only sl. sol. in cold H_2O . Easily sol. in boiling H_2O . Decomp. by hot dil. HCl. (Raschig, B. 1906, 39. 246.)

6.44 pts. are sol. in 100 pts. H_2O at 16.4° .
 7.18 " " " " 100 " H_2O " 17.8° .
 8.05 " " " " 100 " H_2O " 20° .

(Haga, Chem. Soc. 1906, 39. 243.)

Tripotassium — — — — —, $(\text{SO}_3\text{K})\text{ONK}(\text{SO}_3\text{K}) + 2\text{H}_2\text{O}$.

Very sol. in H_2O ; ppt. by alcohol. (Haga.)

Disodium — — — — —, $(\text{SO}_3\text{Na})\text{ONH}(\text{SO}_3\text{Na})$.

Very sol. in H_2O ; insol. in alcohol by which it is ppt. from aqueous solution. (Haga.)

Trisodium — — — — —, $(\text{SO}_3\text{Na})\text{ONNa}(\text{SO}_3\text{Na}) + 2\text{H}_2\text{O}$.

Sol. in H_2O ; ppt. by alcohol. (Haga.)
 $+ 3\text{H}_2\text{O}$. Sol. in 1.3 pts. H_2O at 20° . Less sol. in $\text{NaOH} + \text{Aq}$. (Divers, Chem. Soc. 1894, 65. 546.)

Hydroxylamine trisulphonic acid.

Ammonium hydroxylamine trisulphonate,
 $2(\text{SO}_3\text{NH}_4)\text{ON}(\text{SO}_3\text{NH}_4)_2 + 3\text{H}_2\text{O}$.

Sol. in 0.61 pts. H_2O at 16° . (Haga, Chem. Soc. 1904, 85. 84.)

Potassium — — — — —, $2(\text{SO}_3\text{K})\text{ON}(\text{SO}_3\text{K})_2 + 3\text{H}_2\text{O}$.

1 pt. is sol. in 25.37 pts. H_2O at 18° . (Haga.)

Sodium — — — — —, $(\text{SO}_3\text{Na})\text{ON}(\text{SO}_3\text{Na})_2 + 2\text{H}_2\text{O}$.

Sol. in 2.84 pts. H_2O at 21.5° .

Dihydroxylamine sulphonie aci
 $(\text{HO})_2\text{N}(\text{SO}_3\text{H})$.

"Sulphazinous acid" of Fremy.
 Known only in its salts. (Raschig 161.)

Potassium dihydroxylamine su
 $(\text{HO})_2\text{NSO}_3\text{K}$.

Not obtained in pure state; forms $\text{KO}^-\text{NSO}_3\text{K}$, which is quite sol. in H_2O corresponds to "sulfazite de pot" Fremy (A. ch. (3) 15. 421).

Sol. in H_2O ; insol. in alcohol (Fremy.)

Hydroxylodoplatindiamine sulp
 $(\text{OH})\text{IPt}(\text{NH}_2)_2\text{SO}_4 + \text{H}_2\text{O}$.

Very sl. sol., even in boiling H_2O (Gren, Sv. V. A. F. 47. 312.)

Hydroxylonitratoplatindiamine nit
 $\text{OH} \text{Pt} \begin{matrix} \text{N}_2\text{H}_4\text{NO}_2 \\ \text{NO}_2 \end{matrix} \text{N}_2\text{H}_4\text{NO}_2$.

Sl. sol. in cold, more easily in H_2O containing HNO_3

— **pyrophosphate,**

$\left[\begin{matrix} \text{OH} & \text{N}_2\text{H}_4 \\ \text{NO}_2 & \text{N}_2\text{H}_4 \end{matrix} \text{Pt} \right]_2 \text{P}_2\text{O}_7 + \text{H}_2\text{O}$.

Very sl. sol. in H_2O . (Cleve.)

Hydroxyloplatinamine hydroxid
 $(\text{OH})_2\text{Pt}(\text{NH}_2\text{OH})_2$.

Insol. in H_2O . Easily sol. in dil. HCl , $\text{H}_2\text{O}_2 + \text{Aq}$. Not decomp. by $\text{KOH} + \text{Aq}$. (Gerhardt, Compt. C. 490.)

Hydroxyloplatinamine nitrate,
 $(\text{OH})_2\text{Pt}(\text{NH}_2\text{NO}_2)_2 + 2\text{H}_2\text{O}$.

Sl. sol. in cold, easily in hot H_2O ; attacked by cold $\text{HCl} + \text{Aq}$. (Cleve.)

— **oxalate,** $(\text{OH})_2\text{Pt}(\text{NH}_2)_2\text{C}_2\text{O}_4$.
 Sol. in hot H_2O .

— **sulphate,** $(\text{OH})_2\text{Pt}(\text{NH}_2)_2\text{SO}_4$.
 Difficultly sol. in H_2O . (Cleve.)

Hydroxyloplatindiamine bromid
 $(\text{OH})_2\text{Pt}(\text{NH}_2)_2\text{Br}_2$.

Sl. sol., even in boiling H_2O . (Gren, Sv. V. A. F. 47. 320.)

— **chloride,** $(\text{OH})_2\text{Pt}(\text{NH}_2)_2\text{Cl}_2$.
 Sol. in 206 pts. cold, and 49 pts. H_2O . (Carlgren, Sv. V. A. F. 47.)

— **chromate,** $(\text{OH})_2\text{Pt}(\text{NH}_2)_2\text{CrO}_4$.
 Very sl. sol. in cold or hot H_2O . (Gren, Sv. V. A. F. 47. 319.)

Calcium hypobromite bromide.

Deliquescent, and sol. in H_2O with partial decomp. (Berzelius.)

Potassium hypobromite, $KBrO$.

Known only in solution.

Sodium hypobromite.

Known only in solution.

Strontium hypobromite.

Known only in solution.

Hypochlorous acid, $HClO$.

Miscible with H_2O . Decomposes at 0° in the dark, more rapidly at higher temp. or in light. The stronger the solution the more rapid the decomposition. Moderately strong acid may be distilled without any considerable decomp., a stronger acid distilling over at first, and afterwards an acid weaker than the original acid. Very conc. or very dil. acids decomp. by distillation.

Ammonium hypochlorite.

Known only in aqueous solution, which decomposes at once.

Barium hypochlorite.

Known only in solution.

Calcium hypochlorite, $Ca(OCl)_2 + 4H_2O$.

Deliquescent, and sol. in H_2O . (Kinzgett, Chem. Soc. (2) 13. 404.)

Calcium hypochlorite chloride, etc. (bleaching powder), $Ca(OCl)_2$, $CaCl_2$, $Ca(OH)_2 + H_2O$.

Not deliquescent. Sol. in H_2O . Alcohol does not dissolve out $CaCl_2$. Sol. in 20 pts. H_2O with a slight residue.

Correct formula is $CaOCl_2$ (Lunge and Schäppi; Kraut, A. 214. 354), $Ca \begin{smallmatrix} OCl \\ OH \end{smallmatrix}$ (Stahlschmidt, B. 8. 869), $CaOCl$, Cl (Odling).

$CaCl_2$ is dissolved out by alcohol. Formula = $2Ca \begin{smallmatrix} OH \\ OCl \end{smallmatrix} CaCl + 2H_2O$. (Dreyfuss, Bull. Soc. (2) 41. 600.)

Didymium hypochlorite, $Di(OCl)_2$.

Difficultly sol. in H_2O . Easily sol. in acids. (Frerichs and Smith, A. 191. 348.)

Lanthanum hypochlorite, $La(OCl)_3$.

Easily sol. in H_2O . (Frerichs and Smith.)

Lithium hypochlorite, $LiClO$.

Known only in solution. (Kraut, A. 1882, 214. 356.)

Magnesium hypochlorite.

Known only in solution.

Potassium hypochlorite, $KClO$.

Known only in solution.

Silver hypochlorite, $AgClO$.

Very sol. in H_2O , and decomp. very (Stas, Acad. R. de Belg. 35. 103.)

Sodium hypochlorite, $NaClO$.

Known only in solution.

Hypoiodic acid, I_2O_4 .

See Iodine tetroxide.

Hypoiodous acid, HOI .

Known only in solution which decaying. (Taylor, C. N. 1897, 76. 1)

Calcium hypoiodite iodide, $Ca(OI)_2$.

Not very unstable. (Lunge and S. 15. 1883.)

Hyponitric acid, N_2O_4 .

See Nitrogen tetroxide.

Hyponitrous acid, HNO , or better

Known only in aqueous solution. is quite stable. (van der Plaats, B. 1)

Very deliquescent; sol. in H_2O and sol. in ether, chloroform, benzene; petroleum ether. (Hantzsch and Kaufmann, A. 1896, 292. 323.)

Ammonium hyponitrite, $(NH_4)_2N_2O_2$.

Sol. in H_2O and in alcohol. (Jacks 1893, 68. 266.)

Ammonium hydrogen hyponitrite, NH

Easily sol. in H_2O . The solid decomposes at ord. temp. into ammonia and N_2O . (Hantzsch and Kaufmann 1896, 292. 328.)

Barium hyponitrite, BaN_2O_2 .

Nearly insol. in, but gradually decomposed by H_2O . Sol. in conc. acids with evolution of N_2O , but sol. in dil. HCl , H_2SO_4 , + Aq decomp. (Zorn, B. 15. 1007.)

+ $4H_2O$. Sl. sol. in H_2O ; insol. in ether. (Kirschner, Z. anorg. 424.)

+ rH_2O . Efflorescent. (Maquenne 108. 1303.)

Barium hydrogen hyponitrite, BaH_2

Easily sol. in H_2O . (Zorn, B. 1011.)

Calcium hyponitrite, $CaN_2O_2 + 4H_2O$

Nearly insol. in H_2O ; easily sol. in acids. (Maquenne, C. R. 108. 1303.)

Sl. sol. in H_2O ; insol. in alcohol. (Kirschner, Z. anorg. 1898, 16. 426.)

nitrite, basic, CuN_2O_2 , $\text{Cu}(\text{OH})_2$.
 H_2O ; not decomp. by hot H_2O .
 acids and in ammonia. Decomp.
 (Divers, Chem. Soc. 1899, 75. 121.)
 H_2O . Sol. in dil. acids and in
 aq. (Kirschner, Z. anorg. 1898,

ponitrite, $\text{Cu}_2\text{N}_2\text{O}_2 + 2\text{H}_2\text{O}$.
 Plotow, C. C. 1891, I. 1859.)
 e formed. (Divers, Chem. Soc.
 1.)

itrite, basic, PbN_2O_2 , PbO .
 H_2O . Sol. in dil. acids from which
 ptd. by $\text{NaOH} + \text{Aq}$ or $\text{NH}_4\text{OH} +$
 hner, Z. anorg. 1898, 16. 430.)

itrite, PbN_2O_2 .
 H_2O ; sol. in dil. acids from which
 ptd. by $\text{NaOH} + \text{Aq}$ or $\text{NH}_3 + \text{Aq}$.

yponitrite, basic, 3HgO , HgN_2O_2 .
).
 sol. even in boiling dil. HNO_3 .
 l. in conc., very sol. in warm dil.
 r, Chem. Soc. 1897, 71. 349.)

hyponitrite, $\text{Hg}_2\text{N}_2\text{O}_2$.
 .. HNO_3 with slow decomp. (Ray,
 1907, 91. 1404.)

yponitrite, HgN_2O_2 .
 Cl, and in $\text{NaCl} + \text{Aq}$.
 very dil. alkali. (Divers, Chem.
 75. 119.)

hyponitrite, $\text{K}_2\text{N}_2\text{O}_2$.
 O . (van der Plaats.)
 hen dry.
 0% alcohol, and sl. sol. in abs.
 Divers, Chem. Soc. 1899, 75. 103.)

nitrite (nitrosyl silver),
 O_2 .
 H_2O . Easily sol. in dil. $\text{HNO}_3 +$
 $\text{O}_2 + \text{Aq}$.
 by H_3PO_4 , H_2S , and boiling
 Aq. (van der Plaats.)
 $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$; sol. in NH_4OH
 vers, C. N. 23. 206.)
 il. HNO_3 and H_2SO_4 and in conc.
 Aq; decomp. by HCl . (Kirsch-
 rg. 1898, 16. 431.)

ponitrite, $\text{Na}_2\text{N}_2\text{O}_2 + 6\text{H}_2\text{O}$.
 O . (van der Plaats.)

yponitrite, SrN_2O_2 .
 ol. in H_2O . (Roederer, Bull.
 (3) 35. 715.)
 Nearly insol. in H_2O ; easily sol.

in dil. acids. (Maquenne, C. R. 108. 1303.)
 Sl. sol. in H_2O , insol in alcohol. (Kirsch-
 ner, Z. anorg. 1898, 16. 426.)

Hypophosphomolybdic acid, Mo_2O_7 ,
 $7\text{H}_3\text{PO}_2 + 3\text{H}_2\text{O}$.

Very sl. sol. in cold H_2O . Scarcely sol.
 in cold dil. H_2SO_4 . Sol. in cold conc. H_2SO_4 .
 Sol. in warm conc. HCl . Warm HNO_3
 oxidizes forming clear solution. (Mawrow,
 Z. anorg. 1901, 28. 164.)

Ammonium hypophosphomolybdate,
 $2(\text{NH}_4)_2\text{O}$, $2\text{H}_3\text{PO}_2$, $8\text{MoO}_3 + 2\text{H}_2\text{O}$.

Not very sol. in cold H_2O , readily in hot
 H_2O . (Gibbs, Am. Ch. J. 3. 402.)

Hypophosphoric acid, $\text{H}_4\text{P}_2\text{O}_6$.

Very deliquescent, and sol. in the least
 amount of H_2O . (Joly, C. R. 101. 1058.)
 100 cc. $\text{H}_4\text{P}_2\text{O}_6 + \text{Aq}$, containing 4.1% P_2O_5
 has sp. gr. = 1.036.

100 cc. $\text{H}_4\text{P}_2\text{O}_6 + \text{Aq}$, containing 12.3%
 P_2O_5 has sp. gr. = 1.122.
 (Salzer, A. 1878, 194. 28.)

+ H_2O . (Sanger, A. 232. 14.)

Does not exist. (Joly.)

+ $2\text{H}_2\text{O}$. Appears to be the only stable
 hydrate between 0° and 60° .

Sänger's hydrate, $\text{H}_4\text{P}_2\text{O}_6 + \text{H}_2\text{O}$, and
 Joly's anhydride could not be obtained.
 (Rosenheim, B. 1908, 41. 2711.)

Aluminum hypophosphate, $\text{Al}_4(\text{P}_2\text{O}_6)_3 +$
 $23\text{H}_2\text{O}$.

Easily sol. in mineral acids. Sol. in $\text{Na}_4\text{P}_2\text{O}_6$
 + Aq. (Palm, Dissertation, Rostock, 1890.)

Ammonium hypophosphate, $(\text{NH}_4)_4\text{P}_2\text{O}_6 +$
 H_2O .

Sol. in 30 pts. H_2O . (Salzer, A. 194. 32.)

Ammonium hydrogen hypophosphate,
 $(\text{NH}_4)_2\text{H}_2\text{P}_2\text{O}_6$.

Sol. in 14 pts. cold, and 4 pts. boiling H_2O .
 (Salzer, A. 194. 32.)

Ammonium trihydrogen hypophosphate,
 $\text{NH}_4\text{H}_3\text{P}_2\text{O}_6$.

Sol. in H_2O . (Salzer, A. 211. 1.)

Ammonium magnesium hypophosphate,
 $(\text{NH}_4)_2\text{MgP}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Precipitate. (Salzer, A. 232. 114.)

Barium hypophosphate, $\text{Ba}_2\text{P}_2\text{O}_6$.

Very slightly sol., but not wholly insol. in
 H_2O . Very slightly sol. in acetic acid, but
 more soluble in hydrochloric, and hypophos-
 phoric acids. (Salzer, A. 194. 34.)

Barium hydrogen hypophosphate, $\text{BaH}_2\text{P}_2\text{O}_6$
 + $2\text{H}_2\text{O}$.

Soluble in about 1000 pts. H_2O . Solution
 decomposes by heating. (Salzer, A. 194. 34.)

Bismuth hypophosphate, $\text{Bi}_4(\text{P}_2\text{O}_5)_3 + 8\frac{1}{2}\text{H}_2\text{O}$.

Completely sol. in $\text{HCl} + \text{Aq}$, also in warm $\text{HNO}_3 + \text{Aq}$. Insol. in boiling dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Sl. sol. by long boiling with conc. H_2SO_4 . (Palm, Rostock, 1890.)

Cadmium hypophosphate, $\text{Cd}_2\text{P}_2\text{O}_6 + 2\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in dil. acids. (Drawe, B. 21. 3403.)

Cadmium potassium hydrogen hypophosphate, $\text{CdK}_2(\text{H}_2\text{P}_2\text{O}_6)_2 + 2\frac{1}{2}\text{H}_2\text{O}$.

(Bausa, Z. anorg. 1894, 6. 147.)

Cadmium sodium hypophosphate, $\text{CdNa}_2\text{P}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Insol. in H_2O , but decomp. thereby. Sol. in dil. acids. (Drawe.)

Calcium hypophosphate, $\text{Ca}_2\text{P}_2\text{O}_6 + 2\text{H}_2\text{O}$.

Insol. in H_2O ; difficultly sol. in $\text{HC}_2\text{H}_3\text{O}_2$; easily sol. in $\text{H}_4\text{P}_2\text{O}_6$, or $\text{HCl} + \text{Aq}$. (Salzer, A. 194. 36.)

Calcium hydrogen hypophosphate, $\text{CaH}_2\text{P}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Sol. in 60 pts. H_2O . (Salzer, A. 232. 114.)

Chromic hypophosphate, $\text{Cr}_4(\text{P}_2\text{O}_5)_3 + 34\text{H}_2\text{O}$.

Sol. in $\text{HCl} + \text{Aq}$ on sl. warming, also in $\text{HNO}_3 + \text{Aq}$. Not completely sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, but completely sol. in conc. H_2SO_4 . (Palm, Dissertation, Rostock, 1890.)

Cobaltous hypophosphate, $\text{Co}_2\text{P}_2\text{O}_6 + 8\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in acids. (Drawe, B. 21. 3403.)

Cobaltous potassium hypophosphate, $\text{CoK}_2\text{P}_2\text{O}_6 + 5\text{H}_2\text{O}$.

Ppt. (Bausa, Z. anorg. 1894, 6. 156.)

Cobaltous potassium hydrogen hypophosphate, $\text{CoH}_2\text{P}_2\text{O}_6, 3\text{K}_2\text{H}_2\text{P}_2\text{O}_6 + 15\text{H}_2\text{O}$.

Cobaltous sodium hypophosphate, $\text{CoNa}_2\text{P}_2\text{O}_6 + 1\frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O , but decomp. thereby. Sol. in dil. acids. (Drawe, B. 21. 3403.)

Cupric hypophosphate, $\text{Cu}_2\text{P}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in dil. acids. (Drawe, B. 21. 3403.)

Ppt. (Bausa, Z. anorg. 1894, 6. 145.)

Cupric potassium hydrogen hypophosphate, $\text{CuH}_2\text{P}_2\text{O}_6, 3\text{K}_2\text{H}_2\text{P}_2\text{O}_6 + 15\text{H}_2\text{O}$.

Ppt. (Bausa, Z. anorg. 1894, 6. 152.)

Glucinum hypophosphate, $\text{Gl}_2\text{P}_2\text{O}_6 + 7\text{H}_2\text{O}$.

Insol. in H_2O . Moderately sol. in all mineral acids. (Palm, Rostock, 1890.)
+ $3\text{H}_2\text{O}$. (Rammelsberg.)

Iron (ferrous) hypophosphate, $4\frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in cold Decomp. by hot $\text{HNO}_3 + \text{Aq}$ into Insol. in $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. Somewhat sol. in c but a ppt. separates out on heating Rostock, 1890.)

Iron (ferric) hypophosphate, $\text{Fe}_2\text{P}_2\text{O}_6 + 20\text{H}_2\text{O}$.

Easily sol. in $\text{HCl} + \text{Aq}$. Whol. HNO_3 , and dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Con in conc. H_2SO_4 by warming a short ppt. separates out on boiling. (Pa

Lead hypophosphate, $\text{Pb}_2\text{P}_2\text{O}_6$.

Insol. in H_2O , $\text{HC}_2\text{H}_3\text{O}_2$, or H sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Salzer.)

Lithium hypophosphate, $\text{Li}_2\text{P}_2\text{O}_6$.

Very sl. sol. in H_2O . (Salzer, Sol. in 120 pts. H_2O at ord. tem melsberg, J. pr. (2) 45. 153.)

$\text{Li}_2\text{H}_2\text{P}_2\text{O}_6 + 2\text{H}_2\text{O}$. Deliquesce melsberg.)

Magnesium hypophosphate, $\text{Mg}_2\text{P}_2\text{O}_6 + 12\text{H}_2\text{O}$.

Sol. in 15,000 pts. H_2O ; sl. so easily in hypophosphoric, or mi (Salzer, A. 232. 114.)

+ $24\text{H}_2\text{O}$. (Rammelsberg.)

Magnesium hydrogen hypophosphate, $\text{MgH}_2\text{P}_2\text{O}_6 + 4\text{H}_2\text{O}$.

Sol. in 200 pts. H_2O . (Salzer, A

Manganese hypophosphate, $\text{Mn}_2\text{P}_2\text{O}_6 + 2\frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in mineral in acetic acid. (Palm, Dissertation, 1890.)

Manganous potassium hydrogen phosphate, $\text{MnH}_2\text{P}_2\text{O}_6, \text{K}_2\text{H}_2\text{P}_2\text{O}_6$.

Ppt. (Bausa, Z. anorg. 1894, 6

Manganous sodium hypophosphate, $\text{Na}_2\text{P}_2\text{O}_6 + 11\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in mineral aci

Nickel hypophosphate, $\text{Ni}_2\text{P}_2\text{O}_6 + 1$

Insol. in H_2O . Sol. in dil. acid B. 21. 3401.)

Nickel potassium hypophosphate, $\text{NiK}_2\text{P}_2\text{O}_6 + 6\text{H}_2\text{O}$.

Ppt. (Bausa, Z. anorg. 1894, 6

Nickel potassium hydrogen hyp, $\text{NiH}_2\text{P}_2\text{O}_6, 3\text{K}_2\text{H}_2\text{P}_2\text{O}_6 + 15\text{H}_2\text{O}$.

Ppt. (Bausa, Z. anorg. 1894, 6

m hypophosphate, $\text{NiNa}_2\text{P}_2\text{O}_6 + 3\text{H}_2\text{O}$, but decomp. thereby. Easily diss. (Drawe.)

hypophosphate, $\text{K}_4\text{P}_2\text{O}_6 + 8\text{H}_2\text{O}$. Ppt. H_2O ; insol. in alcohol. (Salzer, A. 211. 1.)

hydrogen hypophosphate, $\text{H}_4\text{P}_2\text{O}_6 + 3\text{H}_2\text{O}$. Ppt. H_2O . (Salzer, A. 211. 1.)

hydrogen hypophosphate, $\text{H}_4\text{P}_2\text{O}_6 + 3\text{H}_2\text{O}$, and $+2\text{H}_2\text{O}$. Sol. cold, and 1 pt. boiling H_2O . (Salzer, A. 211. 1.)

hydrogen hypophosphate, $\text{H}_4\text{P}_2\text{O}_6 + 3\text{H}_2\text{O}$. Pts. cold, and $\frac{1}{2}$ pt. hot H_2O . (Salzer, A. 211. 1.)

entahydrogen dihypophosphate, $\text{O}_6)_2 + 2\text{H}_2\text{O}$. Pts. cold, and $\frac{1}{4}$ pt. boiling H_2O . (Salzer, A. 211. 1.)

dium hypophosphate, $\text{O}_6 + 9\text{H}_2\text{O}$. Ppt. 25 pts. cold, and 3 pts. hot H_2O . (Z. anorg. 1894, 6. 158.)

nc hydrogen hypophosphate, $\text{O}_6, 3\text{K}_2\text{H}_2\text{P}_2\text{O}_6 + 15\text{H}_2\text{O}$. Ppt. H_2O . (Z. anorg. 1894, 6. 148.)

osphate, $\text{Ag}_4\text{P}_2\text{O}_6$. Sol. in H_2O . Easily sol. in HNO_3 , or H_2SO_4 . Very sl. sol. in $\text{H}_4\text{P}_2\text{O}_6 + \text{Aq}$. (Z. anorg. 1894, 6. 114.)

phosphate, $\text{Na}_4\text{P}_2\text{O}_6 + 10\text{H}_2\text{O}$. Ppt. 30 pts. cold, much more easily H_2O . (Salzer.)

ogen hypophosphate, $\text{Na}_2\text{HP}_2\text{O}_6$. Ppt. H_2O . (Salzer.)

drogen hypophosphate, $\text{O}_6 + 6\text{H}_2\text{O}$. Pts. cold, and 5 pts. boiling H_2O . Sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Insol. in alcohol. (A. 187. 331.)

drogen hypophosphate, O_6 . Ppt. H_2O . (Salzer, A. 211. 1.)

drogen dihypophosphate, $\text{O}_6)_2$. Sol. in 15 pts. cold H_2O . (Salzer, A. 211. 1.)

Thallium hypophosphate, $\text{Tl}_4\text{P}_2\text{O}_6$. Sl. sol. in H_2O . Decomp. in sunlight. (Joly, C. R. 1894, 118. 650.)

Thallium hydrogen hypophosphate, $\text{Tl}_2\text{H}_2\text{P}_2\text{O}_6$. Sol. in H_2O . (Joly.)

Zinc hypophosphate, $\text{Zn}_2\text{P}_2\text{O}_6 + 2\text{H}_2\text{O}$. Insol. in H_2O . Easily sol. in dil. acids. (Drawe, B. 21. 3403.)

Hypophosphorosomolybdc acid.

Barium hypophosphorosomolybdate, $\text{BaO}, \text{Mo}_7\text{O}_{24}, 3\text{H}_2\text{PO}_3 + 12\text{H}_2\text{O}$. Very sol. in H_2O and $\text{BaCl}_2 + \text{Aq}$. (Mawrow, Z. anorg. 1902, 29. 156.)

Hypophosphorous acid, H_3PO_3 . Very sol. in H_2O and alcohol. (Rose.)

Aluminum hypophosphite. Not deliquescent, but very sol. in H_2O . (Rose, Pogg. 12. 86.)

Ammonium hypophosphite, $\text{NH}_4\text{H}_2\text{PO}_3$. Sol. in H_2O , less deliquescent than the potassium salt. (Wurtz, A. ch. (3) 7. 193.) Very sol. in absolute alcohol. (Dulong.) Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.) Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4328.)

Barium hypophosphite, $\text{Ba}(\text{H}_2\text{PO}_3)_2 + \text{H}_2\text{O}$. Sol. in 3.5 pts. cold, and 3 pts. boiling H_2O . Insol. in alcohol. (Wurtz, A. 43. 323.)

Bismuth hypophosphite, $\text{Bi}(\text{H}_2\text{PO}_3)_3$. Ppt. (Vanino, J. pr. 1906, (2) 74. 150.) $+ \text{H}_2\text{O}$. Sol. in acid $\text{Bi}(\text{NO}_3)_3 + \text{Aq}$. (Haga, Chem. Soc. 1895, 67. 229.)

Cadmium hypophosphite. Sol. in H_2O . (Rose, Pogg. 12. 91.)

Calcium hypophosphite, $\text{Ca}(\text{PH}_2\text{O}_2)_2$. Sol. in 6 pts. cold, and not much more sol. in hot H_2O . Insol. in strong, very sol. in weak alcohol. (Rose, Pogg. 9. 361.)

Calcium cobaltous hypophosphite. $2\text{Ca}(\text{PH}_2\text{O}_2)_2, \text{Co}(\text{PH}_2\text{O}_2)_2 + 2\text{H}_2\text{O}$. Efflorescent. (Rose, Pogg. 12. 295.)

Calcium ferrous hypophosphite. Sol. in H_2O . (Rose, Pogg. 12. 294.)

Cerous hypophosphite, $\text{Ce}(\text{PH}_2\text{O}_2)_3 + \text{H}_2\text{O}$. Sl. sol. in H_2O . (Rammelsberg, B. A. B. 1872. 437.)

Chromium hypophosphite, $\text{Cr}_2(\text{OH})_2(\text{H}_2\text{PO}_2)_4$.

Anhydrous. Insol. in H_2O or dil. acids.
+ $3\text{H}_2\text{O}$. Sol. in H_2O . (Wurtz, A. ch. (3) 16. 196.)

Cobaltous hypophosphite, $\text{Co}(\text{PH}_2\text{O}_2)_2 + 6\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O . (Rose, Pogg. 12. 87.)

Cupric hypophosphite, $\text{Cu}(\text{PH}_2\text{O}_2)_2$.

Very sol. in H_2O , but very easily decomp. on heating. (Wurtz, A. ch. (3) 16. 199.)

Glucinum hypophosphite.

Sol. in H_2O . (Rose, Pogg. 12. 86.)

Iron (ferrous) hypophosphite, $\text{Fe}(\text{PH}_2\text{O}_2)_2 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Rose, Pogg. 12. 294.)

Iron (ferric) hypophosphite.

Difficultly sol. in H_2O or acids. Decomp. on boiling. Sl. sol. in $\text{H}_3\text{PO}_2 + \text{Aq}$. (Rose.)

Lead hypophosphite, $\text{Pb}(\text{PH}_2\text{O}_2)_2$.

Difficultly sol. in cold, more easily in hot H_2O . Insol. in alcohol. (Rose, Pogg. 12. 288.)

Lithium hypophosphite, $\text{LiH}_2\text{PO}_2 + \text{H}_2\text{O}$.

Sol. in H_2O . (Rammelsberg, B. A. B. 1872. 416.)

Magnesium hypophosphite, $\text{Mg}(\text{PH}_2\text{O}_2)_2 + 6\text{H}_2\text{O}$.

Efflorescent in dry air. Sol. in H_2O . (Rose.)

Manganous hypophosphite, $\text{Mn}(\text{H}_2\text{PO}_2)_2 + \text{H}_2\text{O}$.

Permanent. Very sol. in H_2O . (Wurtz, A. ch. (3) 16. 195.)

Mercurous hypophosphite nitrate,

$\text{HgH}_2\text{PO}_2, \text{HgNO}_2 + \text{H}_2\text{O}$.

Sl. sol. in H_2O with rapid decomp.
Sol. in hot conc. HNO_3 . (Haga, Chem. Soc. 1895, 67. 227.)

Nickel hypophosphite, $\text{Ni}(\text{PH}_2\text{O}_2)_2 + 6\text{H}_2\text{O}$.

Efflorescent. Sol. in H_2O . (Rammelsberg, B. 5. 494.)

Nickel hypophosphite ammonia,

$\text{Ni}(\text{H}_2\text{PO}_2)_2, 6\text{NH}_3$.

(Ephraim, B. 1913, 46. 3111.)

Platinous hypophosphite $\text{Pt}(\text{PH}_2\text{O}_2)_2$.

Insol. in H_2O , HCl , $\text{H}_2\text{SO}_4 + \text{Aq}$, etc. Sol. in $\text{HNO}_3 + \text{Aq}$. Insol. in alcohol. (Engel, C. R. 91. 1068.)

Potassium hypophosphite, KH_2PO_2 .

Very deliquescent. Very sol. in H_2O , sol. in weak, less in absolute alcohol. Insol. in ether. (Wurtz, A. ch. (3) 7. 192.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Sodium hypophosphite, $\text{NaH}_2\text{PO}_2 + \text{H}_2\text{O}$.

Very deliquescent. Somewhat less sol. than the K salt. Very sol. in absolute alcohol. (Dulong.)

Very sol. in H_2O , and somewhat less sol. in alcohol. (Rammelsberg, B. A. B. 1872. 412.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Strontium hypophosphite, $\text{Sr}(\text{PH}_2\text{O}_2)_2$.

Very easily sol. in H_2O . (Dulong.)

Insol. in alcohol. (Wurtz.)

Thallous hypophosphite, TlH_2PO_2 .

Sol. in H_2O . (Rammelsberg, B. A. B. 1872. 492.)

Uranyl hypophosphite, $\text{UO}_2(\text{H}_2\text{PO}_2)_2 + \text{H}_2\text{O}$.

Sl. sol. in H_2O . Easily sol. in HCl or $\text{HNO}_3 + \text{Aq}$. (Rammelsberg, Chem. Soc. (2) 11. 1.)

Divanadyl hypophosphite, $\text{V}_2\text{O}_5(\text{H}_2\text{PO}_2)_2 + 2\text{H}_2\text{O}$.

Insol. in cold, sl. sol. in hot H_2O .

Sol. in hot dil. HCl , H_2SO_4 , and HNO_3 , and in warm conc. HCl and H_2SO_4 .

Insol. in oxalic acid. (Mawrow, Z. anorg. 1907, 55. 147.)

Zinc hypophosphite, $\text{Zn}(\text{H}_2\text{PO}_2)_2 + \text{H}_2\text{O}$.

Sol. in H_2O .

+ $6\text{H}_2\text{O}$. Efflorescent. (Wurtz, A. ch. (3) 16. 195.)

Zirconium hypophosphite, $\text{Zr}(\text{OPH}_2\text{O})_4 + \text{H}_2\text{O}$.

Sensitive to light. Insol. in alcohol, by which it is pptd. from aqueous solution. (Hauser, Z. anorg. 1913, 84. 93.)

Hypophosphotungstic acid.

Potassium hypophosphotungstate, $4\text{K}_2\text{O}, 6\text{H}_2\text{PO}_2, 18\text{WO}_3 + 7\text{H}_2\text{O}$.

Precipitate. Sol. in hot, very al. sol. in cold H_2O . (Gibbs, Am. Ch. J. 5. 361.)

Hyposulpharsenious acid.

Hyposulpharsenites, $\text{As}_2\text{S}_3, \text{M}_2\text{S}$.

Difficultly sol. in H_2O . (Berzelius.)

Do not exist. (Nilson, B. 4. 989.)

Hyposulphuric acid, $\text{H}_2\text{S}_2\text{O}_6$.

See Dithionic acid.

urous acid, $\text{H}_2\text{S}_2\text{O}_3$.

ulphuric acid.

urous (Hydrosulphurous) acid,

only in dil. aqueous solution, decomposes rapidly.

Formula is $\text{H}_2\text{S}_2\text{O}_4$, according to A. 211. 285.)

in alcohol than in H_2O . (Rossler, n. (3) 25. 845.)

hyposulphite, $(\text{NH}_4)_2\text{S}_2\text{O}_4$.

only in solution. (Prudhomme, 1899, (3) 21. 326.)

hydrogen hyposulphite,

H_2O_4 .

only in solution. (Prudhomme, 1899, (3) 21. 326.)

hyposulphite, $\text{CaS}_2\text{O}_4 + 1.5\text{H}_2\text{O}$.

sol. in H_2O . (Bazlen, B. 1905,

hyposulphite, MgS_2O_4 .

R. 1905, 140. 936.)

hyposulphite, $\text{K}_2\text{S}_2\text{O}_4 + 3\text{H}_2\text{O}$.

decomp.

in alcohol. (Bazlen, B. 1905, 38.

hyposulphite, $\text{Na}_2\text{S}_2\text{O}_4$.

Stable in dry air. (Bazlen, 1061.)

10 g. dissolve 24.1 g. of the anhydrous (Jellinek, Z. anorg. 1911, 70. 130.)

Solubility in H_2O .

5 g. of the solution contain at:

1.91 g. $\text{Na}_2\text{S}_2\text{O}_4$

1.67 g. "

1.49 g. "

W. 1061, Z. anorg. 1911, 70. 128.)

in alcohol. (Bazlen, B. 1905, 38.

hyposulphite, $\text{Na}_2\text{S}_2\text{O}_4$, ZnS_2O_4 .

in H_2O than ZnS_2O_4 . (Bazlen, 1060.)

hyposulphite, SrS_2O_4 .

W. 1060, O. (Moissan, C. R. 1902, 135.

hyposulphite, ZnS_2O_4 .

in H_2O ; about 1 pt. in 7 pts. is supersat. solutions readily.

W. 1060, 1905, 38. 1060.)

imido acid, $\text{V}_2\text{O}_5(\text{OH})_4$.

barium tetrahydroxide:

Hypovanadic acid, $\text{H}_2\text{V}_4\text{O}_9$.

See Vanadous acid.

Hypovanadic acid with vanadic acid.

See Vanadico vanadic acid.

Imidodimetaarsenic acid.

Ammonium imidodimetaarsenate,

$(\text{NH}_4)_2\text{O}_2\text{As}_2\text{O}_5\text{NH}_4$.

(Rosenheim and Jacobsohn, Z. anorg. 1906, 50. 307.)

Imidochromic acid.

Ammonium imidochromate,

$\text{NHCrO}(\text{ONH}_4)_2$.

Very sol. in H_2O with decomp. (Rosenheim and Jacobsohn, Z. anorg. 1906, 50. 299.)

Ammonium potassium imidochromate,

$\text{NH}_4\text{KCrO}_5\text{NH}_4$.

Decomp. on solution in H_2O . (Rosenheim, Z. anorg. 1906, 50. 302.)

Imidodiimidochromic acid.

Ammonium imidodiimido chromate,

$\text{NH}[\text{CrO}(\text{NH})\text{ONH}_4]_2$.

(Rosenheim and Jacobsohn, Z. anorg. 1906, 50. 303.)

Imidomolybdic acid.

Potassium imidomolybdate, $\text{NKM}_2\text{O}(\text{OK})_2$.

Unstable in air.

Very hygroscopic. Very sol. in H_2O . (Rosenheim, Z. anorg. 1906, 50. 305.)

Diimidodiphosphoromonamic acid,

$\text{HO}-\text{PO} < \begin{smallmatrix} \text{NH} \\ \text{NH} \end{smallmatrix} > \text{PO}-\text{NH}_2$.

Correct formula for pyrophosphotriamic acid of Gladstone. (Mente, A. 248. 241.)

Imidodiphosphoric acid,

$\text{HO}-\text{PO} < \begin{smallmatrix} \text{O} \\ \text{NH} \end{smallmatrix} > \text{PO}-\text{OH}$.

Correct name for pyrophosphamic acid. (Mente, A. 248. 251.)

Barium imidodiphosphate, $\text{Ba} \begin{smallmatrix} \text{O} \\ \text{O} \\ \text{O} \end{smallmatrix} \begin{smallmatrix} > \text{PO} \\ > \text{PO} \end{smallmatrix} > \text{NH}$.

Sl. sol. in H_2O . (Mente, A. 248. 243.)

Barium imidodiphosphate, basic,

$\text{Ba} \begin{smallmatrix} \text{O} \\ \text{O} \\ \text{O} \end{smallmatrix} \begin{smallmatrix} > \text{PO} \\ > \text{PO} \end{smallmatrix} > \text{N}-\text{Ba}-\text{N} \begin{smallmatrix} \text{PO}-\text{O} \\ \text{O} \\ \text{PO}-\text{O} \end{smallmatrix} \text{Ba} +$

$2\text{H}_2\text{O}$.

Ppt. (Mente.)

Ferric imidodiphosphate.

Sl. sol. in conc. acids: (Mente, A. 248. 241.)

Silver imidodiphosphate, $\text{Ag}_3\text{H}_2\text{P}_2\text{NO}_6$.

Insol. in H_2O . (Stokes, Am. Ch. J. 1896, 18. 660.)

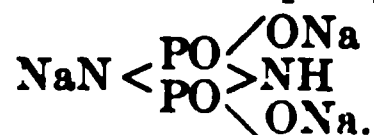
$\text{Ag}_4\text{HP}_2\text{NO}_6$. Ppt. (Stokes.)

Diimidodiphosphoric acid,

Correct name for *pyrophosphodiamic acid*. (Mente, A. 248. 241.)

Barium diimidodiphosphate,

Sl. sol. in dil. acids. (Mente, A. 248. 244.)

Sodium diimidodiphosphate, basic,

Sl. sol. in H_2O . (Mente, A. 248. 245.)

Diimidotriphosphoric acid.**Silver diimidotriphosphate, $\text{Ag}_3\text{H}_4\text{P}_3\text{N}_2\text{O}_8$.**

Insol. in H_2O . Very sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
Rather sl. sol. in dil. HNO_3 . (Stokes, Am. Ch. J. 1896, 18. 657.)

$\text{Ag}_5\text{H}_2\text{P}_3\text{N}_2\text{O}_8$. Insol. in H_2O . Very sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Decomp. by HNO_3 . (Stokes.)

Trisodium diimidotriphosphate,

Sol. in H_2O .

Insol. in alcohol. (Stokes.)

Triimidotetraphosphoric acid.**Silver triimidotetraphosphate.**

Ppt. (Stokes, Am. Ch. J. 1898, 20. 755.)

Sodium triimidotetraphosphate,

Easily sol. in H_2O ; insol. in sodium acetate solution and dil. alcohol. (Stokes, Am. Ch. J. 1898, 20. 754.)

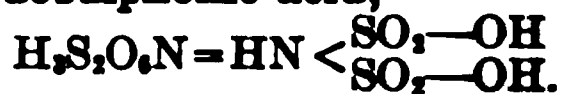
Imidosulphamide, $\text{NH}_2\cdot\text{SO}_2\cdot\text{NH}\cdot\text{SO}_2\cdot\text{NH}_2$.

"Sulphamide" of Traube.

Very sol. in H_2O with decomp. appreciably sol. in cold, easily sol. in hot methyl and ethyl alcohol. Insol. in C_6H_6 and CHCl_3 . Sl. sol. in ether, cold and hot acetic ether and glacial acetic acid. Moderately stable toward alkalis. (Hantzsch and Stuer, B. 1905, 38. 1022.)

Ammonium imidosulphamide,

(Hantzsch and Stuer.)

Imidosulphonic acid,

Ammondisulphonic acid of Claus. Known only in aqueous solution. (Divers and Haga, Chem. Soc. 61. 943.)

Very unstable. (Berglund, B. 9. 252.)

Ammonium imidosulphonate, basic,

Sol. in 9 pts. of H_2O . Solution is stable. Insol. in alcohol.

Sl. sol. in warm conc. H_2SO_4 without decomp. (Rose, Pogg. 1834, 32. 81.)

Much less sol. than the neutral salt. (Berglund, B. 9. 255.)

= "*Parasulphatammon.*"

+ H_2O . Gradually efflorescent. Sol. in H_2O with subsequent decomp. (Divers and Haga.)

Ammonium imidosulphonate, $\text{HN}(\text{SO}_3\text{NH}_4)_2$.

Sol. in H_2O . (Raschig, A. 241. 161.)

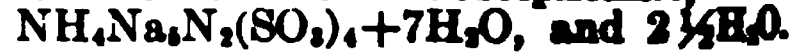
Ammonium barium imidosulphonate,

Very sl. sol. in H_2O . (Divers and Haga.)

$(\text{NH}_4)_2\text{Ba}_2\text{N}_4(\text{SO}_3)_8 + 8\text{H}_2\text{O}$. (D. and H.)

Ammonium calcium imidosulphonate.

(Divers, Chem. Soc. 1892, 61. 968.)

Ammonium sodium imidosulphonate,

Very sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Divers and Haga.)

Ammonium sodium imidosulphonate nitrate,

Very sol. in H_2O . (Divers and Haga.)

Barium imidosulphonate, $\text{Ba}[\text{N}(\text{SO}_3)_2\text{Ba}] + 5\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Berglund, B. 9. 255.)

Sol. in dil. $\text{HNO}_3 + \text{Aq}$ without decomp. (Divers and Haga.)

$\text{HN}(\text{SO}_3)_2\text{Ba} + \text{H}_2\text{O}$. Moderately sol. in H_2O . (D. and H.)

Barium mercury imidosulphonate,

Almost insol. in cold H_2O . (Divers and Haga, Chem. Soc. 1892, 61. 977.)

Barium sodium imidosulphonate,

Sparingly sol. in H_2O . Readily sol. in HNO_3 or HCl . (Divers, Chem. Soc. 1892, 61. 967.)

imidosulphonate, $\text{Ca}[\text{N}(\text{SO}_3)_2\text{Ca}]_2 + n \text{H}_2\text{O}$. (Berglund.)

mercury imidosulphonate, $[(\text{SO}_3)_2\text{Ca}]_2$.
sol. in H_2O . (Divers and Haga, c. 1896, 69. 1629.)

mercury imidosulphonate chloride, $(\text{SO}_3)_2\text{Ca}, \text{Hg}_2\text{Cl} + 12\text{H}_2\text{O}$.
by H_2O . (Divers and Haga, c. 1896, 69. 1629.)

sodium imidosulphonate, $(\text{SO}_3)_2\text{Ca} + 3\text{H}_2\text{O}$.
in cold H_2O . (Divers and Haga, c. 61. 968.)

osulphonate, $(\text{PbOH}\text{SO}_3)_2\text{NPbOH}$. (Berglund.)
 H_2O . (Divers and Haga.)
 $\text{N}(\text{SO}_3)_2$, PbO . Insol. in H_2O ; in dil. $\text{HNO}_3 + \text{Aq}$. (D. and H.)

s imidosulphonate, basic, $(\text{SO}_3)_2\text{Hg}_2\text{O} + 6\text{H}_2\text{O}$.
more sol. in dil. HNO_3 than mer-. Sol. in cold conc. $\text{KI} + \text{Aq}$, lf Hg as metal. (Divers and Haga, c. 1896, 69. 1631.)

imidosulphonate, basic, $(\text{SO}_3)_2\text{HgO}, \text{Hg}$.
decomp. (Divers and Haga.)

mercuric imidosulphonate, $\text{N}(\text{SO}_3)_2\text{Hg}_2\text{O} + 3\text{H}_2\text{O}$.
and Haga.)
 $(\text{SO}_3)_2\text{Hg}_2\text{O}$, $[\text{Hg}_2\text{N}(\text{SO}_3)_2\text{Hg}_2, (\text{SO}_3)_2\text{Hg}_2\text{O} + 6\text{H}_2\text{O}$. (Divers and

sodium imidosulphonate, basic, $\text{N}(\text{SO}_3)_2\text{Na} + 2\text{H}_2\text{O}$.
efflorescent. Decomp. by long with H_2O . Much more readily sol. in HNO_3 or H_2SO_4 and is wholly hereby. (Divers and Haga, Chem. 61. 983.)

sodium imidosulphonate, $(\text{SO}_3\text{Na})_4 + 6\text{H}_2\text{O}$.
ly sol. in cold H_2O .
sol in HNO_3 and in HCl .
by HCl immediately, but not. (Divers and Haga, Chem. Soc. 981.)

imidosulphonate, basic, $(\text{SO}_3\text{K})_2 + \text{H}_2\text{O}$.
 H_2O . (Raschig, A. 241. 161.)
than neutral salt. (Berglund.)

Potassium imidosulphonate, $\text{HN}(\text{SO}_3\text{K})_2$.
Sol. in H_2O . (Raschig, A. 241. 161.)
= Potassium ammondisulphonate of Claus.
Difficultly sol. in cold H_2O , sol. in 64 pts. H_2O at 23° . (Fremy.) Gradually decomp. by boiling. (Claus.)
Sl. sol. in H_2O . (Berglund, B. 9. 255.)

Potassium mercury imidosulphonate, $\text{N}_2\text{Hg}(\text{SO}_3\text{K})_4 + 4\text{H}_2\text{O}$.
See Mercurimidosulphonic acid.

Silver imidosulphonate, $\text{AgN}(\text{SO}_3\text{Ag})_2$.
Sl. sol. in H_2O . (Berglund.)

Silver sodium imidosulphonate, $\text{NaN}(\text{SO}_3\text{Ag})_2$.
Sl. sol. in H_2O . (Divers and Haga.)
 $\text{AgNa}_2\text{N}(\text{SO}_3)_2$. Sl. sol. in H_2O , but more sol. than the two preceding salts. (D. and H.)

Sodium imidosulphonate, $\text{HN}(\text{SO}_3\text{Na})_2 + 2\text{H}_2\text{O}$.
Not efflorescent. Very sol. in H_2O . (Diver and Haga.)
 $\text{NaN}(\text{SO}_3\text{Na})_2 + 12\text{H}_2\text{O}$. Efflorescent. Sl. sol. in cold H_2O , but very sol. in hot H_2O . Sol. in 5.4 pts. H_2O at 27.5° . (Divers and Haga.)

Sodium strontium imidosulphonate, $\text{SrNaNSO}_6 + 3\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Divers, Chem. Soc. 1896, 69. 1625.)

Strontium imidosulphonate, $\text{Sr}[\text{N}(\text{SO}_3)_2\text{Sr}]_2 + 6\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Berglund.)
 $+ 12\text{H}_2\text{O}$. Somewhat sol. in hot H_2O . (Divers, Chem. Soc. 1896, 69. 1623.)

Imidotrisulphoorthophosphoric acid, $\text{NH}:\text{P}(\text{SH})_3$.
Insol. in CS_2 and readily decomp. by H_2O . (Stock, B. 1906, 39. 1991.)

Ammonium imidotrisulphoorthophosphate, $\text{NHP}(\text{SNH}_2)_3$.
Very hygroscopic.
Loses NH_3 in the air.
Somewhat sol. in liquid NH_3 .
Decomp. by any other solvent in which it is sol. (Stock, B. 1906, 39. 1983.)

Diammonium hydrogen imidotrisulphoorthophosphate, $\text{SHP}(\text{SNH}_2)_2\text{NH}$.
(Stock, B. 1906, 39. 1983.)

Ammonium dihydrogen ———, $\text{SNH}_4\text{P}(\text{SH})_2\text{NH}$.
(Stock.)

Disodium hydrogen imidotrisulphoorthophosphate, $\text{SHP}(\text{SNa})_2\text{NH}$.

Very easily sol. in H_2O . Decomp. by H_2O . Somewhat sol. in methyl and ethyl alcohol. (Stock.)

Diimidopentasulphopyrophosphoric acid, $\text{P}_2\text{S}_5\text{N}_2\text{H}_4$.

Not known in pure state. (Stock, B. 1906, 39. 1967.)

Ammonium diimidopentasulphopyrophosphate, $\text{S}[\text{P}(\text{SNH}_2)_2\text{NH}]_2$.

Very hygroscopic.

Sol. in cold H_2O with decomp. (Stock, B. 1906, 39. 1978.)

Inidosulphurous acid.

Ammonium imidosulphite, $\text{HN}(\text{SO}_2\text{NH}_2)_2$.

Somewhat deliquescent.

Very unstable. Easily sol. in H_2O with decomp. into thiosulphate and amidosulphate.

Insol. in alcohol. (Divers and Ogawa, Chem. Soc. 1901, 79. 1100.)

Ammonium barium imidosulphite, $\text{Ba}(\text{SO}_2\text{NHSO}_2\text{NH}_2)_2$.

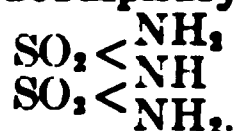
Sol. in H_2O . (Divers, Chem. Soc. 1901, 79. 1102.)

Potassium imidosulphite, $\text{NH}(\text{SO}_2\text{K})_2$.

(Divers and Owaga, Proc. Chem. Soc. 1900, 16. 113.)

Very sol. in H_2O . (Divers, Chem. Soc. 1901, 79. 1101.)

Imidosulphuryl amide, $\text{S}_2\text{O}_4\text{N}_2\text{H}_4 =$



Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by conc. HCl . Insol. in alcohol sat. with NH_3 . (Mente, A. 248. 265.)

Indic acid.

Magnesium indate, $\text{MgIn}_2\text{O}_4 + 3\text{H}_2\text{O}$.

Ppt. Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. (Renz, B. 1901, 34. 2764.)

Indium, In .

Does not decomp. hot H_2O .

Sol. in dil. HCl , and $\text{H}_2\text{SO}_4 + \text{Aq}$. Decomp. by conc. H_2SO_4 . Easily sol. in $\text{HNO}_3 + \text{Aq}$. Insol. in acetic acid. Insol. in $\text{KOH} + \text{Aq}$. (Winkler, J. pr. 102. 273.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0039 g. In in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Indium monobromide, InBr .

Decomp. by hot H_2O . Easily sol. in acid. Easily sol. in cold conc. HCl . (Thiel, Z. anorg. 1904, 40. 328.)

Indium dibromide, InBr_2 .

Decomp. by hot H_2O . Easily sol. in acid. (Thiel, Z. anorg. 1904, 40. 329.)

Indium tribromide, InBr_3 .

Deliquescent. Very sol. in H_2O .

Indium monochloride, InCl .

Deliquescent. Decomp. by H_2O into InCl_3 and In . (Nilson and Pettersson, Chem. Soc. 43. 820.)

Indium dichloride, InCl_2 .

Deliquescent in moist air; decomp. by H_2O into InCl_3 and In . (Nilson and Pettersson, Chem. Soc. 43. 818.)

Indium trichloride, InCl_3 .

Very deliquescent; sol. in H_2O with hissing and great evolution of heat.

Indium lithium chloride.

Extremely deliquescent. Sol. in H_2O . (Meyer, A. 150. 144.)

Indium potassium chloride, $3\text{KCl}, \text{InCl}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Easily sol. in H_2O . (Meyer.)

Indium trifluoride, $\text{InF}_3 + 3\text{H}_2\text{O}$.

Sol. in H_2O ; readily decomp. (Thiel, B. 1904, 37. 175.)

1 l. H_2O dissolves 86.4 g. at 25° . Decomp. on boiling. (Thiel, Z. anorg. 1904, 40. 331.) $+ 9\text{H}_2\text{O}$. Sl. sol. in cold H_2O .

Sol. in HCl and in HNO_3 .

Insol. in alcohol and ether. (Chabrie, C. R. 1905, 140. 90.)

Indium hydrosulphide.

Decomp. by acids. (Meyer.)

Indium hydroxide, $\text{In}_2\text{O}_3\text{H}_2$.

Sol. in acids; also in KOH , or $\text{NaOH} + \text{Aq}$ but the solution clouds up on standing or boiling, with separation of $\text{In}_2\text{O}_3\text{H}_2$. Insol. in NH_4OH , or $\text{NH}_4\text{Cl} + \text{Aq}$.

Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Renz, B. 1904, 37. 2110.)

Sl. sol. in alkylamines but completely ppt. by addition of the hydrochloride of the base. (Renz, B. 1903, 36. 2754.)

Indium monoiodide, InI .

Slowly decomp. in moist air. Not attacked by boiling H_2O .

Sol. in dil. HNO_3 in presence of AgNO_3 . Very slowly sol. in cold, more rapidly sol. in

ith evolution of H_2 . Very sol. in acid.

alcohol. ether and chloroform. norg. 1910, 66. 302.)

ide, InI_3 .

anorg. 1910, 66. 302.)

ide, InI_3 .

ent. (Meyer.)

ICl_3 . Decomp. by xylene. (Thiel, 1904, 40. 330.)

ide, InN .

by heat. (Franz Fischer, B. 1899.)

oxide, InO .

y sol. in $HCl + Aq$. (Winkler, J.

trioxide, In_2O_3 .

sl. in cold, easily in hot acids.

ifications:

ow. Amorphous. Sol. in acids.

de is insol. in ammonia and NH_4Cl .

e. Amorphous. Insol. in acids.

e. Amorphous. Sol. in acids.

de is sol. in ammonia, but pptd. by

tallized. Crystalline modification acids. (Renz, B. 1904, 37. 2112.)

liquid NH_3 . (Gore, Am. Ch. J. 30.)

de, $In_7O_9 = 3InO, 2In_2O_3$ (?).

)
 InO, In_2O_3 (?). (Winkler.)

bromide (?).

omp. by hot acids or alkalies. 150. 137.)

chloride, $InOCl$.

sol. in H_2O . (Thiel, B. 1904, 37.

sol. in cold or hot dil. acids.

l. in hot conc. acids. (Thiel, Z. 40. 327.)

elenide, In_2Se_3 .

rong acids with evolution of H_2Se . norg. 1910, 66. 315.)

ulphide, In_2S_3 .

cids. (Thiel, Z. anorg. 1904, 40.

osulphide, InS .

sl. in HCl with evolution of H_2S . INO_3 with evolution of oxides of (Thiel, Z. anorg. 1910, 66. 314.)

Indium sesquisulphide, In_2S_3 .

Partially sol. in $(NH_4)_2S + Aq$.

Indium potassium sulphide, In_2S_3, K_2S .

Insol. in H_2O ; decomp. by weak acids with separation of In_2S_3 ; sol. in conc. acids. (Schneider, J. pr. (2) 9. 209.)

Indium silver sulphide, In_2S_3, Ag_2S .

Insol. in H_2O . (Schneider, l. c.)

Indium sodium sulphide, $In_2S_3, Na_2S + 2H_2O$.

Insol. in H_2O . (Schneider, l. c.)

Indium monotelluride, $InTe$.

Sol. in HNO_3 ; insol. in $HCl + Aq$. (Thiel, Z. anorg. 1910, 66. 318.)

Infusible white precipitate.

Diiodamine, NHI_2 .

Decomp. by H_2O .

Iodammonium iodide, NIH_2I .

Decomp. by H_2O , caustic alkalies, and acids. Sol. in $KI + Aq$, alcohol, ether. CS_2 , $CHCl_3$. (Guthrie, Chem. Soc. (2) 1. 239.)

Iodauric acid, $HAuI_4$ (?).

Not known with certainty.

Ammonium iodaureate.

Deliquescent. Decomp. by H_2O . (Johnston, Phil. Mag. (3) 9. 266.)

Barium iodaureate.

Sol. in $BaI_2 + Aq$.

Cæsium iodaureate, $CsAuI_4$.

(Gupta, J. Am. Chem. Soc. 1914, 36. 748.)

Ferrous iodaureate.

Sol. in H_2O . (Johnston.)

Potassium iodaureate, $KAuI_4$.

Decomp. by H_2O . Sol. in KI , and $HI + A$. (Johnston.)

Sodium iodaureate.

Very deliquescent. (Johnston.)

Iodauricyanhydric acid, $HAu(CN)_2I_2$.

Known only in its salts.

Barium iodauricyanide, $Ba[Au(CN)_2I_2]_2 + 10H_2O$.

Sl. sol. in cold, easily in hot H_2O . Easily sol. in alcohol. (Lindbom, Lund. Univ. Arsk. 12. No. 6.)

Calcium iodaurycyanide, $\text{Ca}[\text{Au}(\text{CN})_2\text{I}_2]_2 + 10\text{H}_2\text{O}$.
Not stable. (L.)

Cobalt iodaurycyanide, $\text{Co}[\text{Au}(\text{CN})_2\text{I}_2]_2 + 10\text{H}_2\text{O}$.
Most insol. of all iodaurycyanides, and only sl. sol. in warm H_2O . Easily sol. in alcohol.

Potassium iodaurycyanide, $\text{KAu}(\text{CN})_2\text{I}_2 + \text{H}_2\text{O}$.
Sl. sol. in cold, easily sol. in warm H_2O and alcohol. (L.)

Strontium iodaurycyanide, $\text{Sr}[\text{Au}(\text{CN})_2\text{I}_2]_2 + 10\text{H}_2\text{O}$.
Sl. sol. in cold, more easily in hot H_2O .

Iodhydric Acid, HI.
Very easily and quickly absorbed by H_2O , with evolution of much heat.
Solution is decomp. on exposure to the air. 1 vol. H_2O absorbs 450 vols. HI at 10° . (Thomson.)
1 vol. H_2O absorbs 425 vols. HI at 10° . (Berthelot, C. R. 76. 679.)
Weak or strong solutions when boiled in an atmosphere of H leave a residue of constant composition, which distils unchanged at 126° (de Luynes), at 127° (Roscoe, Chem. Soc. 18. 146; Naumann; Topsoë), at 128° (Bineau, A. ch. (3) 7. 266); and has a sp. gr. of 1.67 (Naumann), of 1.70 (Bineau, de Luynes), of 1.708 (Topsoë); and contains 56.26 % HI (Bineau), 57.0% HI (Roscoe), 57.75% HI (Topsoë.)
By conducting dry H gas through the aqueous solution of HI, a constant residue is obtained, containing 60.3–60.7% HI if temp. is $15\text{--}19^\circ$, and 58.2–58.5% HI if temp. is 100° . (Roscoe.)

Solubility of HI in H_2O at t° .

t°	% HI	Solid Phase
–10	20.3	Ice
–20	29.3	
–30	35.1	
–40	39	
–50	42	
–60	44.4	
–70	46.2	
–80	47.9	Ice + HI, $4\text{H}_2\text{O}$
–60	52.6	
–40	59	HI, $4\text{H}_2\text{O}$
–35.5	64	
–40	65.5	"
–49	66.3	
–48	70.3	HI, $4\text{H}_2\text{O}$ + HI, $3\text{H}_2\text{O}$
–56	73.5	HI, $3\text{H}_2\text{O}$ + HI, $2\text{H}_2\text{O}$
–52	74	HI, $2\text{H}_2\text{O}$

(Pickering, B. 1893, 26. 2307.)

Solution in H_2O sat. at 0° has sp. gr. = 1.88 (de Luynes, A. ch. (4) 2. 385); 2.0 (Vignat.)

Sp. gr. of HI + Aq.

Sp. gr.	% HI	Temp.
1.017	2.286	13.5°
1.0524	7.019	13.5
1.077	10.15	13.5
1.095	12.21	13
1.102	13.09	13.5
1.126	15.73	13.5
1.164	19.97	13.5
1.191	22.63	13.8
1.225	25.86	13.8
1.2535	28.41	13.5
1.274	30.20	13.5
1.309	33.07	13
1.347	36.07	13
1.382	38.68	13
1.413	40.45	13
1.451	43.39	13
1.4865	45.71	13
1.528	48.22	13
1.542	49.13	13.5
1.5727	50.75	13
1.603	52.43	12.5
1.630	53.93	14
1.674	56.15	13.7
1.696	57.28	13
1.703	57.42	12.5
1.706	57.64	13.7
1.708	57.74	12

(Topsoe, B. 3. 403.)

Sp. gr. of HI + Aq at 15° .

% HI	Sp. gr.	% HI	Sp. gr.	% HI	Sp. g
1	1.008	21	1.175	41	1.41
2	1.015	22	1.185	42	1.42
3	1.022	23	1.195	43	1.44
4	1.029	24	1.205	44	1.46
5	1.037	25	1.216	45	1.47
6	1.045	26	1.227	46	1.48
7	1.053	27	1.238	47	1.51
8	1.061	28	1.249	48	1.52
9	1.069	29	1.260	49	1.54
10	1.077	30	1.271	50	1.56
11	1.085	31	1.283	51	1.57
12	1.093	32	1.295	52	1.58
13	1.102	33	1.307	53	1.61
14	1.110	34	1.320	54	1.62
15	1.118	35	1.333	55	1.64
16	1.127	36	1.346	56	1.67
17	1.137	37	1.359	57	1.68
18	1.146	38	1.372	58	1.71
19	1.155	39	1.386
20	1.165	40	1.400

(Topsoë, calculated by Gerlach, Z. anal. 316.)

Sp. gr. of HI + Aq at 15°.

HI	Sp. gr.	% HI	Sp. gr.	% HI	Sp. gr.
5	1.045	25	1.239	45	1.533
0	1.091	30	1.296	50	1.650
5	1.138	35	1.361	52	1.700
0	1.187	40	1.438

Only a "moderate degree of accuracy" is aimed for this table. (Wright, C. N. 23. 3.)

+2H₂O. Mpt.—43°. (Pickering, B. 1893, 1. 2308.)

+3H₂O. Mpt.—48°. (Pickering.)

+4H₂O. Mpt.—36.5°. (Pickering.)

dic acid, HIO₃.

Very sol. in H₂O and alcohol.

100 g. H₂O dissolve 286.1 g. HIO₃ at 13.5°.

. gr. of HIO₃ + Aq. = 2.4256.

100 g. H₂O dissolve 293 g. HIO₃ at 18°.

. gr. of HIO₃ + Aq. = 2.4711.

Groschuff, Z. anorg. 1905, 47. 337.)

Solubility of HIO₃ in H₂O at t°.

solid phase	t°	G. HIO ₃ in 100 g. of the solution	G. I ₂ O ₅ in 100 g. of the solution
ice	— 0.30	1.78	1.69
"	— 0.67	4.35	4.13
"	— 1.01	7.17	6.81
"	— 1.90	17.66	16.75
"	— 2.38	27.65	26.22
"	— 4.72	54.19	51.42
"	— 6.32	60.72	57.61
"	— 12.25	71.04	67.40
"	— 13.5	72.2	68.5
"	— 15	73.8	70.0
"	— 19	76.2	72.3
" + HIO ₃	— 14	72.8	69.1
HIO ₃	0	74.1	70.3
"	+ 16	75.6	71.7
"	40	77.7	73.7
"	60	80.0	75.9
"	80	82.5	78.3
"	85	83.0	78.7
"	101	85.2	80.8
" + HI ₂ O ₄	110	86.5	82.1
HI ₂ O ₄	125	87.2	82.7
"	140	88.3	83.8
"	160	90.5	85.9

(Groschuff, Z. anorg. 1905, 47. 343.)

sat. solution has sp. gr. 2.842 at 12.5°, and is at 104°. (Ditte, B. 6. 1533.) Sat. solution has sp. gr. 2.1629 (1.874 pts. I₂O₅ in 1 pt.) at 13°, and boils at 100°. (Kammerer, Z. 128. 400.)

Sp. gr. of HIO₃ + Aq at 15°.

% I ₂ O ₅	Sp. gr.	% I ₂ O ₅	Sp. gr.
1	1.0053	35	1.4428
5	1.0203	40	1.5371
10	1.0525	45	1.6315
15	1.1228	50	1.7356
20	1.2093	55	1.8689
25	1.2773	60	1.9954
30	1.3484	65	2.1269

(Kammerer.)

According to Thomsen (B. 7. 71) solutions of HIO₃ have sp. gr.—

HIO₃ + 10H₂O = 1.6609.

HIO₃ + 20H₂O = 1.3680.

HIO₃ + 40H₂O = 1.1945.

HIO₃ + 80H₂O = 1.1004.

HIO₃ + 160H₂O = 1.0512.

HIO₃ + 320H₂O = 1.0258.

H₂SO₄ at nearly boiling temp. dissolves 1/2 its weight of iodic acid. (Millon.)

Solubility in HNO₃ containing 27.78% HNO₃.

100 g. of the sat. solution contain at:

0°	20°	40°	60°
18	21	27	38 g. HIO ₃ .

Solubility in HNO₃ containing 40.88% HNO₃.

100 g. of the sat. solution contain at:

0°	20°	40°	60°
9	10	14	18 g. HIO ₃ .

(Groschuff, Z. anorg. 1905, 47. 344.)

Less sol. in HNO₃ than H₂O; nearly insol. in anhydrous HNO₃. (Groschuff, Z. anorg. 1905, 47. 347.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 830.)

Unattacked and undissolved by liquid NO₂. (Frankland, Chem. Soc. 1901, 79. 1362.)

Insol. in absolute alcohol. Alcohol of 35° B. dissolves half its weight in HIO₃. (Kammerer.)

+ 4 1/2 H₂O.

HIO₃, I₂O₅. (Groschuff, Z. anorg. 1905, 47. 343.)

Iodates.

The alkali iodates are sol. in H₂O, the others are sl. sol. or insol. therein.

Aluminum iodate, Al(IO₃)₃ (?).

Deliquescent. (Berzelius.)

Ammonium iodate, NH₄IO₃.

Sl. sol. in H₂O. Sol. in 38.5 pts. H₂O at 15°, 6.9 pts. at 100°. (Rammelsberg, Pogg. 44. 555.)

Solubility of NH_4IO_3 in $\text{HIO}_3 + \text{Aq}$ at 30° .

% HIO_3 in the solution	% NH_4IO_3 in the solution	Solid phase
0	4.20	NH_4IO_3
2.54	3.89	"
4.52	3.83	$\text{NH}_4\text{IO}_3 + \text{NH}_4\text{IO}_3, 2\text{HIO}_3$
4.51	3.86	"
4.56	3.75	"
4.73	3.53	$\text{NH}_4\text{IO}_3, 2\text{HIO}_3$
6.57	1.94	"
8.45	1.09	"
9.12	0.89	"
24.00	0.62	"
36.01	0.41	"
44.43	0.39	"
58.12	0.37	"
76.35	0.31	$\text{NH}_4\text{IO}_3, 2\text{HIO}_3 + \text{HIO}_3$
76.70	0	HIO_3

(Meerburg, Z. anorg. 1905, 45. 341.)

+ H_2O . (Ditte, A. ch. (6) 21. 146.)**Ammonium diiodate, $\text{NH}_4\text{H}(\text{IO}_3)_2$.**Sl. sol. in cold H_2O . (Ditte, A. ch. (6) 21. 145.)**Ammonium triiodate, $\text{NH}_4\text{H}_3(\text{IO}_3)_3$.**Sol. in H_2O . (Blomstrand, J. pr. (2) 42. 335.)See also solubility in HIO_3 , under Ammonium iodate. (Meerburg.)**Ammonium cobalt iodate.**Decomp. by H_2O . Insol. in alcohol. (Rammelsberg.)**Ammonium manganic iodate, $\text{Mn}(\text{IO}_3)_4, 2\text{NH}_4\text{IO}_3$.**Ppt. Insol. in H_2O . Insol. in HIO_3 . (Berg, C. R. 1899, 128. 675.)**Ammonium oxydimercuriammonium iodate.**

See Oxydimercuriammonium ammonium iodate.

Ammonium tellurium iodate.

See Iodotellurate, ammonium.

Ammonium iodate selenate.

See Iodoselenate, ammonium.

Barium iodate, $\text{Ba}(\text{IO}_3)_2$.Anhydrous salt is sol. in 1746 pts. H_2O at 15° , and 600 pts. H_2O at 100° (Rammelsberg, Pogg. 44. 577); in 3018 pts. H_2O at 13.5° , and 681 pts. H_2O at 100° . (Kremers, Pogg. 84. 27.)Solubility of $\text{Ba}(\text{IO}_3)_2$ in H_2O .
 $\text{Ba}(\text{IO}_3)_2 + \text{Aq}$ at t° contain
 $\text{Ba}(\text{IO}_3)_2$.

t°	Grams $\text{Ba}(\text{IO}_3)_2$	t°	Grams $\text{Ba}(\text{IO}_3)_2$
Eutectic point $-0.046^\circ \pm 0.002^\circ$	0.008	30°	0.031
+ 10°	0.014	40°	0.041
20°	0.022	50°	0.056
25°	0.028	60°	0.074

* Bpt. at 735 mm. pressure = at 760 mm. pressure.

(Anschütz, Z. phys. Ch. 1906)

1 l. sat. aq. solution contains $\text{Ba}(\text{IO}_3)_2$ at room temp. (Hill Am. Chem. Soc. 1909, 31. 44.)1 l. H_2O dissolves 0.3845 g. 25° . (Harkins and Winningham Chem. Soc. 1911, 33. 1828.)Easily sol. in cold $\text{HCl} + \text{Aq}$; c in warm $\text{HNO}_3 + \text{Aq}$. (Rammelsberg Insol. in H_2SO_4 . (Ditte.)100 cc. $\text{NH}_4\text{OH} + \text{Aq}$ (sp. gr. 0.92) solve 0.0199 g. $\text{Ba}(\text{IO}_3)_2$. (HillSolubility in salts + Aq at

C = concentration of salt in expressed in equivalents per l.

S = solubility of $\text{Ba}(\text{IO}_3)_2$ in expressed in equivalents per l.

Salt	C
$\text{Ba}(\text{NO}_3)_2$	0.001
	0.002
	0.005
	0.020
	0.050
	0.100
	0.200
KNO_3	0.002
	0.010
	0.050
	0.200
KIO_3	0.00010608
	0.0005304
	0.0010608

(Harkins and Winninghof, J. Am. Chem. Soc. 1911, 33. 1829.)

Insol. in alcohol.

100 cc. 95% alcohol dissolves $\text{Ba}(\text{IO}_3)_2$ at room temp. (Hill

Insol. in acetone. (Eidmann II. 1014.)

+ H_2O . Sol. in 3333 pts. H_2O at 15° , and 625 pts. H_2O at 100° . (Gay-Lussac 91. 5.)

Insol. in acetone. (Naumann 37. 4329.)

organic iodate,
 $\text{Ba}(\text{IO}_3)_2$.
 H_2O .
 HIO_3 . (Berg, C. R. 1899, 123.)

late, basic.
 H_2O . Very difficultly sol. in
 (Rammelsberg, Pogg. 44. 568.)
 $-1\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O .

date, $\text{Cd}(\text{IO}_3)_2$.
 ol. in H_2O . Easily sol. in HNO_3 ,
 $+\text{Aq}$. Sol. in $\text{Cd}(\text{C}_2\text{H}_3\text{O}_2)_2 +$
 nelsberg, Pogg. 44. 566.)
 Sl. sol. in H_2O . Very sol. in dil.
 (Ditte, A. ch. (6) 21. 145.)

date ammonia, $\text{Cd}(\text{IO}_3)_2, 2\text{NH}_3$.
 I_2O_5 ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Ditte.)
 1. 145.)
 $2\text{NH}_3 + \text{H}_2\text{O}$. As above. (Ditte.)

ate, CsIO_3 .
 I_2O dissolve 2.6 pts. CsIO_3 at 24° .
 ohol. (Wheeler, Sill. Am. J. 144.)

O_5 . 100 pts. H_2O dissolve 2.5 pts.
 decomp. by hot H_2O . (Wheeler.)
 $\text{O}_5, 2\text{HIO}_3$. Sl. sol. in cold H_2O .
 p. thereby into $2\text{CsIO}_3, \text{I}_2\text{O}_5$.

ate chloride, $\text{CsCl}, \text{HIO}_3$.
 by H_2O into $2\text{CsIO}_3, \text{I}_2\text{O}_5$.

rogen iodate periodate,
 $\text{IO}_4 + 2\text{H}_2\text{O}$.
 in dil. HNO_3 . (Wells, Am. Ch.
 280.)

ate, $\text{Ca}(\text{IO}_3)_2$.
 dissolve 0.22 pt. at 18° , and
 t. 100° . (Gay-Lussac.) Sol. in
 Aq . (Filhol.) Much more sol.
 q than in H_2O . (Rammelsberg.)
 SO_4 . (Ditte.) Scarcely sol. in
 Aq . (Sonstadt, C. N. 29. 209.)

Sat. solution contains at:
 35° 40° 45°
 1.48 0.52 0.54% $\text{Ca}(\text{IO}_3)_2$,
 50° 80° 100°
 1.65 0.79 0.94% $\text{Ca}(\text{IO}_3)_2$.
 and Funk, B. 1897, 30. 1724.)

Efflorescent.
 3 pts. H_2O at 15° , and 75 pts. at
 melsberg.)

Sat. solution contains at:
 0° 10° 18° 30°
 0.1 0.17 0.25 0.42% $\text{Ca}(\text{IO}_3)_2$,
 40° 50° 54° 60°
 0.61 0.89 0.14 1.36% $\text{Ca}(\text{IO}_3)_2$.
 (Mylius and Funk, B. 1897, 30. 1724.)

Much more sol. in $\text{HNO}_3 + \text{Aq}$. Pptd. by
 alcohol from $\text{Ca}(\text{IO}_3)_2 + \text{Aq}$.
 Insol. in H_2SO_4 . (Ditte.)
 Pptd. by alcohol from aqueous solution.
 (Henry.)

Cerous iodate, $\text{Ce}(\text{IO}_3)_3 + 2\text{H}_2\text{O}$.
 Sl. sol. in cold, easily sol. in hot H_2O and in
 acids. (Holzmann, J. pr. 75. 321.)
 Solubility in H_2O . 100 cc. of the sat.
 solution contain 0.1456 g. at 25° . (Rimbach,
 Z. phys. Ch. 1909, 67. 199.)
 Calc. from electrical conductivity of
 $\text{Ce}(\text{IO}_3)_3 + \text{Aq}$, 100 cc. of the sat. solution
 contain 0.1636 g. $\text{Ce}(\text{IO}_3)_3$ at 25° . (Rim-
 bach, Z. phys. Ch. 1909, 67. 199.)

Ceric iodate, $\text{Ce}(\text{IO}_3)_4$.
 Slightly hydrolyzed by H_2O .
 0.34 g. is sol. in 100 cc. hot conc. HNO_3 .
 (Barbieri, Chem. Soc. 1907, 92. (2) 467.)

Cobaltous iodate, $\text{Co}(\text{IO}_3)_2$.
 Anhydrous. Sol. in warm dil. H_3PO_4 , or
 $\text{H}_2\text{SO}_4 + \text{Aq}$. (Ditte, A. ch. (6) 21. 14.)

Solubility in H_2O			
Form	Temp.	% $\text{Co}(\text{IO}_3)_2$	Mols. of water free salt to 100 mols H_2O
$\text{Co}(\text{IO}_3)_2 + 4\text{H}_2\text{O}$	0°	0.54	0.028
"	18°	0.83	0.038
"	30°	1.03	0.046
"	50°	1.46	0.065
"	60°	1.86	0.084
"	65°	2.17	0.098
$\text{Co}(\text{IO}_3)_2 + 2\text{H}_2\text{O}$	0°	0.32	0.014
"	18°	0.45	0.020
"	30°	0.52	0.023
"	50°	0.67	0.030
"	75°	0.84	0.038
"	100°	1.02	0.045
$\text{Co}(\text{IO}_3)_2$	18°	1.03	0.046
"	30°	0.89	0.040
"	50°	0.85	0.036
"	75°	0.75	0.033
"	100°	0.69	0.031

(Meusser, B. 1901, 34. 2435.)
 $+\text{H}_2\text{O}$. Sol. in 148 pts. H_2O at 15° and
 90 pts. at 100° . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Ram-
 melsberg, Pogg. 44. 561.)
 Does not exist. (Meusser, B. 1901, 34.
 2434.)
 $+2\text{H}_2\text{O}$. (Meusser.)
 $+4\text{H}_2\text{O}$. (Meusser.)

Cupric iodate, basic, $6\text{CuO}, 3\text{I}_2\text{O}_5 + 2\text{H}_2\text{O}$.

Insol. in H_2O . (Millon, A. ch. (3) 9. 400.)
 Mixture of CuO and $\text{Cu}(\text{IO}_3)_2$. (Ditte, A. ch. (6) 21. 175.)
 $2\text{CuO}, \text{I}_2\text{O}_5 + \text{H}_2\text{O}$. Slowly sol. in dil. H_2SO_4 . (Granger and de Schulten, Bull. Soc. 1904, (3) 31. 1027.)

Cupric iodate, $\text{Cu}(\text{IO}_3)_2$.

1 l. H_2O dissolves 3.32×10^{-3} mol. $\text{Cu}(\text{IO}_3)_2$ at 25° . (Spencer, Z. phys. Ch. 1913, 83. 295.)

Solubility in $\text{CuSO}_4 + \text{Aq} = 3.28 \times 10^{-3}$ mol. per l. at 25° .

Solubility in $\text{KIO}_3 + \text{Aq} = 3.29 \times 10^{-3}$ mol. per l. at 25° . (Spencer.)

+ H_2O . (Ditte.)
 + $2\text{H}_2\text{O}$. Sol. in 302 pts. H_2O at 15° , and 154 pts. at 100° . Sol. in $\text{HCl} + \text{Aq}$ or $\text{NH}_4\text{OH} + \text{Aq}$. (Millon.)

Cupric iodate ammonia, $\text{Cu}(\text{IO}_3)_2, 2\text{NH}_3 + \text{H}_2\text{O}$.

Insol. in H_2O . (Ditte, A. ch. (6) 21. 145.)
 $\text{Cu}(\text{IO}_3)_2, 4\text{NH}_3 + 2\text{H}_2\text{O}$. Ppt. (Ephraim, B. 1915, 48. 52.)

+ $3\text{H}_2\text{O}$. Partially sol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in alcohol. (Rammelsberg.)

$\text{Cu}(\text{IO}_3)_2, 5\text{NH}_3$. (Ephraim.)
 $\text{Cu}(\text{IO}_3)_2, 8\text{NH}_3 + 4\text{H}_2\text{O}$. Sol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in alcohol. (Ditte, A. ch. (6) 21. 145.)

Decipium iodate, $\text{Dp}(\text{IO}_3)_3 + 3\text{H}_2\text{O} (?)$.

Precipitate; scarcely sol. in H_2O . (Dela-fontaine.)

Didymium iodate, $\text{Di}(\text{IO}_3)_3 + 2\text{H}_2\text{O}$.

Ppt. (Cleve.)

Erbium iodate, $\text{Er}(\text{IO}_3)_3 + 3\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (Hoglund.)

Glucinum iodate.

Deliquescent.

Indium iodate, $\text{In}(\text{IO}_3)_3$.

1 pt. is sol. in 1500 pts. H_2O at 20° .
 1 pt. is sol. in 150 pts. HNO_3 (1:5) at 80° .
 Sol. in HCl with decomp. Sol. in dil. H_2SO_4 . (Mathers, J. Am. Chem. Soc. 1908, 30. 213.)

Iodine iodate, $\text{I}(\text{IO}_3)_3$.

Decomp. by H_2O or by alcohol. (Fichter, Z. anorg. 1915, 91. 142.)

Iron (ferrous) iodate.

Ppt. Sl. sol. in H_2O ; more sol. in $\text{FeSO}_4 + \text{Aq}$. (Geiger, Mag. Pharm. 29. 252.)

Iron (ferric) iodate, $\text{Fe}_2\text{O}_3, \text{I}_2\text{O}_5$.

Insol. in acids. (Ditte, A. ch. (6) 21. 175.)
 $\text{Fe}_2\text{O}_3, 2\text{I}_2\text{O}_5 + 8\text{H}_2\text{O}$. Sol. in 50 parts of $\text{HNO}_3 + \text{Aq}$. Sol. in H_2O . (Geiger.)

$3\text{Fe}_2\text{O}_3, 5\text{I}_2\text{O}_5 + 15\text{H}_2\text{O}$. Sol. in $\text{HNO}_3 + \text{Aq}$. (Rammelsberg.)

Lanthanum iodate, $\text{La}(\text{IO}_3)_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Sl. sol. in cold, easily sol. in warm $\text{HCl} + \text{Aq}$. (Harkins, J. Am. Chem. Soc. 1911, 33. 349.)

100 cc. of the sat. solution in H_2O contains 0.1681 g. at 25° . (Rimbach, Z. anorg. 1909, 67. 199.)

Calc. from electrical conductivity. 100 cc. of the sat. solution contain 0.1871 g. $\text{La}(\text{IO}_3)_3$ at 25° . (Rimbach.)

Lead iodate, basic, $3\text{PbO}, \text{Pb}(\text{IO}_3)_2$.

Ppt. (Strömholm, Z. anorg. 1904, 30. 199.)

Lead iodate, $\text{Pb}(\text{IO}_3)_2$.

Very sl. sol. in H_2O (Pleischl, Z. anorg. 1904, 30. 199.)
 Difficultly sol. in $\text{HNO}_3 + \text{Aq}$. (Rammelsberg.)

Insol. in H_2O and $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in $\text{HNO}_3 + \text{Aq}$, and wholly in $\text{HNO}_3 + \text{Aq}$ after being heated to 100° . (Ditte, A. ch. (6) 21. 169.)

Sl. sol. in H_2O . 1.83×10^{-3} are dissolved in 1 liter of sat. solution at 20° . (Harkins, J. Am. Chem. Soc. 1903, 25. 603.)

1 l. H_2O dissolves 19 mg. $\text{Pb}(\text{IO}_3)_2$. (Kohlrausch, Z. phys. Ch. 1904, 30. 168.)
 17.8 mg. are dissolved in 1 l. of H_2O at 18° . (Kohlrausch, Z. phys. Ch. 1904, 30. 168.)

1 l. H_2O dissolves 0.0307 g. $\text{Pb}(\text{IO}_3)_2$ at 25° . (Harkins, J. Am. Chem. Soc. 1903, 25. 603.)

Solubility of $\text{Pb}(\text{IO}_3)_2$ in salts +

C = concentration of salt in g. expressed in equivalents per l.

S = solubility of $\text{Pb}(\text{IO}_3)_2$ in g. expressed in equivalents per l.

Salt	C	S
$\text{Pb}(\text{NO}_3)_2$	0.0001	0.0001
	0.001	0.001
	0.010	0.010
	0.100	0.100
	0.500	0.500
	3.0	3.0
KNO_3	0.002	0.002
	0.010	0.010
	0.050	0.050
	0.200	0.200
KIO_3	0.00005304	0.00005304
	0.0001061	0.0001061

(Harkins and Winninghof, J. Am. Chem. Soc. 1911, 33. 1830.)

quid NH_3 . (Gore, Am. Ch. J. 5.)

te, $\text{LiIO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

nt, and very sol. in H_2O .

ts. cold, and not much less hot in alcohol. (Rammelsberg, 5.)

olution sat. at $18^\circ = 1.568$, con-
% LiIO_3 . 100 g. H_2O dissolve
). (Mylius and Funk, B. 1897,

ethyl acetate. (Naumann, B. 0.)

ery deliquescent. (Ditte, A. ch.

odate, $\text{Mg}(\text{IO}_3)_2$.

Insol. in H_2O . (Millon, A. 5.)

Very sol. in H_2O . (Ditte.)

pts. H_2O at 15° , and 3.04 pts. (Berzelius.) Very sl. sol. in H_2O . (Ditte.) ch. 45. 279.) Easily sol. in dil. (Ditte.)

. solution contains at:

10°	20°
6.4	7.7% $\text{Mg}(\text{IO}_3)_2$,
63°	100°
12.6	19.3% $\text{Mg}(\text{IO}_3)_2$.

nd Funk, B. 1897, 30. 1722.)

olution at 18° contains 6.44%
6.88 g. are sol. in 100 g. H_2O .
. solution = 1.078. (Mylius and
7, 30. 1718.)

Sat. aq. solution contains at:

30°	35°	50° (m.pt.).
17.4	21.9	67.5% $\text{Mg}(\text{IO}_3)_2$.

nd Funk, B. 1897, 30. 1723.)

odate, $\text{Mn}(\text{IO}_3)_2 + \text{H}_2\text{O}$.

out 200 pts. H_2O . (Rammels-

I_2O and $\text{HNO}_3 + \text{Aq}$, even on
l. in $\text{NH}_4\text{OH} + \text{Aq}$. (Ditte.)

manganic iodate,

, $\text{Mn}(\text{IO}_3)_2$.

I_2O . (Berg, C. R. 1899, 128.

assium iodate,

, 2KIO_3 .

d only sl. attacked by H_2O .

HIO_3 . (Berg, C. R. 1899, 128.

date, $\text{Hg}_2(\text{IO}_3)_2$.

iling H_2O , or cold $\text{HNO}_3 + \text{Aq}$.
dil. $\text{HCl} + \text{Aq}$. Sol. in very

conc. $\text{HIO}_3 + \text{Aq}$. (Lefort, J. Pharm. 1845. 5.)

Mercuric iodate, $\text{Hg}(\text{IO}_3)_2$.

Insol. in H_2O or alcohol. (Millon, A. ch. (3) 18. 367.) Sol. in H_2O . (Berzelius.) Sol. in dil. $\text{HCl} + \text{Aq}$. (Rammelsberg.)

Nearly insol. in H_2O . Easily sol. in HCl , HBr , or $\text{HI} + \text{Aq}$; very sl. sol. in $\text{HNO}_3 + \text{Aq}$; insol. in HF , H_2SiF_6 , or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in alkali chlorides, bromides, iodides, cyanides, and cyanates + Aq ; also in $\text{Na}_2\text{S}_2\text{O}_3$, dil. MnCl_2 , and $\text{ZnCl}_2 + \text{Aq}$. Insol. in KOH , NaOH , NH_4OH , Na_2S , $\text{Na}_2\text{B}_4\text{O}_7$, Na_2HPO_4 , and the alkali chlorates, bromates, and iodates + Aq . (Cameron, C. N. 33. 253.)

Nickel iodate, $\text{Ni}(\text{IO}_3)_2$.

Solubility in H_2O .

Form	Temp.	Percent of $\text{Ni}(\text{IO}_3)_2$ in solution	Mols. water free salt to 100 mols. H_2O
$\text{Ni}(\text{IO}_3)_2 + 4\text{H}_2\text{O}$	0°	0.73	0.033
"	18°	1.01	0.045
"	30°	1.41	0.063
$\alpha\text{Ni}(\text{IO}_3)_2 + 2\text{H}_2\text{O}$	0°	0.53	0.023
"	18°	0.68	0.030
"	30°	0.86	0.039
"	50°	1.78	0.080
$\beta\text{Ni}(\text{IO}_3)_2 + 2\text{H}_2\text{O}$	8°	0.52	0.023
"	18°	0.55	0.0245
"	50°	0.81	0.035
"	75°	1.03	0.045
"	100°	1.12	0.049
$\text{Ni}(\text{IO}_3)_2$	30°	1.135	0.050
"	50°	1.07	0.046
"	75°	1.02	0.045
"	100°	0.988	0.044

(Meusser, B. 1901, 34. 2440.)

+ H_2O . Sol. in 120.3 pts. H_2O at 15° , and 77.35 pts. at 100° . (Rammelsberg, Pogg. 44. 562.)

Sol. in HNO_3 , and dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Ditte.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Does not exist (Meusser.)

+ $2\text{H}_2\text{O}$. See Meusser above.

+ $3\text{H}_2\text{O}$. Insol. in H_2O . Sol. in HNO_3 . (Ditte, A. ch. 1890, (6) 21. 160.)

+ $4\text{H}_2\text{O}$. See Meusser above.

Nickel iodate ammonia, $\text{Ni}(\text{IO}_3)_2, 4\text{NH}_3$.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in alcohol. (Rammelsberg, Pogg. 44. 562.)

$\text{Ni}(\text{IO}_3)_2, 5\text{NH}_3$. Ppt. (Ephraim, B. 1915, 48. 53.)

+ $3\text{H}_2\text{O}$. (Ephraim.)

Potassium iodate, KIO_3 .

1 pt. KIO_3 dissolves in 13 pts. H_2O at 14° . (Gay-Lussac.)

1 pt. KIO_3 dissolves at:

0°	in 21.11 pts. H_2O
20°	" 12.29 "
40°	" 7.76 "
60°	" 5.40 "
80°	" 4.02 "
100°	" 3.10 "

Sat. solution boils at 102°. (Kremers, Pogg. 97. 5.)

Sp. gr. of KIO_3 + Aq containing:

1	2	3	4	5	% KIO_3
1.010	1.019	1.027	1.035	1.044	
6	7	8	9	10	% KIO_3
1.052	1.061	1.071	1.080	1.090	

(Kremers, Pogg. 96. 62.)

Stable at 10° in H_2O or potassium acetate + Aq. (Eakle, C. C. 1896, II. 649.)

Solubility of KIO_3 in HIO_3 + Aq at 30°.

% HIO_3 in the solution	% KIO_3 in the solution	Solid phase
0	9.51	KIO_3
0.64	9.48	$\text{KIO}_3 + \text{KIO}_3, \text{HIO}_3$
0.66	9.52	"
0.65	9.46	"
0.65	8.90	$\text{KIO}_3, \text{HIO}_3$
0.67	6.6	"
1.14	4.57	"
1.69	3.63	"
2.02	3.10	"
3.34	2.14	"
5.00	1.32	"
7.09	1.0	"
8.04	0.85	$\text{KIO}_3, \text{HIO}_3 + \text{KIO}_3, 2\text{HIO}_3$
3.47	3.57	$\text{KIO}_3, 2\text{HIO}_3$ (labile)
4.80	2.90	"
6.45	1.35	"
9.35	0.64	$\text{KIO}_3, 2\text{HIO}_3$
12.04	0.44	"
17.50	0.30	"
31.20	0.52	"
53.64	0.68	"
62.52	0.72	"
76.40	0.80	$\text{KIO}_3, 2\text{HIO}_3 + \text{HIO}_3$
76.70	0	HIO_3

(Meerburg, Z. anorg. 1905, 45. 330.)

More sol. in $\text{KI} + \text{Aq}$ than in H_2O . Sol. in warm $\text{H}_2\text{SO}_4 + \text{Aq}$.

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in alcohol.

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43 314.)

+ $\frac{1}{2}\text{H}_2\text{O}$. (Ditte, C. R. 70. 621.)

Potassium hydrogen iodate, $\text{KH}(\text{IO}_3)_2$.

Sol. in 18.65 pts. H_2O at 17°. (Meineke, A. 261. 360.)

Sol. in 75 pts. H_2O at 15°. 1 alcohol. (Serullas, A. ch. 22. 181.)
See also Meerburg under KIO_3 .

Potassium dihydrogen iodate, KI

Sol. in 25 pts. H_2O at 15°. (Serulla 43. 117.)

See also Meerburg under KIO_3 .

Potassium tellurium iodate.

See Iodotellurate, potassium.

Potassium uranyl iodate,

$\text{KUO}_2(\text{IO}_3)_2 + 3\text{H}_2\text{O}$.

Decomp. by H_2O , dil. salt soluti $\text{UO}_2(\text{NO}_3)_2 + \text{Aq}$. (Artmann, Z. anorg. 79. 340.)

Potassium iodate chloride, $\text{KH}(\text{IO}_3)_2$

Sol. in 19 pts. H_2O at 15° with Cold alcohol dissolves out KCl .

Potassium iodate molybdate, $\text{KIO}_3, 2\text{H}_2\text{O}$.

See Molybdatiodate, potassium.

Potassium iodate selenate.

See Iodoselenate, potassium.

Potassium iodate sulphate, KIO_3, H

Decomp. by H_2O . (Marignac, J. 299.)

$\text{KHIO}_3, \text{KHSO}_4$. More sol. in 1 KHIO_3 . (Serullas.)

Potassium iodate tungstate.

See Tungstoiodate, potassium.

Rubidium iodate, RbIO_3 .

100 pts. H_2O dissolve 2.1 pta. 23°. Easily sol. in cold $\text{HCl} + \text{Aq}$. Sill. Am. J. 144. 123.)

Rubidium hydrogen iodate, $\text{RbH}(\text{IO}_3)_2$

Sl. sol. in cold, more readily in RbIO_3 separating on cooling. Insol. hol. (Wheeler.)

$\text{RbH}_2(\text{IO}_3)_3$. As above. (Whe

Rubidium iodate chloride, $\text{RbIO}_3, \text{HIO}_3, \text{RbCl}$.

Decomp. by cold H_2O . (Wheeler 3 RbCl , 2 HIO_3 . Sol. in H_2O , fr RbIO_3 separates. (Wheeler.)

Rubidium iodate selenate.

See Iodoselenate, rubidium.

Samarium iodate, $\text{Sm}(\text{IO}_3)_3 + 6\text{H}_2\text{O}$

Precipitate. (Cleve.)

ndium iodate, $\text{Sc}(\text{IO}_3)_3 + 10, 13, 15,$ and $18\text{H}_2\text{O}$.

Nearly insol. in H_2O . (Crookes, Phil. Mag. 1910, 210. A, 361.)

per iodate, AgIO_3 .

$.89 \times 10^{-4}$ moles or 5.36×10^{-3} g. AgIO_3 , sol. in 1 liter H_2O at 25° . (Noyes and Br, Z. phys. Ch. 1903, 42. 338.)

3l. sol. in H_2O . 4.35×10^{-2} g. are dissolved in 1 liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

1 l. H_2O dissolves 40 mg. AgIO_3 at 18° . (Ohlrausch, Z. phys. Ch. 1904, 50. 356.)

1 l. H_2O dissolves 0.0275 g. AgIO_3 at 9.43° ; 0.039 g. at 18.4° ; 0.0539 g. at 26.6° . Solubility increases rapidly with temp. (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

1 l. H_2O dissolves 0.039 g. AgIO_3 at 20° . (Hitby, Z. anorg. 1910, 67. 108.)

Not completely insol. in H_2O . (Rose.) Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; sol. in $\text{HNO}_3 + \text{Aq}$. (Naumann, J. B. 1860. 201.) Sol. in conc. $\text{KI} + \text{Aq}$. (Wendenburg, A. 185. 1.)

Sol. in 27,700 pts. H_2O at 25° ; in 42.4 pts. $\text{NH}_4\text{OH} + \text{Aq}$ at 25° ; in 2.1 pts. 10% $\text{NH}_4\text{OH} + \text{Aq}$ at 25° ; in 1044.3 pts. 35% $\text{NH}_4\text{OH} + \text{Aq}$ (sp. gr. 1.21) at 25° . (Longi, Z. ch. it. 18. 87.)

Solubility in $\text{HNO}_3 + \text{Aq}$ at 25° .

Normality HNO_3	G. AgIO_3 dissolved per l.
0.000	0.0503
0.125	0.0864
0.250	0.1075
0.500	0.1414
1.00	0.2067
2.00	0.3319
4.00	0.6985
8.00	1.5875

and Simmons, Z. phys. Ch. 1909, 67. 602.)

sol. in liquid NH_3 . (Gore, Am. Ch. J. 20. 829.)

sol. in methyl acetate. (Bezold, Dissert. 1906; Naumann, B. 1909, 42. 3790); acetate. (Hamers, Dissert. 1906; Naumann, B. 1910, 43. 314.)

r iodate ammonia, $2\text{AgIO}_3, 3\text{NH}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

ry sol. in cold H_2O . (Ditte, A. ch. (6) 15.)

$\text{IO}_3, 2\text{NH}_3$, sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Rosen-A. 1899, 308. 52.)

um iodate, NaIO_3 .

100 pts. H_2O dissolve 7.25 pts. NaIO_3 at 0° ; 9.07 pts. at 20° ; 14.39 pts. at 40° ; 17.7 pts. at 80° ; 33.9 pts. at 100° . (Krem-

ers, Pogg. 97. 5.) Sat. solution boils at 102° (Kremers), 105° (Ditte).

Sol. in warm $\text{H}_2\text{SO}_4 + \text{Aq}$ diluted with $\frac{1}{2}$ vol. H_2O . Crystallizes out on standing over H_2SO_4 . (Ditte.)

Solubility of NaIO_3 in $\text{HIO}_3 + \text{Aq}$ at 30° .

% HIO_3 in the solution	% NaIO_3 in the solution	Solid phase
0	9.36	$\text{NaIO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$
1.98	9.52	"
4.86	10.22	"
5.86	11.04	"
7.40	11.60	"
9.73	14.73	" } labile
6.76	11.18	$\text{NaIO}_3 + 1\frac{1}{2}\text{H}_2\text{O} + \text{Na}_2\text{O}, 2\text{I}_2\text{O}_5$
6.66	11.28	"
7.80	10.30	$\text{Na}_2\text{O}, 2\text{I}_2\text{O}_5$
9.15	9.00	"
9.93	8.71	"
11.20	7.54	"
11.89	7.21	$\text{Na}_2\text{O}, 2\text{I}_2\text{O}_5 + \text{NaIO}_3, 2\text{HIO}_3$
11.75	7.18	"
14.62	5.65	$\text{NaIO}_3, 2\text{HIO}_3$
23.23	3.69	"
32.68	2.91	"
40.91	2.64	"
46.62	2.67	"
55.48	2.12	"
65.47	1.83	"
76.19	1.42	$\text{NaIO}_3, 2\text{HIO}_3 + \text{HIO}_3$
76.70	0	HIO_3

(Meerburg, Z. anorg. 1905, 45. 334.)

Insol. in alcohol. Sol. in dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)
 $+ 1\frac{1}{2}\text{H}_2\text{O}$. See Meerburg above.

Sodium diiodate, $\text{Na}_2\text{O}, 2\text{I}_2\text{O}_5$.

See Meerburg under NaIO_3 .

Sodium triiodate, $\text{NaIO}_3, 2\text{HIO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Very sol. in H_2O . (Blomstrand, J. pr. (2) 42. 337.)
See also Meerburg under NaIO_3 .

Sodium iodate bromide, $\text{NaIO}_3, 2\text{NaBr} + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Rammelsberg.)

Sodium iodate chloride, $\text{NaIO}_3, \text{NaCl} + 4\text{H}_2\text{O}$, and $2\text{NaIO}_3, 3\text{NaCl} + 18\text{H}_2\text{O}$.

Cold H_2O dissolves out NaCl .

Sodium iodate iodide, $\text{NaIO}_3, \text{NaI}$.

Hot H_2O or alcohol dissolves out NaI .
 $+ 8\text{H}_2\text{O}$.
 $+ 10\text{H}_2\text{O}$.

$2\text{NaIO}_3, 3\text{NaI} + 20\text{H}_2\text{O}$. (Penny, A. 37. 202.)

Stable in a solution of $\text{NaI} + \text{NaOH} + \text{Aq.}$ (Eakle, C. C. 1896, II. 650.)

Strontium iodate, $\text{Sr}(\text{IO}_3)_2$.

Anhydrous. Insol. in H_2SO_4 (Ditte); easily sol. in cold $\text{HCl} + \text{Aq.}$ (Rammelsberg, Pogg. 44. 575.)

+ H_2O . Difficultly sol. in H_2O .

+ $6\text{H}_2\text{O}$. Sol. in 416 pts. H_2O at 15° , and 138 pts. at 100° (Gay-Lussac); 342 pts. at 15° , and 110 pts. at 100° . Difficultly sol. in warm $\text{HNO}_3 + \text{Aq.}$ (Rammelsberg, Pogg. 44. 575.)

Thallous iodate, TlIO_3 .

Difficultly sol. in warm H_2O . (Oettinger.)

Insol. in H_2O ; difficultly sol. in $\text{HNO}_3 + \text{Aq.}$ (Rammelsberg.)

Sl. sol. in H_2O .

0.58×10^{-1} g. are dissolved in 1 liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

2.12×10^{-3} mols. = 0.667 g. are sol. in 1 l. H_2O at 25° . (Spencer, Z. phys. Ch. 1912, 80. 707.)

Sol. in a little $\text{NH}_4\text{OH} + \text{Aq.}$ also in boiling HNO_3 , H_2SO_4 , or $\text{HCl} + \text{Aq.}$ Insol. in alcohol. (Oettinger.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Very sl. sol. in H_2O or dil. boiling acids. (Ditte, A. ch. (6) 21. 145.)

Thallic iodate, basic, $\text{Tl}(\text{OH})(\text{IO}_3)_2 + \text{H}_2\text{O} = \text{Tl}_2\text{O}_3, 2\text{I}_2\text{O}_5 + 3\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in cold $\text{HCl} + \text{Aq.}$ and warm dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Ditte, A. ch. (6) 21. 145.)

Thallic iodate, $\text{Tl}(\text{IO}_3)_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O ; sl. sol. in $\text{HNO}_3 + \text{Aq.}$ Decomp. by alkalis. (Rammelsberg.)

+ $12\text{H}_2\text{O}$. Difficultly sol. in H_2O .

Easily sol. in dil. acids. (Gewecke, Z. anorg. 1912, 75, 275.)

Thorium iodate, $\text{Th}(\text{IO}_3)_4$.

Precipitate. (Cleve.)

Tin (stannous) iodate.

Ppt. Sol. in $\text{SnCl}_2 + \text{Aq.}$; insol. in $\text{NaIO}_3 + \text{Aq.}$

Tin (stannic) iodate.

Ppt.

Uranous iodate.

Precipitate. Very unstable. (Rammelsberg.)

Uranyl iodate, $\text{UO}_2(\text{IO}_3)_2$.

Sol. or insol. in HNO_3 and $\text{H}_3\text{PO}_4 + \text{Aq.}$ according to method of preparation. (Ditte.)

+ H_2O . Sl. sol. in $\text{HNO}_3 + \text{Aq.}$ (Rammelsberg.)

Ytterbium iodate, $\text{Yb}(\text{IO}_3)_3 + 6\text{H}_2\text{O}$.

Ppt. (Cleve, Z. anorg. 1902, 32. 136.)

Yttrium iodate, $\text{Y}(\text{IO}_3)_3 + 3\text{H}_2\text{O}$.

Sol. in 190 pts. H_2O . (Berlin.)

Zinc iodate, $\text{Zn}(\text{IO}_3)_2$.

Anhydrous. (Ditte, A. ch. (6) 21. 145.)

+ $2\text{H}_2\text{O}$. Sol. in 114 pts. cold, and 76 pts. hot H_2O . (Rammelsberg, Pogg. 43. 665.)

Sol. in HNO_3 , and $\text{NH}_4\text{OH} + \text{Aq.}$

Exists also in a very sol. modification (Mylius and Funk, B. 1897, 30. 1723.)

Zinc iodate ammonia, $3\text{Zn}(\text{IO}_3)_2, 8\text{NH}_3$.

Decomp. by H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ from which it is pptd. by alcohol. (Rammelsberg, Pogg. 44. 563.)

$\text{Zn}(\text{IO}_3)_2, 2\text{NH}_3$. Insol. in H_2O . (Ditte, A. ch. (6) 21. 145.)

$\text{Zn}(\text{IO}_3)_2, 3\text{NH}_3 + \text{H}_2\text{O}$. Insol. in H_2O . (Ditte.)

$\text{Zn}(\text{IO}_3)_2, 4\text{NH}_3$. (Ditte, A. ch. 1890, 21. 164.) (Ephraim, B. 1915, 48. 53.)

Periodic acid.

See Periodic acid.

Iodides.

The iodides are in general easily sol. in H_2O ; exceptions are HgI_2 , PbI_2 , AgI , CuI , and BiI_3 , also the iodides of the Pt metals, all of which are insol. SnI_4 , SbI_3 , and TlI_3 decomp. by H_2O . Many iodides are more sol. in solutions of salts than in H_2O , and some are sol. in alcohol or ether.

See under each element.

Iodine, I_2 .

Sol. in 5524 pts. H_2O at $0-12^\circ$. (Wittstein, Z. anorg. 1897, 123.)

Sol. in 7000 pts. H_2O . (Gay-Lussac.)

Sol. in 3800 pts. H_2O at 15° . (Bassé.)

Sol. in 5000 pts. H_2O . (Jacquelin.)

Sol. in 7196.4 pts. B_2O at 18.75° . (Abl.)

Pure H_2O dissolves 0.01519173 g. I_2 per litre, or I_2 is sol. in 6582 pts. H_2O at 15° . (Dossius and Weith, Zeit. Ch. 12. 378.)

Sol. in about 4500 pts. H_2O . (H. Comm. 1883.)

Sol. in 7000 pts. H_2O . (Cap and Gar, Pharm. (3) 26. 80.)

1 l. H_2O at 25° dissolves 0.3387 g. I_2 . (Jakowkin, Z. phys. Ch. 1895, 18. 590.)

1 l. H_2O dissolves 1.342 millimoles of I_2 at 25° . (Noyes, Z. phys. Ch. 1898, 27. 1.)

When iodine is shaken with H_2O at 1 pt. dissolves in 3750 pts. H_2O ; when I_2 and H_2O are heated together and then cooled to 15° , 1 pt. iodine dissolves in 3500 pts. H_2O .

At 30° , 1 pt. is sol. in 2200 pts. H_2O . (Dietz, Chem. Soc. 1899, 76, (2) 150.)

1 l. H_2O dissolves 0.279 grams I_2 at 15° . (McLauchlan, Z. phys. Ch. 1903, 44. 1.)

ubility of I₂ in H₂O at t°.

	g. I ₂ per l. H ₂ O
	0.2765
	0.3395
	0.4661
	0.6474
	0.9222

y, Chem. Soc. 1908, 93. 744.)

ubility of I₂ in H₂O at t°.

g. per l.	milliat. per l.
0.1649	1.30
0.2941	2.30
0.5684	4.56

tieff, Z. anorg. 1910, 69. 30.)

mol I₂ are sol. in 1 l. H₂O. (Bray' m. Soc., 1910, 32. 938.)
d from electrical conductivity of 1 l. H₂O dissolves 0.0006383 mols. ones, J. Am. Chem. Soc. 1915, 37.

SO₄, HCl, HNO₃, H₃PO₄, tartaric, or citric acids+Aq dist give it up to CS₂ on shaking (Tessier, Z. anal. 11. 313.)
50 pts. H₂SO₄ on warming, but out in part on cooling. (Kraus.)
re sol. in HBr+Aq than in pure +Aq of sp. gr. 1.486 dissolves neu.)
HCl+Aq. Easily sol. in even l.
N-HCl+Aq sat. with I₂ contains (Bray and Mackay, J. Am. Chem. 2. 1919.)
-HNO₃+Aq sat. with I₂ contains (Sammet, Z. phys. Ch. 1905, 53.
-H₂SO₄+Aq sat. with I₂ contains (Sammet.)
SO₃+Aq with decomp.
J. H₂BO₃ dissolves 0.300 g. I₂ at auchlan, Z. phys. Ch. 1903, 44.

f a 10% solution of BaBr₂ dis- g. I₂ at 13.5°. (Meyer, Z. anorg. 4.)
a 10% solution of BaCl₂ dissolve t 18.5°. (Meyer.)
a 10% solution of BaI₂ dissolve t 13.5°. (Meyer.)
f a 10% solution of CaBr₂ dis- g. I₂ at 13.5°. (Meyer.)
a 10% solution of CaCl₂ dissolve t 18.5°. (Meyer.)
a 10% solution of CaI₂ dissolve t 13.5°. (Meyer.)
l. in boiling dil. HgCl₂+Aq.

Solubility in HgCl₂+Aq at 25°.

10 ccm. of the solution contain:	
millimols I ₂	millimols Hg
0.0134	0
0.1294	0.9444
0.1460	1.2442
0.1806	1.9542
0.2543	3.3460

(Hers and Paul, Z. anorg. 1914, 85. 214.)

Sol. in solutions of soluble iodides.
100 pts. KI+200 pts. H₂O dissolve 153 pts. I; from this solution H₂O precipitates 1/2 the dissolved I. 100 pts. KI+400 pts. H₂O dissolve quickly 76.5 pts. I. If more water is present, the solution takes place more slowly. (Baup.)
CS₂ extracts the I from the above solutions.

Solubility of I in KI+Aq at 7-7.3°.

% KI in KI+Aq	Pts I dissolved	Sp. gr. of solution
1.802	1.173	1.0234
3.159	2.303	1.0433
4.628	3.643	1.0668
5.935	4.778	1.0881
7.201	6.037	1.1112
8.663	7.368	1.1382
10.036	8.877	1.1637
11.034	9.949	1.1893
11.893	11.182	1.2110
12.643	12.060	1.2293

(Dossius and Weith, Zeit. Ch. (2) 5. 379.)

Solubility of I₂ in KI+Aq at room tempera- ture, 14.5°-15.1°.

% KI	% I	I/KI
1.80	1.17	0.651
3.16	2.30	0.729
4.63	3.64	0.786
5.93	4.78	0.805
7.20	6.04	0.839
8.66	7.37	0.851
10.04	8.88	0.884
11.03	9.95	0.902
11.89	11.18	0.940
12.64	12.06	0.954

(Weith and Dossius, Z. phys. Ch. 1898, 26. 150.)

Solubility of I ₂ in KI+Aq at 15°		
% KI	ccm. 1/10-n. iodine in 5cm. of the solution	I/KI
10	35.0	35.0
8	27.1	33.9
6	19.7	32.8
4	12.7	31.8
2	6.25	31.2
1	3.04 *	30.4

* Obtained with 1/100-normal iodine.
(Bruner, Z. phys. Ch. 1898, 26. 151.)

Solubility of I ₂ in KI+Aq at 25°.	
Millimols KI per liter	Millimols dissolved iodine per liter
106.3	55.28
53.15	28.03
26.57	14.68
13.29	8.003
6.643	4.667
3.322	3.052
1.661	2.235
0.8304	1.814

(Noyes and Seidenstricker, Z. phys. Ch. 1898, 27. 359.)

Solubility in KI+Aq at 25°.	
KI mol./l.	I G. atoms/l.
1.91	3.29
2.85	5.45
4.51	11.52
5.36	17.12
5.55	17.16

(Abegg, Z. anorg. 1906, 50. 427.)

Solubility of I ₂ in KI+Aq at 25°.	
Millimol KI per l.	Millimol I ₂ dissolved
100	51.35
50	25.77
20	11.13
10	6.185
5	3.728
2	2.266
1	1.788

(Bray and MacKay, J. Am. Chem. Soc. 1910, 32. 919.)

Solubility in KI+Aq at 25°.				
Sp. gr.	Analysis of liquid phase		Analysis of solid phase together with adhering mother liquor	
	% KI	% I	% KI	% I
(a) In equilibrium with excess of KI.				
1.733	60.39	0.0		0.0
1.888	54.415	11.63	84.92	4.05
2.066	49.045	23.085	85.94	6.32
2.216	44.82	31.01	80.46	10.84
2.539	38.065	44.56	78.56	15.23
2.560	37.655	45.55	77.32	16.73
2.665	35.805	49.61		
3.232	29.71	62.81	39.99	56.10
3.246	27.92	66.45	38.78	56.27
(b) In equilibrium with excess of I.				
1.349	16.025	18.49	3.04	85.43
1.516	19.705	26.16	4.48	83.87
1.769	22.88	36.06	3.70	89.33
1.910	23.55	40.515	6.49	83.62
2.403	24.78	53.605	8.62	83.81
2.904	24.995	63.125	4.82	92.41
3.082	25.18	66.04	4.00	94.39
(c) Invariant point. Excess of KI and I.				
3.316	26.05	68.06		
	25.96	68.01	16.14	83.77
	26.04	68.16		
	25.92	68.13	11.32	86.56

(Parsons and Whittemore, J. Am. Chem. 1911, 33. 1934.)

Solubility in KI+Aq at 0°.			
KI +Aq		KI +Aq. sat. with I ₂	
Wt. norm.	Sp. gr. 0°/4°	G. I ₂ in 1 g. of solution	Sp. gr. 0°/4°
0.09871	(1.0123)	0.01199	(1.0219)
0.09861	1.01231	0.01199	1.02187
0.04969	(1.0061)	0.006094	(1.0109)
0.04966	1.00610	0.006083	1.01089
0.01992	1.00236	0.002535	1.00429
0.01983	(1.0024)	0.0025325	(1.0044)
0.00998	(1.0011)	0.0013532	(1.0020)
0.00992	(1.0011)	0.0013585	(1.0020)
0.004999	(1.0005)	0.0007609	(1.0010)
0.004991	(1.0005)	0.0007577	(1.0011)
0.002000	(1.0001)	0.0004137	(1.0004)
0.002000	(1.0001)	0.0004015	(1.0004)
0.000999	(0.9999)	0.0002839	(1.0002)
0.000992	(1.0000)	0.00028125	(1.0002)

Values in parentheses are found by interpolation.
(Jones and Hartman, J. Am. Chem. Soc. 1915, 37. 247.)

1 mol. KI in alcohol dissolves 2 atoms I, and the solution does not give up I to CS₂.
(Jørgensen, J. pr. (2) 2. 347.)

Solubility in KI+60% alcohol at 25°.

T.	Analysis of liquid phase		Analysis of solid phase together with adhering mother liquor	
	% KI	% I	% I	% KI
(a) In equilibrium with excess KI.				
8	30.93	0.0		0.0
1	29.87	4.51	89.13	0.71
5	28.39	12.48	86.60	2.27
8	28.00	18.60	87.30	3.21
7	27.60	21.80	85.75	4.25
3	27.00	28.00	84.39	6.05
6	25.90	40.52	81.05	10.30
0	24.90	52.42	76.21	16.73
7	24.40	58.93	73.20	21.04
5	22.49	65.75	71.66	24.15
	21.50	68.95	70.04	26.42

(b) In equilibrium with excess I.				
4	0.0	23.04	0.0	I
0	7.36	43.05	1.40	88.76
1	10.60	49.38	2.50	88.21
	12.44	55.33	3.72	87.10
	13.74	59.26	4.41	86.60
	15.20	62.66	5.80	85.20
	17.72	69.10	7.15	85.49
	19.30	71.90	7.45	88.96

(c) Invariant point. Excess KI and I.				
2	20.11	72.51
	20.03	72.46	21.84	74.64
				KI+I
	20.05	72.54
	19.98	72.44	7.40	89.81 I
	20.08	72.51	20.61	74.09
				KI+I
	20.06	72.44
	20.05	72.48	33.46	63.19 KI

(Parsons and Corliss, J. Am. Chem. Soc. 1910, 32. 1370.)

Solubility in KI+40% alcohol at 25°.

T.	Analysis of liquid phase		Analysis of solid phase together with adhering mother liquor.	
	% KI	% I	% KI	% I.
(a) In equilibrium with excess KI.				
39	42.10	0.0		0.0
77	40.83	3.76	89.21	0.70
55	38.94	10.09	88.80	1.90
32	37.41	15.71	88.19	3.02
05	36.25	20.52	87.04	4.21
55	35.38	24.44	86.08	5.11
47	33.26	33.62	83.61	8.41
24	31.71	39.99	82.06	10.76
69	30.59	44.76	80.80	12.35
58	28.56	55.30	75.90	18.63
84	26.95	60.27	74.77	20.86
.	24.52	65.93	72.98	23.61
.	23.04	69.93	72.45	25.04

Solubility in KI+40% alcohol at 25°.—
Continued.

Sp. gr.	Analysis of liquid phase		Analysis of solid phase together with adhering mother liquor	
	% KI	% I	% KI	% I
(b) In equilibrium with excess I.				
0.962	0.0	2.97	0.0	
1.292	8.45	28.70	1.85	84.51
1.581	12.56	40.63	3.41	84.02
...	15.20	49.95	4.98	83.81
2.000	16.02	52.95	5.60	82.96
2.173	17.18	57.38	6.61	83.60
1.749	19.20	66.89	8.45	85.16
2.902	20.12	69.10	7.08	88.81
(c) Invariant point. Excess KI and I.				
3.246	22.50	70.79	19.48	76.24
...	22.43	70.88	69.37	26.14

(Parsons and Corliss, J. Am. Chem. Soc. 1910, 32. 1372.)

See also under KI.

Sol. in KI+nitrobenzene. (Dawson, Chem. Soc. 1902, 81. 529.)

Solubility in KIO₃+Aq is the same as in H₂O. (Lami, C. A. 1909. 1622.)

Solubility of I₂ in KBr+Aq at 25°.

G. KBr per l.	G. atoms I ₂ per l.
60.6	0.0176
106.9	0.0278
175.9	0.0415
229.8	0.0532
281.9	0.0628
330.6	0.0717
377.1	0.0797
411.0	0.0864
461.7	0.0948
509.8	0.1006
548.0	0.1062
567.9 sat.	0.1094

(Bell and Buckley, J. Am. Chem. Soc. 1912, 34. 13.)

Solubility in NaBr+Aq at 25°.

G. NaBr per l.	G. atoms I ₂ per l.
96.4	0.0266
187.7	0.0425
271.8	0.0538
357.4	0.0598
422.4	0.0638
499.1	0.0648
569.9	0.0644
632.0	0.0622
679.7	0.0595
750.5	0.0551
756.1 sat.	0.0550

(Bell and Buckley, J. Am. Chem. Soc. 1912, 34. 13.)

100 cc. of a 10% solution of SrBr_2 dissolve 0.270 g. I_2 at 13.50. (Meyer, Z. anorg. 1902, 30. 114.)
100 cc. of a 10% solution of SrCl_2 dissolve 0.066 g. I_2 at 18.5°. (Meyer.)
100 cc. of a 10% solution of SrI_2 dissolve 6.616 g. I_2 at 13.5°. (Meyer.)

Solubility in salts + Aq at 25°.

Salt + Aq	Grams I_2 sol. in 1 liter	Salt + Aq	Grams I_2 sol. in 1 liter
$\frac{1}{2}\text{N. Na}_2\text{SO}_4$	0.160	N. NaCl	0.575
$\frac{1}{2}\text{N. K}_2\text{SO}_4$	0.238	N. KCl	0.658
$\frac{1}{2}\text{N. (NH}_4)_2\text{SO}_4$	0.246	N. NH_4Cl	0.735
N. NaNO_3	0.257	N. NaBr	3.29
N. KNO_3	0.266	N. KBr	3.801
N. NH_4NO_3	0.375	N. NH_4Br	4.003

(McLauchlan, Z. phys. Ch. 1903, 44. 617.)

1.14 g. are sol. in 100 ccm. liquid H_2S . (Antony, Gazz. ch. it. 1905, 35, (1) 206.)
Sol. in liquid NH_3 . (Franklin, Am. ch. J. 1898, 20. 822.)
Sl. sol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54. 674.)
Sol. in liquid SO_2 (Sestini), and SO_2 (Weber).
100 pts. AsCl_3 dissolve 8.42 pts. I at 0°; 11.88 pts. I at 15°; 36.89 pts. I at 96°. (Sloan, C. N. 46. 194.)
Sol. in liquid SO_2 , AsCl_3 , SO_2Cl_2 , and acetaldehyde. (Walden, Z. phys. Ch. 1903, 43. 407.)
Very sol. in liquid NO_2 . (Frankland, Chem. Soc. 1901, 79. 1361.)
Sol. in 10–12 pts. alcohol. (Wittstein.)
Sol. in wood-spirit. (Playfair.)
Abundantly sol. in amyl (Pelletan), and hexyl alcohol (Bouis).
Iodine is sol. in 20 pts. alcohol, 110 pts. oil, 7000 pts. H_2O , 100 pts. glycerine. (Cap and Garot, J. Pharm. (3) 26. 80.)

Solubility of I_2 in $\text{C}_2\text{H}_5\text{OH} + \text{Aq}$ at room temperature (14.5°–15.1°).

Volumes of $\text{C}_2\text{H}_5\text{OH}$ in 100 volumes of $\text{C}_2\text{H}_5\text{OH} + \text{H}_2\text{O}$	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	61.7
90	29.4
80	16.6
70	9.2
60	4.45
50	3.4
40	1.0
30	0.4
20	0.25
10	0.2
0	0.0

(Bruner, Z. phys. Ch. 1898, 26. 150.)

Solubility of I_2 in $\text{C}_2\text{H}_7\text{OH} + \text{Aq}$ at room temperature (14.5°–15.1°).

Volumes of $\text{C}_2\text{H}_7\text{OH}$ in 100 volumes of $\text{C}_2\text{H}_7\text{OH} + \text{H}_2\text{O}$	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	58.8
90	36.0
80	23.6
70	16.1
60	10.7
50	6.4
40	3.7
30	1.56
20	0.42
10	0.19
0

(Bruner, Z. phys. Ch. 1898, 26. 150.)

Solubility in ethyl alcohol + Aq at 25°.

Molecules of $\text{C}_2\text{H}_5\text{OH}$ in 100 molecules $\text{C}_2\text{H}_5\text{OH} + \text{H}_2\text{O}$	Molecules of H_2O in 100 molecules $\text{C}_2\text{H}_5\text{OH} + \text{H}_2\text{O}$	Normality of the iodine solution
0.0	100	0.0022
0.03	99.7(?)	0.0024
0.06	99.4(?)	0.0024
1.12	98.88	0.0023
1.83	98.27(?)	0.0025
9.40	90.60	0.0059
13.48	86.52	0.0111
23.80	76.20	0.0617
50.80	49.20	0.4326
100	0	1.590

(McLauchlan, Z. phys. Ch. 1903, 44. 627.)

Solubility in acetic acid + Aq at 25°.

Molecules of CH_3COOH in 100 molecules $\text{CH}_3\text{COOH} + \text{H}_2\text{O}$	Molecules of H_2O in 100 molecules $\text{CH}_3\text{COOH} + \text{H}_2\text{O}$	Normality of the iodine solution
0.0	100	0.0022
6.98	93.02	0.0049
16.40	83.60	0.0112
31.90	68.10	0.0331
55.70	44.30	0.0882
100	0	0.205

(McLauchlan, Z. phys. Ch. 1903, 44. 627.)

Very sol. in ether, chloroform, and bromo form.

Solubility in ether.

100 g. of the sat. solution contain at:

—83° —90° —108°
15.39 14.58 15.09 g. I_2 .

(Arctowski, Z. anorg. 1896, 11. 276.)

About as sol. in all fatty oils as in CHCl_3 etc. (Gruel, Arch. Pharm. 223. 431.)

in 56.6 pts. chloroform at 10°. (Dun-
harm. J. Trans. 51. 544.)
ibility in CHCl₃.
g. of the sat. solution contain at:
—55.5° —60° —69.5° —73.5°
0.144 0.129 0.089 0.080 g. I₂.
Lretowski, Z. anorg. 1896, 11. 276.)
y sol. in methylene iodide. (Retgers,
org. 3. 343.)

ibility of I₂ in C₆H₆+CHCl₃ at room
temperature (14.5°—15.1°).

es of C ₆ H ₆ in 100 s of C ₆ H ₆ +CHCl ₃	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	41.05
90	38.8
80	34.6
70	30.5
60	27.4
50	24.4
40	21.0
30	19.2
20	17.8
10	16.0
0	14.3

Bruner, Z. phys. Ch. 1898, 26. 147.)

bility of I₂ in CS₂+CHCl₃ at room
temperature (14.5°—15.1°).

mes of CS ₂ in 100 mes of CS ₂ +CHCl ₃	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	69.4
90	62.7
80	55.9
70	47.9
60	42.0
50	35.8
40	30.4
30	25.3
20	20.8
10	17.0
0	14.3

(Bruner.)

olubility of I₂ in C₆H₆+CCl₄ at room
temperature (14.5°—15.1°).

mes of C ₆ H ₆ in 100 mes of C ₆ H ₆ +CCl ₄	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	41.05
90	37.2
80	33.6
70	29.6
60	26.1
50	22.4
40	19.25
30	16.1
20	13.4
10	10.75
0	8.1

(Bruner.)

Solubility of I₂ in CS₂+CCl₄ at room
temperature (14.5°—15.1°).

Volumes of CS ₂ in 100 volumes of CS ₂ +CCl ₄	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	69.1
90	56.9
80	48.6
70	40.7
60	33.9
50	26.9
40	21.8
30	17.7
20	13.25
10	10.2
0	8.1

(Bruner.)

Solubility of I₂ in C₂H₅OH+CHCl₃ at room
temperature (14.5°—15.1°).

Volumes of C ₂ H ₅ OH in 100 volumes of C ₂ H ₅ OH + CHCl ₃	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	61.7
90	37.1
80	34.2
70	30.7
60	27.9
50	26.1
40	24.6
30	22.7
20	19.9
10	17.1
0	14.25

(Bruner.)

Solubility of I₂ in C₂H₇OH+CHCl₃ at room
temperature (14.5°—15.1°).

Volumes of C ₂ H ₇ OH in 100 volumes of C ₂ H ₇ OH +CHCl ₃	Ccm. of 1/10-normal iodine in 5 cc. of the solution
100	58.8
90	51.9
80	44.2
70	35.4
60	31.8
50	30.8
40	27.9
30	25.3
20	21.8
10	17.8
0	14.25

(Bruner.)

Sol. in acetone. (Naumann, B. 1904, 37.
4328); (Eidmann, C. C. 1999, II. 1014.)
Sol. in methyl acetate (Naumann, B. 1909,
42. 3790); ethyl acetate. (Naumann, B.
1904, 37. 3601.)
Sol. in allyl mustard oil, phenyl mustard

Partition of I₂ between CHCl₃ and other solvents.
C = millimols iodine in 10 ccm. of the CHCl₃ layer.
W = millimols iodine in 10 ccm. of the other layer.

Other Solvent	C	W	C/W
Water	0.338	0.0025	134.6
	1.546	0.0120	129.0
	2.318	0.0184	126.3
	3.207	0.0242	132.8
	3.439	0.0259	132.8
75% by vol. H ₂ O + 25% by vol. glycerine	1.217	0.0183	66.32
	1.893	0.0290	65.33
	2.434	0.0367	66.31
	3.219	0.0483	66.65
50% by vol. H ₂ O + 50% by vol. glycerine	1.217	0.0405	30.0
	1.835	0.0609	30.1
	2.376	0.0782	30.4
	3.294	0.1020	32.2
25% by vol. H ₂ O + 75% by vol. glycerine	1.188	0.116	10.25
	1.806	0.173	10.45
	2.656	0.249	10.66
	2.859	0.265	10.80
	3.400	0.312	10.93

(Herz, Z. Elektrochem. 1910, 16. 870.)

Distribution of I₂ between benzene and glycerine at t°.
M₁ = concentration of I₂ in benzene layer expressed in g.-mol. per l.
M₂ = concentration of I₂ in glycerine layer expressed in g.-mol. per l.

t°	M ₁	M ₂
25°	0.00757	0.001604
	0.01610	0.002664
	0.02719	0.004115
	0.04024	0.005794
	0.06255	0.00834
	0.07923	0.01033
	0.10243	0.01324
	0.12201	0.01559
	0.13342	0.01668
	0.16734	0.02081
40°	0.008545	0.00181
	0.01544	0.002593
	0.04432	0.006242
	0.095004	0.012013
	0.13271	0.01632
	0.18508	0.02193
50°	0.00865	0.00184
	0.01523	0.00253
	0.02683	0.00390
	0.04413	0.00576
	0.0620	0.00744
	0.07832	0.00942
	0.10153	0.01214
	0.12166	0.0145
	0.13199	0.01560
	0.18438	0.02122

(Landau, Z. phys. Ch. 1910, 73. 202.)

Distribution of I₂ between glycerine and CCl₄ at t°.
M₁ = concentration of I₂ in CCl₄ layer expressed in g.-mol. per l.
M₂ = concentration of I₂ in glycerine layer expressed in g.-mol. per l.

t°	M ₁	M ₂
25°	0.002230	0.0014386
	0.0024113	0.0014506
	0.0048227	0.0027014
	0.010452	0.005581
	0.038973	0.019959
	0.04598	0.023948
	0.05820	0.030097
40°	0.00227	0.00127
	0.00239	0.00138
	0.00461	0.00272
	0.01092	0.00482
	0.02540	0.01116
	0.04091	0.01749
	0.06074	0.02701
50°	0.00257	0.00118
	0.00500	0.00225
	0.01363	0.00596
	0.02549	0.01050
	0.04167	0.01693
	0.06309	0.02502

(Landau, Z. phys. Ch. 1910, 73. 203.)

Distribution of I₂ between ether and ethylglycol at t°.
M₁ = concentration of I₂ in ether layer expressed in g.-mol. per l.
M₂ = concentration of I₂ in C₂H₅O₂ layer expressed in g.-mol. per l.

t°	M ₁	M ₂
0°	0.00843	0.00571
	0.03082	0.01713
	0.06551	0.03736
	0.08105	0.04606
	0.12528	0.07148
	0.31511	0.17524
25°	0.00870	0.00571
	0.01677	0.01001
	0.02710	0.01586
	0.03046	0.01713
	0.06385	0.03594
	0.11951	0.06726
	0.30820	0.17524

(Landau, Z. phys. Ch. 1910, 73. 206.)

Iodine monobromide, IBr.
Slowly sol. in H₂O with slight decol.
Sol. in CHCl₃, CS₂, ether, and alcohol.
+5H₂O. (Löwig, Pogg. 14. 485.) I not exist. (Bornemann, A. 189. 183.)

pentabromide, $\text{IBr}_5(?)$.
in H_2O with separation of iodine.
, Pogg. 14. 485.)

monochloride, ICl .

mp. by H_2O ; sol. without decomp. in
, ether, and $\text{HCl} + \text{Aq.}$
in CS_2 .

hydrogen chloride, ICl , HCl .

able. Sol. in ether. (Schützenberger,
14. 389.)

trichloride, ICl_3 .

uescent. With H_2O , a part is dissolved
t decomp., and the rest is decomp.
queous solution contains more und
d ICl_3 , the more conc. it is. (Serullas.)
tated from aqueous solution by H_2SO_4 .
 $\text{HCl} + \text{Aq.}$ Sol. in warm conc. H_2SO_4 .
t decomp. Sol. in alcohol, and ben-
Decomp. by small amount of CS_2 .
omanos, B. 10. 434.) Ether does not
it from aqueous solution. (Serullas.)

lithium chloride, ICl_3 , $\text{LiCl} + 4\text{H}_2\text{O}$.

lithium chloriodide.

trichloride magnesium chloride, 2ICl_3 ,
 $\text{MgCl}_2 + 5\text{H}_2\text{O}$.

deliquescent and easily decomposed.
, J. Pharm. 25. 442.)

I_2O . Hydroscopic. (Weinland, Z.
1902, 30. 141.)

trichloride manganous chloride, 2ICl_3 ,
 $\text{MnCl}_2 + 8\text{H}_2\text{O}$.

rosopic. (Weinland, Z. anorg. 1902,
1.)

trichloride nickel chloride, 2ICl_3 ,
 $\text{NiCl}_2 + 8\text{H}_2\text{O}$.

rosopic. CCl_4 dissolves out ICl_3
and, Z. anorg. 1902, 30. 138.)

monochloride phosphorus pentachloride,
 ICl , PCl_5 .

r deliquescent; decomp. by H_2O .

potassium chloride, ICl_3 , KCl .

in H_2O with decomp.
x dissolves out ICl_3 . (Filhol, J. Pharm.
3, 506.)

Potassium chloriodide.

sodium chloride, ICl_3 , $\text{NaCl} + 2\text{H}_2\text{O}$.

Sodium chloriodide.

trichloride strontium chloride, 2ICl_3 ,
 $\text{SrCl}_2 + 8\text{H}_2\text{O}$.

rosopic. (Weinland, Z. anorg. 1902,
2.)

Iodine trichloride sulphur tetrachloride, ICl_3 ,
 8Cl_4 .

Very deliquescent in air; decomp. by H_2O .
Decomp. with formation of clear solution by
dil. $\text{HNO}_3 + \text{Aq.}$ (Weber, Pogg. 128. 459.)

SCl_2 , 2ICl_3 . (Jaillard, J. B. 1860. 95.)

Correct formula is as above. (Weber, l. c.)

2ICl_3 , SCl_4 . Sol. in SO_2Cl_2 , SOCl_2 , POCl_3 ,
warm SCl_2 , petroleum ether, ligroin, CHCl_3 ,
 CCl_4 , CS_2 , and abs. ether. (Ruff, B. 1904, 37.
4519.)

Iodine trichloride zinc chloride, 2ICl_3 , $\text{ZnCl}_2 +$
 $8\text{H}_2\text{O}$.

Unstable. Hydroscopic. (Weinland, Z.
anorg. 1902, 30. 140)

Iodine pentafluoride, IF_5 .

Fumes in air; decomp. with H_2O . (Gore,
C. N. 24. 291.)

Decomp. by H_2O into iodic acid and HF .
Decomp. by solutions of the alkalies. (Mois-
san, C. R. 1902, 136. 564.)

Iodine trioxide, I_2O_3 .

Decomp. by H_2O . (Ogier, C. R. 85. 957;
86. 722.)

Probably a mixture.

Iodine tetroxide, $\text{I}_2\text{O}_4(?)$.

Insol. in cold, decomp. by hot H_2O ; insol.
in alcohol. Decomp. by $\text{HNO}_3 + \text{Aq.}$ Sol. in
 H_2SO_4 . (Millon, J. pr. 34. 319, 337.)

Iodine pentoxide, I_2O_5 .

Very sol. in H_2O , and in dil. alcohol.
Insol. in absolute alcohol, ether, CS_2 , chloro-
form, and hydrocarbons.

Forms hydrates, iodic acid HIO_3 , and
 $3\text{I}_2\text{O}_5$, H_2O ; insol. in ordinary alcohol.

For sp. gr. of aqueous solution, see *iodic*
acid

Iodine oxides, $\text{I}_{10}\text{O}_{19}$, I_8O_{13} .

The compounds, $\text{I}_{10}\text{O}_{19}$ (Millon, J. pr. 34.
336), and I_8O_{13} (Kämmerer, J. pr. 83. 81),
are probably mixtures.

Millon's oxides are impure I_2O_5 . (Kap-
peler, B. 1911, 44. 3496.)

Iodine sulphur oxide, $5\text{I}_2\text{O}_5$, 8SO_3 .

Decomp. by H_2O . (Kämmerer.)

I_2O_5 , 3SO_3 . Decomp. by H_2O ; al. sol. in
hot SO_4 . (Weber, B. 20. 86.)

$= (\text{IO})_2(\text{SO}_4)_3$. Iodyl sulphate (?).

Iodine oxyfluoride, $\text{IOF}_3 + 5\text{H}_2\text{O}$.

Fumes in the air. (Weinland, Z. anorg.
1908, 60. 163.)

Iodine sulphide, S_2I_2 .

Sol. in CS_2 . (Linebarger, Am. Ch. J. 1895,
17. 57.)

Iodine sulphoxide, $I_2SO_2(?)$.

Decomp. by H_2O . (Schultz-Sellack.)
 $I_2(SO_2)_2(?)$. Decomp. by H_2O . (Weber, J. pr. (2) 25. 224.)
 $I_2(SO_2)_2(?)$. As above. (Weber.)
 See also Iodosulphuric anhydride.

Iodiridic acid.**Ammonium iodiridate, $(NH_4)_2IrI_4$.**

Very easily sol. in cold H_2O , decomp. on warming. Insol. in alcohol. (Oppler, J. B. 1857. 263.)

Potassium iodiridate, K_2IrI_4 .

Very easily sol. in H_2O . Insol. in alcohol.

Sodium iodiridate, Na_2IrI_4 .

Insol. in cold, sl. sol. in hot H_2O . Easily sol. in acids. (Oppler.)

Iodiridous acid.**Ammonium iodiridite, $(NH_4)_2Ir_2I_{12} + H_2O$.**

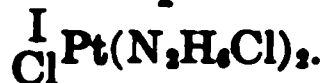
Very sol. in H_2O , but decomp. on warming. (Oppler.)

Potassium iodiridite, $K_2Ir_2I_{12}$.

Insol. in H_2O , or alcohol. Slowly sol. in acids; easily in warm alkalies + Aq.

Silver iodiridite, $Ag_2Ir_2I_{12}$.

Ppt.

Iodochloroplatindiamine chloride,

Sl. sol. in H_2O .

Iodochromic acid.**Potassium iodochromate, $KCrO_3I$.**

Decomp. by boiling H_2O . (Guyot, C. R. 73. 46.)

See also Chromiodic acid.

Iodomolybdic acid.

See Molybdoiodic acid.

Iodonitratoplatinmonodiamine bromide,

Very sl. sol. in H_2O . (Cleve.)

Iodonitritoplatindiamine nitrate,

Quite easily sol. in hot H_2O . (Cleve.)

Iodopalladous acid.**Potassium iodopalladite.**

Deliquescent. (Lassaigne.)

Iodophosphoric acid.

See Phosphiodic acid.

Iodoplatinamine iodide, $I_2Pt(NH_5I)_2$.

Sol. in H_2O , especially easily if boil (Cleve.)

Iodoplatindiamine iodide, $I_2Pt(N_2H_5I)_2$.

Sol. in H_2O , especially when hot. (Cleve.)

— mercuric iodide, $I_2Pt(N_2H_5I)_2, 2HgI_2$.

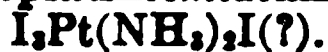
Extremely difficultly sol. in cold H_2O , partly decomp. by boiling. (Jørgensen, K. 3. 1214.)

— nitrate, $I_2Pt(N_2H_5NO_2)_2$.

More sol. in hot than cold H_2O .

— sulphate, $I_2Pt(N_2H_5)_2SO_4$.

Very sl. sol. in H_2O . (Jørgensen, J. pr. 15. 429.)

Iodoplatinsemidiamine iodide,

Sl. sol. in H_2O . (Jørgensen, J. pr. (2) 345.)

— periodide, $I_2Pt(NH_5)_2I, I_2$.

Moderately sl. sol. in H_2O . (Cleve.)

Iododiplatinamine iodide, $I_2Pt_2(N_2H_5)_4$.

Insol. in H_2O .

Iododiplatindiamine anhydroiodide,

Insol. in $NH_4OH + Aq$.

— anhydronitrate, $I_2Pt_2(N_2H_5)_4O(NO_2)_2$.

Easily sol. in warm $H_2SO_4 + Aq$. (Cleve.)

— iodide, $I_2Pt_2(N_2H_5)_4I_4$.

Ppt.

— nitrate, $I_2Pt_2(N_2H_5)_4(NO_2)_4 + 4H_2O$.

Sl. sol. in cold, moderately sol. in hot H_2O (Cleve.)

— phosphate, $I_2Pt_2(N_2H_5)_4[O_3P(OH)]_2$.

Nearly insol. in H_2O .

— sulphate, $I_2Pt_2(N_2H_5)_4(SO_4)_2$.

Nearly insol. in H_2O .

— platodiamine sulphate, $I_2Pt_2(N_2H_5)_4Pt(NH_5)_2SO_4$.

Very sl. sol. in H_2O . (Carlgren Sv. V. A. 47. 306.)

Iodoplatinic acid, $H_2PtI_6 + 9H_2O$.

Deliquescent. Easily sol. in H_2O , with comp. into PtI_4 and HI on standing or warming. (Topsoë.)

1 iodoplatinate, $(\text{NH}_4)_2\text{PtI}_6$.
sl. in H_2O . (Topsoë.)
 PtI_6 . Sl. sol. in H_2O ; insol. in
Lassaigne, A. ch. (2) 51. 128.)

2 iodoplatinate, BaPtI_6 .
cent, but less so than Na_2PtI_6
herwise resembles. (Lassaigne.)

3 iodoplatinate, $\text{CaPtI}_6 + 12\text{H}_2\text{O}$.
deliquescent as Na salt.

4 iodoplatinate, $\text{CoPtI}_6 + 9\text{H}_2\text{O}$.
deliquescent.

5 iodoplatinate, $[\text{PtI}_4(\text{OH})_2]\text{PbI}_2$.
elluci, C. C. 1902, I. 625.)

6 iodoplatinate, $\text{MgPtI}_6 + 9\text{H}_2\text{O}$.
deliquescent.

7 iodoplatinate, $\text{MnPtI}_6 + 9\text{H}_2\text{O}$.
deliquescent.

8 triiodoplatinate, $[\text{PtI}_4(\text{OH})_2]\text{HgI}_2$.
elluci, C. C. 1902, I. 625.)

9 iodoplatinate, $\text{NiPtI}_6 + 9\text{H}_2\text{O}$.
deliquescent.

10 iodoplatinate, K_2PtI_6 .
sl. in H_2O . Insol. in alcohol. Not
by cold conc. H_2SO_4 .

11 iodoplatinate, $\text{Pt}[\text{I}_4(\text{OH})_2]\text{Ag}_2$.
elluci, C. C. 1902, I. 625.)

12 iodoplatinate, $\text{Na}_2\text{PtI}_6 + 6\text{H}_2\text{O}$.
deliquescent, but easily sol. in H_2O
sol. (Vauquelin.) Deliquescent.
)

13 triiodoplatinate, $[\text{PtI}_4(\text{OH})_2]\text{TiI}_2$.
elluci, C. C. 1902, I. 625.)

14 iodoplatinate, $\text{ZnPtI}_6 + 9\text{H}_2\text{O}$.
sl. in H_2O .

15 dicyanhydric acid, $\text{H}_2\text{Pt}(\text{CN})_4\text{I}_2$.
16 iodoplatinocyanhydric acid.

17 platinumocyanide, $\text{Ag}_2(\text{PtI}_2(\text{CN})_4)_2$.
liolati, Gazz. ch. it. 1900, 30. 588.)

18 iodoplatinocyanide platinumocyanide,
 $2(\text{CN})_4\text{I}_2, 108\text{rPt}(\text{CN})_4 + x\text{H}_2\text{O}$.

Iodopurpleochromium chloride,
 $\text{ICr}(\text{NH}_3)_3\text{Cl}_2$.
Quite sol. in H_2O . (Jørgensen, J. pr. (2)
25. 83.)

— **chloroplatinate, $\text{ICr}(\text{NH}_3)_3\text{PtCl}_6$.**
Precipitate. (Jørgensen, l. c.)

— **iodide, $\text{ICr}(\text{NH}_3)_3\text{I}_3$.**
Difficultly sol. in H_2O . Insol. in HI , or
 $\text{KI} + \text{Aq}$; insol. in alcohol. (Jørgensen, l. c.)

— **nitrate, $\text{ICr}(\text{NH}_3)_3(\text{NO}_3)_3$.**
Much less sol. in H_2O than the chloride.
(Jørgensen, l. c.)

Iodopurpleocobaltic iodide, $\text{CoI}(\text{NH}_3)_3\text{I}_3$.
(Claudet.)
Does not exist. (Jørgensen, J. pr. (2) 25.
94.)

Iodopurpleorhodium chloride,
 $\text{IRh}(\text{NH}_3)_3\text{Cl}_2$.
Relatively easily sol. in H_2O ; insol. in HCl
 $+ \text{Aq}$ and alcohol. Insol. in $\text{KI} + \text{Aq}$. (Jør-
gensen, J. pr. (2) 27. 433.)

— **fluosilicate, $\text{IRh}(\text{NH}_3)_3\text{SiF}_6$.**
Nearly insol. in cold H_2O .

— **iodoplatinate, $\text{IRh}(\text{NH}_3)_3\text{PtI}_6$.**
Ppt.

— **iodide, $\text{IRh}(\text{NH}_3)_3\text{I}_3$.**
Very sl. sol. in cold H_2O ; more sol. in hot
 H_2O ; insol. in dil. $\text{HI} + \text{Aq}$, and alcohol.
(Jørgensen, J. pr. (2) 27. 433.)

— **nitrate, $\text{IRh}(\text{NH}_3)_3(\text{NO}_3)_3$.**
Sl. sol. in H_2O , more easily sol. in hot H_2O ;
insol. in dil. $\text{HNO}_3 + \text{Aq}$, and alcohol.

— **sulphate, $\text{IRh}(\text{NH}_3)_3\text{SO}_4$, and $+3\text{H}_2\text{O}$.**
Sl. sol. in even hot H_2O . (Jørgensen.)

Iodoselenic acid.

**Ammonium iodoselenate, $2(\text{NH}_4)_2\text{O}, \text{I}_2\text{O}_5,$
 $2\text{SeO}_3 + \text{H}_2\text{O}$.**
Decomp. by H_2O . (Weinland, B. 1903, 36.
1400.)

$2(\text{NH}_4)_2\text{O}, 3\text{I}_2\text{O}_5, 2\text{SeO}_3 + 5\text{H}_2\text{O}$. Sol. in
 H_2O with decomp.(?). (Weinland.)

**Potassium iodoselenate, $2\text{K}_2\text{O}, \text{I}_2\text{O}_5, 2\text{SeO}_3 +$
 H_2O .**

Decomp. by H_2O . (Weinland.)
 $2\text{K}_2\text{O}, 3\text{I}_2\text{O}_5, 2\text{SeO}_3 + 5\text{H}_2\text{O}$. Sol. in H_2O
with decomp.(?) (Weinland.)

Rubidium iodoselenate, $2\text{Rb}_2\text{O}$, $3\text{I}_2\text{O}_5$, 2SeO_2 , $+5\text{H}_2\text{O}$.

Sol. in H_2O . (Weinland.)

Iodostannous acid.

Data concerning solubility of SnI_2 in $\text{HI} + \text{Aq}$ indicate formation of this compound. (Young, J. Am. Chem. Soc. 1897, 19. 853.)

Iodosulphobismuthous acid.

Cuprous iodosulphobismuthite, $2\text{Cu}_2\text{S}$, Bi_2S_3 , 2BiSI .

Decomp. by H_2O at ord. temp. Decomp. by mineral acids with evolution of H_2S . (Ducatte, C. R. 1902, 134. 1213.)

Lead iodosulphobismuthite, PbS , Bi_2S_3 , 2BiSI .

Insol. in H_2O . Partially decomp. by boiling H_2O . Decomp. by dil. mineral acids with evolution of H_2S . (Ducatte.)

Iodosulphuric acid.

Ammonium iodosulphate, $(\text{NH}_4)_2\text{SO}_4\text{I}_2(?)$.

Very sol. in H_2O . (Zinno, N. Rep. Pharm. 20. 449.)

Mercuric iodosulphate, $\text{Hg}_2(\text{SO}_4)\text{I}_2$.

See **Mercuric sulphate iodide**.

Potassium iodosulphate, $\text{K}_2\text{SO}_4\text{I}_2(?)$.

Sol. in 7.14 pts. H_2O at 15° . (Zinno, N. Rep. Pharm. 20. 449.)

Sodium iodosulphate, $\text{Na}_2\text{SO}_4\text{I}_2 + 10\text{H}_2\text{O}$.

Sol. in 3.64 pts. H_2O at 15° and in dil. alcohol. (Zinno, N. Rep. Pharm. 20. 449.)

Does not exist. (Michaelis and Koethe, B. 6. 999.)

Iodosulphuric anhydride, ISO_3 .

Decomp. very violently by H_2O . (Weber, J. pr. (2) 25. 224.)

Diiodosulphuric anhydride, I_2SO_3 .

Decomp. with H_2O , but not so violently as ISO_3 . (Weber, J. pr. (2) 25. 224.)

Iodo/trisulphuric anhydride, $\text{I}(\text{SO}_3)_3$.

Decomp. by H_2O . (Weber, J. pr. (2) 25. 224.)

Iodotelluric acid.

Ammonium iodotellurate, $(\text{NH}_4)_2\text{O}$, I_2O_5 , $2\text{TeO}_2 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Weinland, Z. anorg. 1901, 28. 52.)

$(\text{NH}_4)_2\text{O}$, I_2O_5 , $2\text{TeO}_2 + 8\text{H}_2\text{O}$. Sol. in H_2O . (Weinland, B. 1900, 33. 1017.)

Cæsium iodotellurate, Cs_2TeI_4 .

Insol. in CsI , or $\text{HI} + \text{Aq}$. Decom. by cold, rapidly by hot H_2O . (Wh. Am. J. 145. 267.)

Potassium iodotellurate, $\text{K}_2\text{TeI}_4 + 2$

Sl. efflorescent. Somewhat sol. in and dil. $\text{HI} + \text{Aq}$. (Wheeler.)

K_2O , I_2O_5 , $\text{TeO}_2 + 3\text{H}_2\text{O}$. Sol. Partially decomp. on recryst. fr. (Weinland, Z. anorg. 1901, 28. 53.)

K_2O , I_2O_5 , $2\text{TeO}_2 + 6\text{H}_2\text{O}$. Sol. without decomp. (Weinland.)

Rubidium iodotellurate, Rb_2TeI_4 .

Sl. sol. in HI , or $\text{RbI} + \text{Aq}$. D. H_2O . Somewhat sol. in alcohol.

Rb_2O , I_2O_5 , $2\text{TeO}_2 + 6\text{H}_2\text{O}$. So (Weinland.)

Iodotetramine chromium iodide

$\text{ICr}(\text{NH}_3)_4\text{I}_2 + \text{H}_2\text{O}$.

Sol. in H_2O . Pptd. by alcohol.

Iodotetramine cobaltic sulphate

$\text{ICo}(\text{NH}_3)_4\text{SO}_4$.

(Vortmann and Blasberg, B. 22.

Iodotungstic acid.

See **Tungstoiodic acid**.

Iodous acid, I_2O_3 .

See **Iodine trioxide**.

Iodovanadic acid, I_2O_5 , $\text{V}_2\text{O}_5 + 5$

Very easily sol. in H_2O .

$2\text{V}_2\text{O}_5$, $3\text{I}_2\text{O}_5 + 18\text{H}_2\text{O}$. (Ditte, 757.)

Ammonium iodovanadate, $3(\text{NH}_4)$

$5\text{I}_2\text{O}_5 + 20\text{H}_2\text{O}$.

Sol. in H_2O . (Ditte, C. R. 102.

Irididiamine compounds, $\text{Cl}_2\text{Ir}(\text{CN})_4$

See **Chloriridiamine compounds**.

Iridic acid.

Potassium iridate (?).

Sol. in H_2O and $\text{HCl} + \text{Aq}$.

Iridicyanhydric acid, $\text{H}_2\text{Ir}(\text{CN})_6$

Easily sol. in H_2O , still more in alcohol, less in ether. (Martius, A.

Barium iridicyanide, $\text{Ba}_2[\text{Ir}(\text{CN})_6]$

Efflorescent. Easily sol. in hot alcohol. Nearly insol. in alcohol. Not d acids.

iridicyanide ammonia, $\text{Cu}_2\text{Ir}_2(\text{CN})_{12}$, $+4\text{H}_2\text{O}$.
decomp. in air. (Rimbach, Z. anorg. 13.)

iridicyanide, $\text{K}_2\text{Ir}(\text{CN})_6$.
sol. in H_2O .

iridicyanide ammonia, $\text{Ag}_2\text{Ir}(\text{CN})_6$, $+3\text{H}_2\text{O}$.
decomp. in the light. (Rimbach, Z. 7, 52. 414.)

Ir.
sol. in all acids, including aqua regia, even in finely divided state, as "black," when it is sol. in aqua regia. (Pr. 42. 251.)

iridium ammonia compounds.

iridium diamine comp., $\text{ClIr}(\text{NH}_3)_2\text{X}$.
amine " $\text{Ir}(\text{NH}_3)_2\text{X}_2$.
amine " $\text{Ir}(\text{NH}_3)_4\text{X}_2$.
amine " $\text{Ir}(\text{NH}_3)_6\text{X}_3$.
pentamine " $\text{Ir}(\text{NH}_3)_5(\text{OH})\text{X}_2$.
amine " $\text{Ir}(\text{NH}_3)_6\text{X}_3$.
amine " $\text{Ir}(\text{NH}_3)_4\text{X}_2$.

iridium bromide, $\text{IrBr}_3 + 4\text{H}_2\text{O}$.
sol. in H_2O . Insol. in alcohol or ammonia.

iridium bromide, IrBr_3 , or H_2IrBr_4 .
cent. Sol. in H_2O and alcohol.
iridic acid.

iridium hydrogen sesquibromide, 3HBr , $+ \text{H}_2\text{O} = \text{H}_2\text{IrBr}_4 + 3\text{H}_2\text{O}$.
iridous acid.

iridium bromide with MBr.
iridite, M.

iridium bromide with MBr.
iridate, M.

iridium phosphorous bromide, IrBr_3 , 3PBr_3 .
decomp. by H_2O into a sol., and
iridification. Sol. in PBr_3 . (Geisen-

iridium bromide. Not easily attacked by H_2O .
 2Cl_2 .
iridium phosphorous chlorobromide.

iridium chloride, $\text{IrCl}_3(?)$.

Ir.
sol. toward H_2O , acids, and alcohol. (B. 1909, 42. 1773.)

iridium monochloride, IrCl .

Insol. in acids and bases. (Wöhler, B. 1913, 46. 1584.)

iridium dichloride, IrCl_2 .

Insol. in acids and bases. (Wöhler, B. 1913, 46. 1585.)

iridium trichloride, IrCl_3 .

Insol. in acids or alkalies. (Claus, C. C. 1861. 690.)

Insol. in H_2O , acids and alkalies, (Leidié, C. R. 1899, 129. 1251.)
 $+4\text{H}_2\text{O}$. Sol. in H_2O . (Claus.)

iridium tetrachloride, IrCl_4 , or $\text{H}_2\text{IrCl}_6(?)$.

Deliquescent, and easily sol. in H_2O .

iridium trichloride with MCl.

See Chloriridite, M.

iridium tetrachloride with MCl.

See Chloriridate, M.

iridium chloride with potassium chloride and sulphite.

See Chloriridosulphite, potassium.

iridium phosphorus chloride, IrP_2Cl_6 .

Insol. in cold H_2O . Sl. decomp. by hot H_2O . (Geisenheimer, A. ch. (6) 23. 254.)

$\text{IrP}_2\text{Cl}_{10}$. Very sol. in chloroform. (G.)
 $\text{IrP}_2\text{Cl}_{12}$. Easily sol. in PCl_5 or CHCl_3 , also in CS_2 with gradual decomp. Sl. sol. in cold H_2O . Decomp. by boiling into IrCl_3 , $3\text{H}_2\text{PO}_4$. Sl. sol. in benzene, ligroin and CCl_4 . (Strecker, B. 1909, 42. 1772.)

$+ \text{H}_2\text{O}$. Insol. in PCl_5 at 100° . Very slowly sol. in boiling H_2O . (Geisenheimer, A. ch. (6) 23. 266.)

$\text{IrP}_2\text{Cl}_{12}$. Decomp. by H_2O into 2IrCl_3 , $3\text{H}_2\text{PO}_4$, $3\text{H}_2\text{PO}_4$. Violently decomp. by alcohol. Sl. sol. in cold, more in hot POCl_3 , without decomp. Very sol. in PCl_5 with decomp. into $\text{IrP}_2\text{Cl}_{12}$; similarly in PBr_3 . Sol. in AsCl_3 with combination. Sol. in CS_2 with decomp. Sol. in SCl_2 with combination. Easily sol. in cold C_6H_6 with decomp. Insol. in CCl_4 . Sol. in CHCl_3 with decomp. (Geisenheimer, A. ch. (6) 23. 254.)

iridium phosphorus arsenic chloride,
 $2\text{IrP}_2\text{Cl}_{12}$, 5AsCl_3 .

Sol. in H_2O with decomp. into corresponding acid. (Geisenheimer, C. R. 110. 1336.)

IrCl_3 , 2PCl_5 , 2AsCl_3 . Very sol. in H_2O with decomp. Sol. in AsCl_3 ; insol. in CCl_4 . (Geisenheimer.)

iridium phosphorus sulphur chloride, IrCl_3 , 2PCl_5 , 2SCl_2 .

Very sol. in sl. amt. H_2O , with decomp. into an acid analogous to chlorophosphoiridic acid. Sol. in SCl_2 . (Geisenheimer.)

Iridium phosphorus chlorobromide, $\text{IrBr}_4, 2\text{PCl}_3$.

(Geisenheimer, C. R. 111. 40.)

Iridium dihydroxide, $\text{IrO}_2, 2\text{H}_2\text{O} = \text{IrO}_4\text{H}_4$.

Insol. in dil. HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq}$. Slowly but completely sol. in $\text{HCl} + \text{Aq}$. Sol. in KOH , and $\text{NaOH} + \text{Aq}$. (Claus, J. pr. 39. 104.)

Iridium sesquihydroxide, $\text{Ir}_2\text{O}_3\text{H}_4$.

Not attacked by acids, except slightly by conc. $\text{HCl} + \text{Aq}$. (Claus, C. C. 1861. 690.)

Iridium triiodide, IrI_3 .

Very sl. sol. in cold, somewhat more in hot H_2O . Insol. in alcohol. (Oppler, J. B. 1857. 263.)

Iridium tetraiodide, IrI_4 .

Insol. in H_2O or acids. (Lassaigne.)

Sol. in solutions of iodides. (Oppler.)

Iridium triiodide with MI.

See Iodiridite, M.

Iridium tetraiodide with MI.

See Iodiridate, M.

Iridium dioxide, IrO_2 .

Very sl. sol. in acids.

Freshly pptd. Sol. in conc. H_2SO_4 , hot 2-N H_2SO_4 , HNO_3 , HCl . Insol. in 2-N KOH and sl. sol. in hot 1-N KOH .

Dried in a dessicator. Sol. in HCl . Insol. in H_2SO_4 , HNO_3 , and KOH .

Dried at 100° . Sol. in hot conc. HCl . Insol. in H_2SO_4 , HNO_3 , and KOH . (Wöhler, Z. anorg. 1908, 57. 334.)

See also Iridium dihydroxide.

Iridium trioxide, IrO_3 .

Unstable. (Wöhler, Z. anorg. 1908, 57. 340.)

Iridium sesquioxide, Ir_2O_3 .

Insol. in acids.

Sol. in conc. H_2SO_4 , and hot conc. HCl . Forms colloidal solution with dil. HCl . Conc. HNO_3 converts it into the dioxide.

Insol. in $\text{KOH} + \text{Aq}$. (Wöhler, Z. anorg. 1908, 57. 339.)

Iridium oxybromide, $\text{Ir}_2\text{Br}_5\text{O}_2 = 2\text{IrBr}_4, \text{IrO}_2$.

Not decomp. by H_2O . (Geisenheimer, A. ch. (6) 23. 286.)

Iridium phosphide, Ir_3P .

(Clarke and Joslin, Am. Ch. J. 5. 231.)

Iridium sesquiselenide, Ir_2Se_3 .

Insol. in HNO_3 ; slowly sol. in hot aqua regia. (Chabrie and Bouchonnet, C. R. 1903, 137. 1060.)

Iridium monosulphide, IrS .

Insol. in $\text{HNO}_3 + \text{Aq}$, and very at all in aqua regia. (Berzelius.)

Sol. in K_2S , and $\text{KSH} + \text{Aq}$. $+x\text{H}_2\text{O}$. Sl. sol. in H_2O ; sol. $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{NH}_4\text{Cl} + \text{acids}$. More sol. in $\text{K}_2\text{S} + \text{Aq}$ t (Berzelius.)

Iridium disulphide, IrS_2 .

Not attacked by H_2O , but dec exposed moist in air. Not attack $\text{HCl} + \text{Aq}$ or by conc. $\text{HNO}_3 + \text{oxidised}$ by fuming $\text{HNO}_3 + \text{Aq}$, regia. Insol. in NH_4 sulphides, phides + Aq . Slowly sol. in alk phides + Aq . (Antony, Gazz. ch. 190.)

Iridium sesquisulphide, Ir_2S_3 .

Sl. sol. in H_2O . Sol. in HNO_3 , or

Iridotriamine chloride, $\text{Ir}(\text{NH}_3)_3$

Sl. sol. in H_2O . Not attacked H_2SO_4 . (Palmaer, B. 22. 15.)

Iridotetraamine chloride, $\text{Ir}(\text{NH}_3)_4$

Very sol. in H_2O . (Palmaer, B.

— chlorosulphate, $[\text{Ir}(\text{NH}_3)_4\text{Cl}_2]\text{S}_4\text{H}_2\text{O}$.

(Palmaer.)

Iridopentamine bromide, $\text{Ir}(\text{NH}_3)_5$

Sol. in 352 pts. H_2O at 12.5° . (I 23. 3817.)

— bromochloride, $\text{Ir}(\text{NH}_3)_5\text{ClBr}$

Sol. in H_2O . (Palmaer, B. 24. 20)

— bromonitrite, $\text{Ir}(\text{NH}_3)_5\text{Br}(\text{NO})$

Sol. in 17.9 pts. H_2O at 18° . (I

— bromosulphate, $\text{Ir}(\text{NH}_3)_5\text{BrS}$

Sol. in H_2O . (Palmaer.)

— carbonate, $[\text{Ir}(\text{NH}_3)_5](\text{CO}_3)_2$

Sol. in H_2O . (Claus, J. pr. 63.)

— trichloride, $\text{Ir}(\text{NH}_3)_5\text{Cl}_3$.

Sol. in 153.1 pts. H_2O at 15.1° . B. 23. 3810.)

Sol. in hot H_2O containing HCl . pr. 69. 30.)

— chlorobromide, $\text{Ir}(\text{NH}_3)_5\text{ClBr}$

Sol. in 213.6 pts. H_2O at 15° . (I

— chloriodide, $\text{Ir}(\text{NH}_3)_5\text{ClI}_2$.

Sol. in 104.5 pts. H_2O at 15° . (I

— chlorooxalate, $\text{Ir}(\text{NH}_3)_5\text{ClC}_2\text{O}_4$

Sl. sol. in H_2O . (Palmaer.)

amine chloronitrate,
 $\text{H}_3\text{IrCl}_2(\text{NO}_2)_2$.
 51.54 pts. H_2O at 15.4° . (Palmaer.)

ronitrite,
 $\text{H}_3\text{IrCl}(\text{NO}_2)_2$.
 sol. in H_2O . (Palmera.)

roplatinate, $\text{Ir}(\text{NH}_3)_5\text{Cl}_2$, PtCl_4 .
 . sol. in H_2O . (Palmeaar.)

rosulphate, $\text{Ir}(\text{NH}_3)_5\text{ClSO}_4 + 2\text{H}_2\text{O}$.
 134.5 pts. H_2O at 15° . (Palmaer.)

oxide, $\text{Ir}(\text{NH}_3)_5(\text{OH})_2$.
 only in solution, which decomp. on
 on. (Claus.)

te, $\text{Ir}(\text{NH}_3)_5(\text{NO}_2)_2$.
 tely sol. in H_2O . (Claus.)
 349 pts. H_2O at 16° . (Palmaer.)

iate, $[\text{Ir}(\text{NH}_3)_5]_2(\text{SO}_4)_3$.
 H_2O . (Claus.)

pentamine bromide,
 $\text{Ir}_2(\text{OH})_2\text{Br}_2$.
 4 pts. H_2O . Pptd. from aqueous
 by $\text{HBr} + \text{Aq}$. (Palmaer, B. 24.)

ide, $\text{Ir}(\text{NH}_3)_5(\text{OH})_2\text{Cl}_2$.
 1.2 to 1.5 pts. H_2O at ord. temp.
 $\text{HCl} + \text{Aq}$ from aqueous solution.
 B. 24. 2090.)

te, $\text{Ir}(\text{NH}_3)_5(\text{OH})_2(\text{NO}_2)_2$.
 about 10 pts. H_2O at 17° . Pptd.
 ous solution by $\text{HNO}_3 + \text{Aq}$. (Pal-

us acid, $\text{H}_4\text{Ir}_2(\text{NO}_2)_{12}$.
 ol. in H_2O . (Gibbs, B. 4. 281.)

n iridonitrite, $(\text{NH}_4)_4\text{Ir}_2(\text{NO}_2)_{12}$.
 insol. in cold H_2O ; decomp. by hot
 evolution of N_2 . Decomp. by hot
 O_2 or HCl . Insol. in sat. $\text{NH}_4\text{Cl} +$
 lié, C. R. 1902, 134. 1583.)

donitrite iridochloride,
 $(\text{NO}_2)_{12}$, $\text{Ba}_2\text{Ir}_2\text{Cl}_{12}$.
 I_2O . (Lang.)

ridonitrite, $\text{Hg}_2\text{Ir}_2(\text{NO}_2)_{12}$.
 H_2O . (Gibbs, B. 4. 280.)

ridonitrite, $\text{K}_4\text{Ir}_2(\text{NO}_2)_{12}$.
 n cold, more sol. in boiling H_2O .
 y hot HCl or H_2SO_4 . Insol. in
 (Leidié, Bull. Soc. 1902, (3) 27.

Moderately sol. in H_2O .

Potassium iridonitrite iridochloride,
 $\text{K}_4\text{Ir}_2(\text{NO}_2)_{12}$, $\text{K}_4\text{Ir}_2\text{Cl}_{12}$.
 Sol. in H_2O .

Silver iridonitrite, $\text{Ag}_4\text{Ir}_2(\text{NO}_2)_{12}$.
 Difficultly sol. in cold, more easily in hot
 H_2O .

Sodium iridonitrite, $\text{Na}_4\text{Ir}_2(\text{NO}_2)_{12} + 2\text{H}_2\text{O}$.
 Sl. sol. in H_2O . Sol. in cold H_2O . Decomp.
 by hot conc. H_2SO_4 or HCl . (Leidié, C. R.
 1902, 134. 1583.)

Sodium iridonitrite iridochloride,
 $\text{Na}_4\text{Ir}_2\text{Cl}_2(\text{NO}_2)_8 + 2\text{H}_2\text{O}$.
 Sl. sol. in H_2O . (Gibbs.)
 $\text{Na}_4\text{Ir}_2(\text{NO}_2)_{12}$, $\text{Na}_4\text{Ir}_2\text{Cl}_4$. Insol. in cold, sl.
 sol. in hot H_2O . (Lang.)

Iridosamine chloride, $\text{Ir}(\text{NH}_3)_5\text{Cl}_2$.
 Nearly insol. in H_2O . (Skoblikoff, A. 34.
 275.)

— **sulphate**, $\text{Ir}(\text{NH}_3)_5\text{SO}_4$.
 Easily sol. in H_2O . (Skoblikoff.)

Iridosodiamine chloride, $\text{Ir}(\text{N}_2\text{H}_5)_2\text{Cl}_2$.
 Insol. in cold, decomp. by hot H_2O . (Skob-
 likoff.)

— **nitrate**, $\text{Ir}(\text{N}_2\text{H}_5\text{NO}_2)_2$.
 Easily sol. in H_2O .

— **sulphate**, $\text{Ir}(\text{N}_2\text{H}_5)_2\text{SO}_4$.
 Sl. sol. in cold, easily in boiling H_2O . Sl.
 sol. in alcohol.

Iridosulphuric acid.

Potassium iridosulphate, $\text{K}_4\text{Ir}_2(\text{SO}_4)_6$.
 Sol. in H_2O . (de Boisbaudran, C. R. 96.
 1406.)

Iridosulphurous acid.

Ammonium iridosulphite, $(\text{NH}_4)_4\text{Ir}_2(\text{SO}_3)_6 +$
 $6\text{H}_2\text{O}$.
 Slightly sol. in H_2O . (Birnbbaum, A. 136.
 179.)

Potassium iridosulphite, $\text{K}_4\text{Ir}_2(\text{SO}_3)_6 + 6\text{H}_2\text{O}$.
 Slightly sol. in H_2O .

Sodium iridosulphite, $\text{Na}_4\text{Ir}_2(\text{SO}_3)_6 + 8\text{H}_2\text{O}$.
 Scarcely sol. in H_2O .

Iron, Fe.

Permanent in dry air; oxidises only slowly
 in moist air, but rapidly when in contact
 with air and H_2O simultaneously.

Fe does not rust in contact with air and

H₂O containing alkalies even in very small amounts. (Payen, A. ch. 50. 305.)

Not attacked at ord. temp. by H₂O free from air. More easily oxidised by NH₄ salts + Aq than by H₂O when exposed to air simultaneously. (Persoz, A. ch. (3) 24. 506.)

Iron is slowly attacked by distilled H₂O in presence of air. 100 ccm. distilled water removed 29 mg. from 11.8 sq. cm. iron in one week, while air free from CO₂ was passed through the solution. In presence of CO₂, 54 mg. were removed. (Wagner, Dingl. 221. 260.)

CO₂ acts as a catalyst for the solution of Fe by H₂O. (Whitney, J. Am. Chem. Soc. 1903, 25. 394.)

Iron is most easily oxidised when it is exposed to air, and H₂O is deposited on it at the same time in liquid form.

100 l. sea water dissolve 27.37 g. from 1 sq. metre Fe; 29.16 g. from 1 sq. metre steel; 1.12 g. from 1 sq. metre galvanised Fe. (Calvert and Johnson, C. N. 11). 171.)

Readily sol. in HCl, dil. H₂SO₄ + Aq, and most other acids.

Action of H₂SO₄ + Aq (1:12) is very much accelerated by a few drops of PtCl₄ + Aq; the addition of As₂O₃ arrests the action completely. Tartar emetic and HgCl₂ diminish the action, but do not arrest it. CuSO₄ + Aq strongly accelerates the action, and Ag₂SO₄ + Aq also to a less extent.

In the case of HCl + Aq, the addition of small amts. of metallic salts also influences the action. Weak HC₂H₃O₂ + Aq has but little action, and the addition of PtCl₄ increases it; As₂O₃ stops it; other solutions have no effect. With racemic and tartaric acids the phenomena are the same.

With oxalic acid, PtCl₄ prevents the action. Saline solutions and even distilled H₂O, when mixed with PtCl₄, have slight solvent action. (Millon, C. R. 21. 45.)

Above phenomena are due to galvanic action from metal deposited on the iron. (Barreswill, C. R. 21. 292.)

H₂SO₄ has only sl. action on cast-iron at ord. temp. with exclusion of air.

Weak acids have a strong action at higher temperatures.

Charcoal pig-iron, and case-hardened cast-iron are much less attacked by weak acids at h.-pt. than other sorts of Fe. Scotch pig-iron is most strongly attacked.

99.8% H₂SO₄ has very sl. action on iron at ord. temp. when air is excluded. (Lunge, Dingl. 261. 131.)

Resistance against dil. H₂SO₄ + Aq is greatly increased by increase in amt. of C if chemically combined, less so by P or Si. (Ledebur, Dingl. 223. 326.)

Passive Iron. -When Fe is treated with pure conc. HNO₃ + Aq of 1.512-1.419 sp. gr., it soon becomes coated with a bluish or black coating, apparently FeO, and when thus

covered Fe is not attacked by HNO₃ + Aq of any strength at ord. temp. or at the temp. of a freezing mixture; but action occurs on heating. Nor is Fe attacked at ord. temp. by acid of 1.401 sp. gr. or even somewhat weaker acid, though action begins at once on heating. Very dil. HNO₃ + Aq attacks Fe at ord. temp. with formation of NH₄NO₃ and Fe(NO₃)₃. The action of HNO₃ + Aq is influenced by PtCl₄. If acid containing 45 equivalents of H₂O is diluted with 2-3 vol. H₂O, and then poured on Fe turnings, they dissolve at once with evolution of nitrous fumes and formation of ferric salt, but if to the acid one drop of PtCl₄ be added, only H gas is evolved, and NH₄NO₃ and Fe(NO₃)₃ are formed. (Millon, C. R. 21. 47.)

The more H₂O the acid contains the lower will be the temp. at which the Fe remains passive. Shaking the wire hastens the passivity. Contact with Pt, Au, or C does not prevent it. Fe wire becomes passive by remaining 10 min. in HNO₃ vapour. (Renard, C. R. 79. 159.)

Iron may be made passive by HClO₄, HBrO₃, HIO₃, H₂CrO₄, in the same way as by HNO₃.

Iron may also be made passive by moderate ignition.

Passivity occurs with HNO₃ + Aq of 1.38 sp. gr. after a short time at 31°; but if temp. is 32°, passivity does not occur.

Colourless HNO₃ + Aq of 1.42 sp. gr. produces passivity at 55° but not at 56°. Red fuming HNO₃ + Aq of 1.42 sp. gr. produces passivity at 82° but not at 83°. (Ordway, Sill. Am. J. (2) 40. 316.)

The passivity of Fe is destroyed when it is placed in a magnetic field at a much lower temperature than when in normal condition. (Nichols and Franklin, Sill. Am. J. (3) 34. 419.)

Passivity depends on a coating of NO which hinders the action of the acid. All operations which remove this layer terminate the passivity, as shaking, rubbing, placing in a vacuum, etc. (Varenne, C. R. 89. 783.)

When Fe is plunged in HNO₃ + Aq of 1.42 sp. gr. there is a sudden evolution of gas which ceases after 3 to 20 seconds, and the surface becomes bright. The same phenomena take place with a more dilute acid, if of not less than 1.32 sp. gr. In the latter case, there is an immediate evolution of gas, which suddenly ceases and the metal becomes bright, but soon the acid begins to act again at a single point, and the action gradually spreads over the whole surface; this, however, soon ceases again, and we have an "intermittent passivity."

If a part of a piece of iron is immersed in strong acid, the whole of it is made passive. This is explained by the NO spreading over the whole surface by capillarity.

The passivity ceases when the Fe is placed in dil. acid, after a longer or shorter time,

According to the dilution of the acid,—when acid has sp. gr. = 1.30, after 11 days
 " " " 1.28 " 5 "
 " " " 1.26 " 32 hours
 " " " 1.16 " 12 "

Iron may also be made passive by long standing in NO gas under pressure. (Varenne, R. 90. 998.)

Fe is made passive by a coating of Fe_3O_4 , by NO. (Schonbein, Pogg. 39. 342.)
 etc., Pogg. 67. 286.) (Ramann, B. 14. 0.)

Passivity may also be caused by NH_4NO_3 , aq., ammoniacal AgNO_3 +Aq, $\text{Fe}(\text{NO}_3)_3$, NO_3 , $\text{Al}(\text{NO}_3)_3$, $\text{Co}(\text{NO}_3)_2$, $\text{Ni}(\text{NO}_3)_2$, +Aq instead of HNO_3 +Aq. (Ramann, 4. 1933.)

Hardly attacked by either dil. or conc. acids when they are under high pressure. (Cailletet 2. 68. 395.)

Iron is dissolved by HNO_3 +Aq, even when conc., but no gas is evolved and the process is very slow.

NO_3 +Aq of the following sp. gr. dissolves given amts. from strips of pure Fe.

Sp. gr. of acid	Diminution of weight in 24 hours
1.28	0.82%
1.34	0.75
1.38	0.29
1.48	0.34
1.53	5.80

Gautier and Charpy, C. R. 113. 1451.)

Sol. in liquid chlorine below 90° . (Lange, Angew. Ch. 1900, 13. 686.)

Sol. in liquid NH_3 . (Gore, Am. ch. J. 3, 20. 828.)

Not attacked by alkalis.

Sol. in NaOH +Aq (34%) when air is blown through the liquid. (Zirnité, Ch. Ztg. 12.)

NaOH +Aq attacks iron and steel. (Vena-Dingl. 261. 133.)

NaOH +Aq has slight action on Fe between 50° and 100° . (Lunge, Dingl. 261. 131.)

Presence of alkalis prevent rusting entirely, and fats and oils greatly hinder it. (Wagner.)

Sol. in alkali hydrogen carbonates+Aq. (Zerelius.)

Sat. NaCl +Aq has sl. but perceptible action on Fe. NH_4Cl +Aq has stronger action than NaCl +Aq. (Lunge.)

100 ccm. H_2O containing 0.5 g. NaCl or 1 removed 42 mg. from 11.8 sq. cm. iron in one week, while air free from CO_2 was led through the solution, and 72 mg. in presence of CO_2 .

100 ccm. H_2O containing 1 g. NH_4Cl removed 45 mg., and 76 mg. respectively under above conditions.

100 ccm. H_2O containing 0.8 g. MgCl_2 re-

moved 49 mg., and 65 mg. respectively under the above conditions.

Not attacked by 100 ccm. H_2O containing 1 g. Na_2CO_3 , or by CaO, H_2 +Aq. (Wagner, Dingl. 221. 260.)

Action of KClO_3 +Aq. KClO_3 +Aq (6.3% KClO_3) oxidised 11.21 g. cast iron and 20.1 g. pure iron from a surface of 1 sq. metre in 7 hours; KClO_3 +Aq (25% KClO_3) oxidised 24.59 g. cast, and 44.90 g. pure Fe under above conditions; $\text{Ca}(\text{ClO}_3)_2$, CaCl_2 +Aq (20° Baume) obtained by passing Cl through CaO, H_2 +Aq oxidised 85 g. cast, and 95 g. pure Fe under the above conditions. (Lunge and Deggeler, J. Soc. Chem. Ind. 4. 32.)

Easily sol. in organic acids.

Comparative action of oils on Fe.

	Amount Fe dissolved
Neatsfoot oil	0.0875 grains
Colza "	0.0800 "
Sperm "	0.0460 "
Lard "	0.0250 "
Olive "	0.0062 "
Linseed "	0.0050 "
Seal "	0.0050 "
Castor "	0.0048 "
Paraffine "	0.0045 "
Almond "	0.0040 "
"Lubricating" oil	0.0018 "

(Watson, C. N. 42. 190.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0097 g. Fe in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Fe dissolves in albumen solution to the extent of 1 to 2 per cent. (Buchner, Arch. Pharm. (3) 20. 417.)

Attacked by sugar+Aq at 115 – 120° , also by inverted sugar or malt extract, not by glycerine or mannite+Aq. (Klein and Berg, C. R. 102. 1170.)

Iron arsenide, FeAs_2 .

Min. *Löllingite*. Sol. in HNO_3 +Aq with separation of As_2O_3 .

Fe_3As_4 . Min. *Leucopyrite*.

Iron arsenide sulphide, $\text{FeAs}_2, \text{FeS}_2$.

Min. *Arsenopyrite*. Sol. in HNO_3 +Aq with separation of S and As_2O_3 ; wholly sol. in aqua regia; not attacked by HCl +Aq.

Iron boride, Fe_2B .

Decomp. by H_2O . Sol. in hot dil. HCl or H_2SO_4 , and in hot conc. HCl or H_2SO_4 . Sol. in hot dil., or cold conc. HNO_3 . (Jassonneix, C. R. 1907, 145. 122.)

FeB . Sol. in molten alkali carbonates; not sol. in dil. or conc. H_2SO_4 in the cold; sol. in boiling H_2SO_4 and in HNO_3 . (Moissan, Bull. Soc. 1895, (3) 13. 958.)

Stable in dry air. Decomp. by aqua regia,

but not readily sol. in conc. H_2SO_4 and HCl . (Moissan, C. R. 1895, 120. 176.)

FeBr_2 . Decomp. by H_2O . Sol. in HNO_3 and in hot conc. HCl . (Jassonneix, C. R. 1907, 145. 122.)

Iron (ferrous) bromide, FeBr_2 .

Sol. in H_2O . Decomp. by heating on air.

Sat. $\text{FeBr}_2 + \text{Aq}$ contains at:

—21°	—7°	+10°	21°
47.0	48.3	52.3	53.7% FeBr_2 ,
37°	50°	65°	95°
56.0	58.0	59.4	63.3% FeBr_2 .

(Étard, A. ch. 1894, (7) 2. 541.)

+4 H_2O . Very sol. in H_2O ; pptd. from cooled aq. solution. (Volkmann, C. C. 1894, II. 611.)

+6 H_2O . Sol. in H_2O . (Löwig.)

+9 H_2O . (Volkmann.)

Ferric bromide, FeBr_3 .

Deliquescent. Sol. in H_2O , alcohol, and ether. (Löwig.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

+6 H_2O . Sol. in alcohol and ether. (Bolschakoff, C. C. 1898, II. 660.)

Ferrous mercuric bromide.

Deliquescent. (v. Bonsdorff.)

Ferric rubidium bromide, $\text{Rb}_2\text{FeBr}_4 + \text{H}_2\text{O}$.

Sol. in H_2O . (Walden, Z. anorg. 1894, 7. 332.)

Ferrous stannic bromide.

See Bromostannate, ferrous.

Ferrous bromide nitric oxide, $3\text{FeBr}_2, 2\text{NO}$.

Sol. in H_2O . Not isolated. (Thomas, C. R. 1896, 123. 944.)

Ferric bromochloride, FeCl_2Br .

Very deliquescent, and sol. in H_2O , alcohol, and ether. Notably sol. in chloroform, benzene, and toluene. Insol. in CS_2 . (Lenormand, C. R. 116. 820.)

Iron carbide, Fe_3C .

(Gurlt, J. B. 1856, 781.)

Mixture of Fe and FeC_4 . (Tunner, Polyt. Centralbl. 1861. 1227.)

Fe_3C . (Karsten, J. pr. 40. 229.)

Fe_3C . Sol. in hot conc. HCl ; oxidized slowly by moist air. (Campbell, Am. Ch. J. 1896, 18. 840–841.)

Fe_3C_2 . (Rammelsberg, C. C. 1847. 60.)

Iron molybdenum carbide, $\text{Fe}_3\text{C}, \text{Mo}_3\text{C}$.

Sol. in hydracids; insol. in HNO_3 . (Williams, C. R. 1898, 127. 484.)

Iron tungsten carbide, $2\text{Fe}_3\text{C}, 3\text{W}_3\text{C}$.

Insol. in H_2O and hydracids; sol. in HNO_3 and H_2SO_4 . (Williams, C. R. 1898, 127. 411.)

Iron carbonyl, $\text{Fe}(\text{CO})_5$.

Slowly decomp. on air. Not attacked by dil. H_2SO_4 , HNO_3 , or $\text{HCl} + \text{Aq}$. Conc. HNO_3 , $\text{Cl}_2 + \text{Aq}$, or $\text{Br}_2 + \text{Aq}$ decomp. easily. Sol. in alcoholic solution of KOH or NaOH with subsequent decomp. Sol. in alcohol, ether, benzene, mineral oils, etc. (Mond and Langer, Chem. Soc. 59. 1090.)

$\text{Fe}_2(\text{CO})_9$. Decomp. on air. Not attacked by H_2SO_4 or $\text{HCl} + \text{Aq}$. Sol. in alcoholic potash. Very much less sol. in organic solvents than $\text{Fe}(\text{CO})_5$. (Mond and Langer.)

Ferrous chloride, FeCl_2 .

Deliquescent. Easily sol. in H_2O with evolution of heat, or in alcohol. Insol. in ether. (Jahn.)

Sol. in 2 pts. H_2O at 18.75°. (Abl.)

Sol. in 1 pt. strong alcohol. (Wenzel.)

Sp. gr. of $\text{FeCl}_2 = \text{Aq}$ at 15.5°.

Sp. gr.	% FeCl_2	% $\text{FeCl}_2, 4\text{H}_2\text{O}$
1.05	5.40	8.45
1.06	6.43	10.00
1.07	7.47	11.60
1.08	8.48	13.20
1.09	9.49	14.86
1.10	10.47	16.41
1.11	11.45	17.86
1.12	12.42	19.46
1.13	13.37	20.96
1.14	14.31	22.41
1.15	15.24	23.87
1.16	16.15	25.31
1.17	17.05	26.73
1.18	17.94	28.13
1.19	18.83	29.51
1.20	19.68	30.85
1.21	20.50	32.14
1.22	21.39	33.53
1.23	22.24	34.84
1.24	23.05	36.11
1.25	23.86	37.38
1.26	24.68	38.67
1.27	25.44	39.87
1.28	26.19	41.04
1.29	26.98	42.29
1.30	27.75	43.49
1.31	28.49	44.65
1.32	29.23	45.81
1.33	29.96	46.94
1.34	30.68	48.08
1.35	31.39	49.18
1.36	32.10	50.30
1.37	32.79	51.39
1.38	33.47	52.46
1.39	34.14	53.50
1.40	34.80	54.55
1.41	35.46	55.57
1.42	36.09	56.58
1.43	36.73	57.55
1.44	37.33	58.51

(Dunn, J. Soc. Chem. Ind. 1902, 21. 300.)

Insol. in liquid HF. (Franklin, Z. anorg. 1905, 46. 2.)
Sol. in acetone; insol. in methylal. (Eidmann, C. C. 1899, II. 1014.)
Sol. in acetone. (Naumann, B. 1904, 37. 4328.)
Sl. sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)
Sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)
Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)
Yellow modification is sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)
Mol. weight determined in pyridine. (Wermer, Z. anorg. 1897, 15. 21.)
+2H₂O. (Jonas.)
+4H₂O. Deliquescent. Easily sol. in alcohol. Sol. in 0.68 pt. cold H₂O. (Reimann, Mag. Pharm. 17. 215.)
Sat. aq. solution contains at:
16° 18° 25° 28° 43°
10.5 40.9 41.0 42.5 44.4% FeCl₂,
50° 53° 72° 89° 96° 118°
15.0 45.9 49.2 51.3 51.0 51.7% FeCl₂.
(Étard, A. ch. 1894, (7) 2. 537.)
100 g. FeCl₂, 4H₂O + Aq contain 17.54 g. Fe at 22.8°; 18.59 g. at 43.2°. (Boecke, N. Jahrb. Min. 1911, I, 61.)

More sol. in water containing NO than in pure H₂O. (Gay, Bull. Soc. (2) 44. 175.)
Sol. in hot HCl + Aq. (Sabatier, Bull. Soc. 1895, (3) 13. 599.)
[Sabatier could not obtain FeCl₂ + 6H₂O of Lescœur.]
Ferroferric chloride, Fe₂Cl₃ + 18H₂O.
Deliquescent. (Lefort, J. Pharm. (4) 10. 85.)
Ferric chloride, Fe₂Cl₃ or FeCl₃.
Very deliquescent, and sol. in H₂O with evolution of great heat.
100 mols. H₂O dissolve mols. anhydrous Fe₂Cl₃ at t°.

t°	Mols. Fe ₂ Cl ₃	t°	Mols. Fe ₂ Cl ₃
66	29.20	80	29.20
70	29.42	100	29.75
75	28.92

(Rooseboom, Z. phys. Ch. 10. 477.)
See also hydrated salts below.
Solution in H₂O is decomp. into colloidal Fe₂O₃, xH₂O and HCl, upon heating if conc., and on simple standing if dil.

Krecke (J. pr. (2) 3. 286) gives the following table.

% Fe ₂ Cl ₃ in solution	Temp. at which Graham's colloidal hydrate is formed	Temp. at which Saint Gilles' colloidal hydrate is formed	Temp. at which oxychlorides are formed	Temp. at which Fe ₂ O ₃ is formed
32	100-130°	...	100° +	140°
16	100-120	...	"	120
8	100-110	...	"	110
4	90-100	...	90	...
2	87	...	87	...
1	83	100-130°
0.5	75	"
0.25	64	"
0.125	54	"
0.0625	36	"

Sp. gr. of Fe₂Cl₃ + Aq.

% Fe ₂ Cl ₃	Sp. gr. at 4.8°	Sp. gr. at 9.7°	Sp. gr. at 14.6°	Sp. gr. at 19.7°
9.61	1.5609	1.5575	1.5540	1.5497
1.00	1.4413	1.4387	1.4361	1.4335
3.95	...	1.3847	1.3824	1.3800
1.25	1.3381	1.3359	1.3339	1.3317
1.60	1.2351	1.2334	1.2318	1.2298
1.54	1.2140	1.2129	1.2107	1.2090
1.79	1.1534	1.1521	1.1507	1.1491
1.45	1.0939	1.0930	1.0918	1.0901
1.65	1.0382	...
1.70	1.0221	...

(Schult, from Gerlach, Z. anal. 27. 278.)

Sp. gr. of Fe₂Cl₃ + Aq increases or diminishes between 8° and 24° for a decrease or increase of temp. of 1° by the following amts.

% Fe ₂ Cl ₃	Corr.	% Fe ₂ Cl ₃	Corr.
50-60	0.0008	30-39	0.0005
45-49	0.0007	20-29	0.0004
40-44	0.0006	10-19	0.0003

(Hager, l. c.)

Sp. gr. of $\text{Fe}_2\text{Cl}_6 + \text{Aq}$ at 17.5° .						Solubility of Fe_2Cl_6 in $\text{HCl} + \text{Aq}$.			
$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	t°	Sat. solution contains per 100 mola. H_2O		Solid phase
							mola. HCl	mola. FeCl_3	
1	1.0073	21	1.1644	41	1.3746	30	0	12.70	$\text{Fe}_2\text{Cl}_6 + 12$
2	1.0146	22	1.1746	42	1.3870	"	5.92	16.07	
3	1.0219	23	1.1848	43	1.3994	"	0	20.90	
4	1.0292	24	1.1950	44	1.4118	25	0	10.90	
5	1.0365	25	1.2052	45	1.4242	"	2.33	23.72	
6	1.0438	26	1.2155	46	1.4367	"	0	24.50	
7	1.0513	27	1.2258	47	1.4492	20	0	10.20	
8	1.0587	28	1.2365	48	1.4617	"	5.60	23.60	
9	1.0661	29	1.2464	49	1.4742	"	0	25.70	
10	1.0734	30	1.2568	50	1.4867	10	0	9.10	
11	1.0814	31	1.2673	51	1.5010	"	8.75	8.00	
12	1.0894	32	1.2778	52	1.5153	"	16.70	16.65	
13	1.0974	33	1.2883	53	1.5296	"	13.80	23.05	
14	1.1054	34	1.2988	54	1.5439	0	0	8.25	
15	1.1134	35	1.3093	55	1.5582	"	7.52	6.51	
16	1.1215	36	1.3199	56	1.5729	"	13.37	6.33	
17	1.1297	37	1.3305	57	1.5876	"	16.80	8.70	
18	1.1378	38	1.3411	58	1.6023	"	18.45	10.23	
19	1.1458	39	1.3517	59	1.6170	"	20.40	15.40	
20	1.1542	40	1.3623	60	1.6317	"	20.10	16.00	
(Frans, J. pr. (2) 5. 283.)						"	19.95	17.70	$\text{Fe}_2\text{Cl}_6 + 7$
Sp. gr. of $\text{Fe}_2\text{Cl}_6 + \text{Aq}$ at 17.5° .						"	19.00	22.75	
$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	"	18.05	23.40	
1	1.0028	21	1.191	41	1.428	-10	0	7.40	
2	1.016	22	1.202	42	1.441	"	19.46	10.37	
3	1.025	23	1.212	43	1.454	"	20.48	20.54	
4	1.033	24	1.223	44	1.469	-12 5	20.25	21.56	
5	1.042	25	1.234	45	1.481	-15	0	8.08	
6	1.051	26	1.245	46	1.494	"	21.30	9.65	
7	1.060	27	1.256	47	1.507	-20	0	6.56	
8	1.069	28	1.268	48	1.520	"	7.50	4.90	
9	1.078	29	1.280	49	1.533	"	15.30	5.09	
10	1.087	30	1.292	50	1.547	"	20.56	7.08	
11	1.095	31	1.304	51	1.560	30	0	25.20	$\text{Fe}_2\text{Cl}_6 + 7$
12	1.104	32	1.316	52	1.573	"	4.25	27.80	
13	1.113	33	1.328	53	1.587	"	0	30.24	
14	1.123	34	1.340	54	1.600	25	0	23.50	
15	1.131	35	1.352	55	1.612	"	2.33	23.72	
16	1.140	36	1.364	56	1.624	"	7.50	29.75	
17	1.150	37	1.376	57	1.636	"	0	31.50	
18	1.160	38	1.390	58	1.648	20	0	22.50	
19	1.170	39	1.403	59	1.659	"	5.60	23.60	
20	1.180	40	1.415	60	1.670	"	11.05	26.20	
(Hager, Comm. 1883.)						"	11.05	29.20	
Sp. gr. of conc. $\text{Fe}_2\text{Cl}_6 + \text{Aq}$ at $20-21^\circ$.						"	0	22.00	
$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	$\frac{\% \text{Fe}_2\text{Cl}_6}{100}$	Sp. gr.	15	10.75	23.50	
60	1.669	65	1.715	70	1.758	"	14.90	28.15	
61	1.679	66	1.724	71	1.766	10	13.80	23.35	
62	1.688	67	1.733	72	1.774	"	17.80	27.75	
63	1.697	68	1.742	73	1.782	"	17.80	27.75	
64	1.706	69	1.750	74	1.790	0	18.05	23.40	
(Hager, l. c.)						"	19.50	25.93	

Solubility of Fe_2Cl_6 in $\text{HCl} + \text{Aq.}$ —Continued.

	Sat. solution contains per 100 mols. H_2O		Solid phase	t°	Sat. solution contains per 100 mols. H_2O		Solid phase
	mols. HCl	mols. FeCl_3			mols. HCl	mols. FeCl_3	
	0	35.00	$\text{Fe}_2\text{Cl}_6 + 5\text{H}_2\text{O}$	33	30.45	48.70	$\text{Fe}_2\text{Cl}_6 + 4\text{H}_2\text{O}$
	3.25	39.95		30	17.15	36.75	
	0	33.50		"	31.20	43.49	
	3.04	33.80		"	33.80	47.80	
	10.62	34.64		"	32.60	49.93	
	11.50	35.60		■	19.50	35.25	
	10.70	38.00		"	20.60	35.34	
	0	32.40		"	31.34	41.58	
	13.40	37.45		"	33.00	43.00	
	0	31.00		"	34.65	44.80	
	15.70	37.06		■	21.25	34.25	
	0	30.24		"	28.81	37.57	
	17.20	34.00		"	34.23	42.02	
	17.15	36.75		"	35.40	43.16	
	0	29.00		15	29.40	36.50	
	7.50	29.75		"	33.60	40.03	
	19.50	35.25		10	24.50	32.75	
	0	27.90		"	35.04	39.95	
	11.05	29.20		■	26.00	32.16	
	15.80	30.68		"	34.60	38.11	
	21.25	34.25		-10	27.30	32.05	
	14.90	28.35		"	33.56	36.25	
	16.40	29.32		-20	30.08	32.76	
	17.80	27.75		"	32.65	35.44	
	18.80	28.70	.	45	0	58.00	Fe_2Cl_6 anhydrous
	24.50	32.75		"	31.28	50.08	
	24.12	30.04		"	40.65	48.60	
	26.00	32.16		40	0	58.00	
	24.95	29.60		"	27.00	50.80	
	26.05	30.50		"	42.01	48.64	
	27.30	32.05		35	0	58.00	
			$\text{Fe}_2\text{Cl}_6 + 4\text{H}_2\text{O}$	"	29.01	50.33	
70	0	44.0		"	37.04	49.20	
"	6.75	50.00		30	0	58.00	
"	■	55.80		"	32.60	49.93	
65	0	42.50		"	34.40	49.72	
"	10.25	50.00		40	42.50	47.52	$\text{Fe}_2\text{Cl}_6, 2\text{HCl},$ $4\text{H}_2\text{O}$
"	3.75	57.25		"	42.01	48.64	
"	0	59.00		35	39.47	46.57	
60	0	41.40		"	37.04	49.20	
"	14.25	50.00		30	40.21	42.54	
"	10.70	55.25		"	38.20	44.70	
"	■	61.00		"	35.55	47.30	
55	0	40.64		"	34.40	49.72	
"	19.00	50.72		25	40.41	40.25	
"	16.71	53.60		"	39.03	41.38	
"	0	62.00		"	35.74	45.24	
50	0	39.92		20	39.50	39.25	
"	3.25	39.95		"	35.40	43.16	
"	21.24	49.33		10	38.62	37.48	
"	20.04	52.50		"	37.46	38.33	
44	0	39.00		"	36.30	38.70	
"	10.70	38.00		"	35.04	39.93	
"	14.80	38.70		0	37.27	36.60	
"	24.14	50.10		"	34.60	38.11	
40	13.40	37.45		-10	37.92	35.32	
"	27.00	50.80		"	34.54	36.00	
35	15.70	37.06		-20	33.56	36.25	
"	29.20	42.70		"	37.80	34.50	
"	31.08	46.85		"	34.10	34.84	
"	30.81	47.65		"	32.56	35.44	

Solubility of Fe_2Cl_6 in $\text{HCl} + \text{Aq.}$ —Continued.

t°	Sat. solution contains per 100 mols. H_2O		Solid phase
	mols. HCl	mols. FeCl_3	
—4.5	20.50	24.50	$\text{Fe}_2\text{Cl}_6, 2\text{HCl},$ $8\text{H}_2\text{O}$
"	20.66	25.74	
"	23.42	27.40	
—6	29.10	24.73	
"	26.18	21.75	
"	24.41	21.50	
"	23.25	21.35	
"	21.73	21.84	
"	19.73	25.50	
"	24.42	28.45	
"	28.20	27.04	
—10	20.48	20.54	
"	24.90	18.94	
"	28.75	20.34	
"	31.42	28.53	
"	28.25	30.25	
"	26.05	30.50	
—15	24.50	15.83	
—15	28.40	31.89	
—20	19.44	12.10	
"	22.83	11.63	
"	25.20	11.60	
"	27.20	11.31	
"	31.08	11.51	
"	34.13	12.90	
"	33.93	31.77	
"	30.08	32.76	
"	28.70	32.88	
—10	12.01	11.99	$\text{Fe}_2\text{Cl}_6, 2\text{HCl},$ $12\text{H}_2\text{O}$
"	19.78	14.02	
"	20.95	16.20	
"	20.25	20.20	
"	17.73	20.70	
"	15.44	19.65	
—12.5	22.14	16.69	
—15	21.30	9.65	
"	24.50	15.83	
—20	9.96	9.94	
"	13.32	8.57	
"	16.90	7.35	
"	18.97	7.16	
"	20.56	7.08	
"	23.40	7.20	
"	24.85	9.88	
"	25.20	11.60	
"	25.40	12.37	
"	25.59	13.39	

(Rooseboom and Schreinemakers, Z. phys. Ch. 1894, 15, 633.)

Solubility of $\text{Fe}_2\text{Cl}_6 + \text{NH}_4\text{Cl}$.
See $\text{NH}_4\text{Cl} + \text{Fe}_2\text{Cl}_6$ under NH_4Cl .

Solubility of Fe_2Cl_6 in CaCl_2 .
See $\text{CaCl}_2 + \text{Fe}_2\text{Cl}_6$ under CaCl_2 .

Solubility of $\text{FeCl}_3 + \text{KCl}$ in H_2O at 21°.

Substance added		Pta. by weight wt. in 100 pta. of solution	
FeCl_3 grams	KCl grams	FeCl_3	KCl
0	25	0	34.97
13	28	13.44	24.45
18	21	23.18	16.54
3	18.5	28.05	11.66
28	16	35.72	11.19
31	10.5	36.62	13.67
36.2	9	37.35	7.88
41.5	8	42.03	7.54
46.5	6	51.69	0
52	0.5		
155	0	83.89	0

(Hinrichsen and Sachse, Z. phys. Ch. 1904, 50, 95.)

 $\text{FeCl}_3 + \text{NaCl}$.Solubility of $\text{FeCl}_3 + \text{NaCl}$ in H_2O at 21°.

Substance added		Pta. by weight wt. in 100 pta. of solution	
FeCl_3 grams	NaCl grams	FeCl_3	NaCl
0	3.6	0	36.10
1.8	3.0	24.27	9.10
3.6	2.5	25.40	8.45
5.5	2.0	26.40	5.25
7.2	1.5	38.15	
9.0	1.0	43.38	2.45
10.8	0.5	46.75	2.11
10.8	0	83.39	0

(Hinrichsen and Sachse, Z. phys. Ch. 1904, 50, 94.)

Solubility of FeCl_3 in $\text{NaCl} + \text{Aq}$ at t°.

t°	Substance added		γ_{\pm} of Fe in the solution
	FeCl_3	NaCl	
10	100	20	15.2
10	60	20	15.2
10	100	10	15.16
20	60	20	16.2
20	80	10	16.18
20	100	20	16.2
30	70	30	17.7
30	90	30	17.6
30	110	30	17.67
50	30	20	23.5
50	45	20	23.9
40	35	30	25.4
40	50	30	25.5
30	30	20	23.8
30	45	20	24.0
17.6	30	20	24.47
17.6	50	20	24.5

(Hinrichsen and Sachse, Z. phys. Ch. 1904, 50, 95.)

Difficultly sol. in AsBr_3 . (Walden, Z. anorg. 1902, 29. 374.)

Attacked by liquid NO_2 in the presence of traces of moisture. (Frankland, Chem. Soc. Trans. 79. 1361.)

Sol. in liquid SO_2 . (Walden, B. 1899, 32. 54.)

Sol. in alcohol ether, acetic ether (Cann. R. 102. 363), and acetone (Krug and Eroy, J. anal. Ch. 6. 184).

Sol. in ethylamine. (Shinn, J. phys. chem. 1907, 11. 538.)

Sol. in benzonitrile. (Naumann, B. 1914, 1369.)

Sol. in methyl acetate. (Naumann, B. 9, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 9, 43. 314.)

g. FeCl_3 is sol. in 1.59 g. acetone at 18° . gr. of sat. solution $18^\circ/4^\circ = 1.160$. (Naumann, B. 1904, 37. 4333.)

sol. in acetone and in methylal. (Eidmann, C. 1899, II. 1014.)

sol. in quinoline. (Beckmann and Gabel, inorg. 1906, 51. 236.)

l. sol. in CS_2 . (Arctowski, Z. anorg. 1894, 57.)

Vol. weight determined in pyridine. (Arner, Z. anorg. 1897, 15. 22.)

Sublimed.

sol. in AsCl_3 , POCl_3 , SO_2Cl_2 and PBr_3 ; sl. in PCl_3 . (Walden, Z. anorg. 1900, 25. 1.)

The salts with different amts. of crystal water have different solubilities. (Roozeboom. $+4\text{H}_2\text{O}$. Melts in crystal H_2O at 73.5° .)

100 mols. H_2O dissolve mols. Fe_2Cl_6 from $\text{Fe}_2\text{Cl}_6 + 4\text{H}_2\text{O}$ at t° .

t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6
0	19.96	69	21.53	72.5	26.15
5	20.32	72.5	23.35	70	27.90
10	20.70	73.5	25.00	66	29.20

(Roozeboom, Z. phys. Ch. 10. 477.)

$+5\text{H}_2\text{O}$. Correct formula for $+6\text{H}_2\text{O}$ salt.

100 mols. H_2O dissolve mols. Fe_2Cl_6 from $\text{Fe}_2\text{Cl}_6 + 5\text{H}_2\text{O}$ at t° .

t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6
2	12.87	30	15.12	55	19.15
0	13.95	35	15.64	56	20.00
7	14.85	50	17.50	55	20.32

(Roozeboom.)

Melts in crystal H_2O at 31° (Engel, C. R. 1708); at 56° (Roozeboom).

$+6\text{H}_2\text{O}$. Very deliquescent. Sol. in alcohol. Ether dissolves out Fe_2Cl_6 .

M.-pt. is 31° . (Ordway.) Contains only $5\text{H}_2\text{O}$. (Roozeboom.)

$+7\text{H}_2\text{O}$. Melts in crystal H_2O at 32.5° .

100 mols. H_2O dissolve mols. Fe_2Cl_6 from $\text{Fe}_2\text{Cl}_6 + 7\text{H}_2\text{O}$ at t° .

t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6
20	11.35	32	13.55	30	15.12
27.4	12.15	32.5	14.99	25	15.54

(Roozeboom.)

$+12\text{H}_2\text{O}$. Less deliquescent than Fe_2Cl_6 or $\text{Fe}_2\text{Cl}_6 + 5\text{H}_2\text{O}$.

100 mols. H_2O dissolve mols. Fe_2Cl_6 from $\text{Fe}_2\text{Cl}_6 + 12\text{H}_2\text{O}$ at t° .

t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6	t°	Mols. Fe_2Cl_6
-55	2.75	30	5.93	27.4	11.20
-41	2.81	35	6.78	20	12.15
-27	2.98	36.5	7.93	10	12.83
0	4.13	37	8.33	8	13.70
10	4.54	36	9.29
20	5.10	30	10.45

(Roozeboom.)

Sol. in alcohol. Ether dissolves out Fe_2Cl_6 . Melts in crystal H_2O at 37° (Roozeboom); at 35.5° (Ordway).

Ferric hydrogen chloride, $\text{FeCl}_3, \text{HCl} + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Sabatier, Bull. Soc. (2) 197.)

More sol. in H_2O than FeCl_3 . (Engel, C. R. 104. 1708.)

For solubility, see $\text{FeCl}_3 + \text{HCl}$, under ferric chloride.

$+6\text{H}_2\text{O}$. (Roozeboom and Schreinemakers.)

For solubility, see $\text{FeCl}_3 + \text{HCl}$, under ferric chloride.

Ferrous lithium chloride, $\text{FeCl}_2, \text{LiCl} + 3\text{H}_2\text{O}$.

(Chassevant, A. ch. (6) 30. 17.)

Ferric magnesium chloride, $\text{FeCl}_3, \text{MgCl}_2 + \text{H}_2\text{O}$.

Deliquescent. (Neumann, B. 18. 2890.)

Ferrous mercuric chloride, $\text{FeCl}_2, \text{HgCl}_2 + 4\text{H}_2\text{O}$.

Deliquescent. (v. Bonsdorff.)

Ferric nitrosyl chloride, $\text{FeCl}_3, \text{NOCl}$.

Very deliquescent. (Weber, Pogg. 118. 477.)

Ferric phosphoric chloride, $\text{FeCl}_3, \text{PCl}_5$.

Decomp. by H_2O . (Baudrimont, A. ch. (4) 2. 15.)

Iron (ferrous) potassium chloride, FeCl_2 , $2\text{KCl} + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Berzelius.)

Ferric potassium chloride, FeCl_3 , $2\text{KCl} + \text{H}_2\text{O}$.

A little H_2O dissolves out FeCl_3 . (Fritzsche J. pr. 18. 483.)

Sol. in H_2O . (Walden, Z. anorg. 1894, 71. 332.)

Ferric rubidium chloride, FeCl_3 , 3RbCl .

Easily sol. in H_2O . Insol. in $\text{HCl} + \text{Aq}$. (Godeffroy, Arch. Pharm. (3) 9. 343.)

FeCl_3 , $2\text{RbCl} + \text{H}_2\text{O}$. Decomp. by H_2O . (Neumann, A. 244. 329.)

Sol. in H_2O . (Walden, Z. anorg. 1894, 7. 332.)

Ferric sulphur chloride, FeCl_3 , 8SCl_4 .

Very sensitive toward heat and moisture. (Ruff, B. 1904, 37. 4518.)

Ferric thallium chloride, FeCl_3 , 3TlCl .

Decomp. by H_2O . Can be crystallised from $\text{HCl} + \text{Aq}$. (Wöhler, A. 144. 250.)

Ferrous chloride ammonia, 3FeCl_2 , 2NH_3 .

Decomp. by H_2O . (Rogstadius, J. pr. 86. 310.)

FeCl_2 , 6NH_3 . Loses 4NH_3 at 100° . (Miller, Am. Ch. J. 1895, 17. 577.)

FeCl_2 , 2NH_3 . Decomp. in the air (Miller).

Ferric chloride ammonia, FeCl_3 , NH_3 .

Slowly deliquescent. Sol. in H_2O with evolution of heat. (Rose, Pogg, 24. 302.)

FeCl_3 , 6NH_3 . Not deliquescent; not sol. in H_2O ; sol. in HCl with decomp. (Miller, Am. Ch. J. 1895, 17. 577.)

Loses NH_3 to give FeCl_3 , 5NH_3 , and FeCl_2 , 4NH_3 .

Ferric chloride cyanhydric acid, FeCl_3 , 2HCN .

Deliquescent. (Klein, A. 74. 85.)

Ferrous chloride nitric oxide, FeCl_2 , NO .

Sol. in H_2O without evolution of gas. (Thomas, C. R. 1895, 121. 204.)

$+2\text{H}_2\text{O}$. Sol. in cold H_2O without decomp. (Thomas, C. R. 1895, 120. 448.)

2FeCl_2 , NO . Very hygroscopic. (Thomas, C. R. 1895, 121. 129.)

10FeCl_2 , NO . Very hygroscopic. (Thomas C. R. 1895, 121. 128.)

Ferric chloride nitric oxide, Fe_2Cl_6 , NO .

Very hygroscopic. Loses NO when exposed to the air.

$2\text{Fe}_2\text{Cl}_6$, NO . Very hygroscopic. In contact with H_2O gives off NO . (Thomas, C. R. 1895, 120. 447.)

Iron (ferrous) fluoride, FeF_2 .

Sl. sol. in H_2O ; insol. in alcohol and Partly sol. in hot $\text{HCl} + \text{Aq}$; slowly cold, easily in hot HNO_3 ; decomp. by (Poulenc, C. R. 115. 941.)

$+8\text{H}_2\text{O}$. Difficultly sol. in H_2O easily if it contains HF . (Berzelius.)

Ferroferric fluoride, FeF_3 , $\text{FeF}_2 + 7\text{H}_2\text{O}$.

Sol. in dil. $\text{HF} + \text{Aq}$. (Weinland, Z. 1899, 22. 268.)

Ferric fluoride, FeF_3 .

Sl. sol. in H_2O ; insol. in alcohol (Sl. attacked by HNO_3 , HCl , or H_2S (Poulenc, C. R. 115. 941.)

$+4\frac{1}{2}\text{H}_2\text{O}$. More sol. in hot than cold. Insol. in alcohol. (Scheurer-Kestner (3) 68. 472.)

Ferric nickel fluoride, FeF_3 , $\text{NiF}_2 + 7\text{H}_2\text{O}$.

Sl. sol. in dil. $\text{HF} + \text{Aq}$. (Weinland, Z. anorg. 1899, 22. 268.)

Ferrous potassium fluoride, FeF_2 , KF .

(Wagner, B. 19. 896.)

FeF_2 , 2KF . Sl. sol. in H_2O . (Berzelius.)

Ferric potassium fluoride, FeF_3 , 2KF .

Somewhat sol. in H_2O , especially (Berzelius.)

$+ \text{H}_2\text{O}$. (Christensen, J. pr. (2) 35.)

FeF_3 , 3KF . Properties as above. (Berzelius.)

Ferric sodium fluoride, FeF_3 , $2\text{NaF} + 7\text{H}_2\text{O}$.

Rather easily sol. in H_2O . Solut. comp. on heating. Very sol. in FeCl_3 (Nicklès, J. Pharm. (4) 10. 14.)

FeF_3 , 3NaF . (Wagner, B. 19. 896.)

Ferric thalious fluoride, 2FeF_3 , 3TlF .

Sol. in hot H_2O , less sol. in cold. Sol. in HF . (Ephraim, Z. anorg. 19. 239.)

Ferrous titanium fluoride.

See Fluotitanate, ferrous.

Ferric zinc fluoride, FeF_3 , $\text{ZnF}_2 + 7\text{H}_2\text{O}$.

Sl. sol. in dil. $\text{HF} + \text{Aq}$. (Weinland, Z. anorg. 1899, 22. 269.)

Ferrous hydroxide, FeO_2H_2 .

Sol. in 150,000 pts. H_2O . (Bineau, C. R. 41. 509.)

Insol. in KOH , or $\text{NaOH} + \text{Aq}$. NH_4 salts $+ \text{Aq}$. Sl. sol. in $\text{Na}_2\text{C}_2\text{H}_3\text{O}_7$ (Mercer.)

Not pptd. in presence of Na citrate . in boiling cane sugar $+ \text{Aq}$, but sl. sol. if KOH has been added. Not pptd. in presence of much $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$. (Rose.)

solubility in glycerine+Aq containing
out 60% by vol. of glycerine.
100 ccm. of the solution contain 1.0 g. FeO .
(Müller, Z. anorg. 1905, 43. 322.)

(ferric) hydroxides, $\text{Fe}_2\text{O}_3 \cdot x\text{H}_2\text{O}$.

Many indefinite compounds of Fe_2O_3 and
H₂O are known, and uncertainty exists as to
their composition.

According to van Bemmelen (R. t. c. 7. 106)
there are probably no true definite compounds
 Fe_2O_3 and H_2O .

According to Tommasi (B. 12. 1924, 2334),
there are two series of Fe hydroxides, α , red
hydroxides, and β , yellow hydroxides.

Hydroxides. $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ (unstable), $\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$ (loses H_2O at 50°), and $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ (loses
H₂O at 92°).

sol. in dil. acids and in Fe_2Cl_6 +Aq, and
d. from the latter solution by Na_2SO_4 or
 K_2SO_4 +Aq.

Hydroxides. $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ (stable below 70°),
 O_2 , $2\text{H}_2\text{O}$ (loses H_2O at 105°), $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$
(loses H_2O at 150°).

l. sol. in acids, and insol. in Fe_2Cl_6 +Aq.
(Tommasi.)

The following more or less uncertain data
given.

$\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$. Sol. in HCl +Aq. Very sl.
in HNO_3 +Aq. (Davies, Chem. Soc.
4. 69.)

Min. *Turgite*.

$\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$. Insol. in cold acids, difficultly
in warm HCl and H_2SO_4 +Aq, and especi-
ally in warm HNO_3 +Aq. (Schiff, A. 114. 199.)

Min. *Göthite*.

$\text{Fe}_2\text{O}_3 \cdot 3\text{H}_2\text{O}$. Sl. sol. in tartaric, citric,
acetic acids, but easily sol. in HCl +Aq.
(Wittstein.)

Scarcely attacked by conc. HNO_3 , or HCl +
Aq. Sol. in acetic acid or dil. HNO_3 , or HCl +
Aq, from which solution it is pptd. by trace of
alkali salts. (St. Gilles.)

Min. *Limonite*.

$3\text{Fe}_2\text{O}_3 \cdot 5\text{H}_2\text{O}$. (Muck.)

$\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$. Easily sol. in HCl +Aq.

Min. *Xanthosiderite*.

$\text{Fe}_2\text{O}_3 \cdot 3\text{H}_2\text{O}$. Sl. sol. in acetic acid of 1.03
gr., but easily sol. if of 1.076 sp. gr. Sol.
mineral acids. (Limberger, J. B. 1883. 70.)

Pptd. $\text{Fe}_2\text{O}_3 \cdot x\text{H}_2\text{O} = \text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ (?). Insol.
H₂O, or in solutions of the alkalis or NH_4
salts. When recently pptd. is easily sol. in
acids. (Fresenius.)

Sl. sol. in NH_4OH , and NH_4 salts+Aq.
(Lind.)

Apparently insol. in NH_4Cl , or $(\text{NH}_4)_2\text{CO}_3$
+Aq. (Brett, 1837.)

Sl. sol. in conc., but insol. in dil. KOH +Aq.
(Hodnew, J. pr. 23. 221.)

Sl. sol. in very conc. KOH +Aq free from
H₂O. (Völcker, A. 59. 34.)

Not at all sol. in pure conc. KOH +Aq,
solubility noticed by previous observers being
masked by the presence of silicic acid. (Sand-
ström.)

Sl. sol. in conc. alkali carbonates+Aq.

When freshly pptd., it is not acted upon by
conc. K_2CO_3 +Aq. (Grotthaus.)

Readily sol. in conc. $(\text{NH}_4)_2\text{CO}_3$ +Aq, but
pptd. by addition of H_2O .

Sol. in excess of $(\text{NH}_4)_2\text{CO}_3$ +Aq when
pptd. by that reagent. (Wohler.)

Sol. in solutions of the alkali bicarbonates.
(Berzelius.)

Sol. in aqueous solutions of water-glass.
(Ordway.)

Immediately dissolved by H_2SO_4 +Aq.

Sol. in NH_4F +Aq. (Helmholtz, Z. anorg.
3. 124.)

Sol. in conc. $\text{Al}_2(\text{SO}_4)_3$ +Aq. (Schneider,
B. 23. 1352.)

Sl. sol. in a solution of MgCO_3 (?). (Bis-
choff.)

Insol. in ethylamine, or amylamine+Aq.
(Wurts, A. ch. (3) 30. 472.)

Sol. in boiling solution of $\text{Bi}(\text{NO}_3)_3$, with
pptn. of Bi_2O_3 . (Persoz.)

Sol. in Cr_2Cl_6 +Aq; after 3 months 15 mols.
 $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ were dissolved by 1 mol. Cr_2Cl_6 .
(Béchamp, A. ch. (3) 57. 296.)

Insol. in fumaric acid, even when freshly
pptd.

When recently pptd., it is easily sol. in
 $\text{KHC}_4\text{H}_4\text{O}_6$ +Aq, but after drying it is dif-
ficultly sol. therein.

When moist easily sol. in $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$ +Aq,
but after drying is scarcely sol. therein when
cold, and only sl. sol. when hot. (Werther.)

Easily sol. in acetic, citric, and other acids.
(Wittstein.)

Solubility in glycerine+Aq containing
about 60% by vol. of glycerine.

100 ccm. of the solution contain 0.8 g.
 Fe_2O_3 . (Müller, Z. anorg. 1905, 43. 322.)

Easily sol. in aqueous solution of sucra-
tes of Ca, Ba, Sr, K, Na. (Hunton, 1837.)

Unacted upon by cane sugar+Aq. (Glad-
stone.)

Sl. sol. in cane sugar+Aq, from which it is
pptd. by $(\text{NH}_4)_2\text{S}$ +Aq, but not by NH_4OH ,
or $\text{K}_4\text{FeC}_6\text{N}_8$ +Aq. (Peschier.)

Solubility of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ in sugar solutions. 1 l.
of sugar solution of given strength dis-
solves mg of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$.

% Sugar	Mg. $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$		
	at 17.4°	at 45°	at 75°
10	3.4	3.4	6.1
30	2.3	2.7	3.8
50	2.3	1.9	3.4

(Stolle, Z. Ver. Zuckerind. 1900, 50. 340.)

Not pptd. from solutions by alkalis or
alkali carbonates in presence of many organic
substances, as tartaric acid, sugar, etc.

Not pptd. by NH_4OH from solutions con-
taining $\text{Na}_4\text{P}_2\text{O}_7$. (Rose, Pogg. 76. 19.)

Not pptd. by NH_4OH in presence of Na
citrate. (Spiller.)

Soluble. (a) *By dialysis.* Solutions containing 1% can be concentrated somewhat, whereupon they gelatinise. They also gelatinise by cold, or addition of traces of H_2SO_4 , alkalis, alkali carbonates or sulphates, or neutral salts, not, however, by HCl , HNO_3 , alcohol, or sugar. (Graham, A. 121. 46.)

When a dil. solution of a solid organic acid, or an alkali, or salt is added to a dialysed solution of $\text{Fe}_2\text{O}_3\cdot\text{H}_2\text{O}$, a coagulum sol. in H_2O is formed, but if the solutions are conc. the separating coagulum is no longer sol. in H_2O . (Athenstädt, C. C. 1871. 822.)

(b) *Pean St. Gilles' hydroxide, or meta-iron hydroxide.* Sol. in H_2O . Pptd. from solution by traces of H_2SO_4 , HCl , HNO_3 + Aq, and alkalis; the ppt. is insol. in cold acids, but sol. in pure H_2O . (Pean St. Gilles, A. ch. (3) 46. 47.)

See also table by Krecke in the article on ferric chloride.

Iron (Ferroferric) hydroxide, $\text{Fe}_2\text{O}_3\cdot\text{H}_2\text{O}$ (?).

Sol. in acids.

$\text{Fe}_2\text{O}_3\cdot 4\text{H}_2\text{O}$. (Lefort.)

Ferrous iodide, FeI_2 .

Very deliquescent. Sol. in H_2O . Solution decomp. on evaporating.

+ $4\text{H}_2\text{O}$. Very deliquescent; sl. sol. in H_2O ; sol. in ether. (Jackson, Am. Ch. J. 1900, 24. 19.)

+ $5\text{H}_2\text{O}$. Deliquescent. Sol. in alcohol. Sol. in sugar + Aq, and solution is much more stable than aqueous solution. Easily sol. in glycerine.

Insol. in methylene iodide. (Retgers, Z. anorg. 3. 343.)

+ $6\text{H}_2\text{O}$, and + $9\text{H}_2\text{O}$. Very sol.; pptd. from cooled aq. solution. (Volkmann, C. C. 1894, II. 611.)

Ferric iodide, FeI_3 .

Has not been isolated. Solution of I in FeI_2 + Aq in the molecular ratio of I : FeI_2 , probably contains FeI_3 .

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Ferrous mercuric iodide, $\text{FeI}_2\cdot 2\text{HgI}_2\cdot 6\text{H}_2\text{O}$.

As the corresponding Mg salt. (Duboin, C. R. 1907, 145. 714.)

Ferrous iodide ammonia, $\text{FeI}_2\cdot 6\text{NH}_3$.

Decomp. by H_2O . (Jackson, Am. Ch. J. 1900, 24. 27.)

Ferrous mercuric iodide.

Very deliquescent. Decomp. by H_2O ; sol. in $\text{HC}_2\text{H}_3\text{O}_2$, or alcohol.

Iron molybdenide, FeMo_2 .

Attacked by HCl + Aq with difficulty. Sol. in hot conc. H_2SO_4 . (Steinacker.)

Iron nitride.

Easily decomp. by H_2O when fine dered. (Rossel, C. R. 1895, 121. 942.)

Fe_3N . Easily sol. in HNO_3 , HCl , + Aq. Very slowly decomp. by H_2O . (Schmidt, Pogg. 125. 37.)

Sol. in HCl with decomp.; dec. steam and by H_2S at 200° . (Fowle 1894, 68. 152.)

Fe_4N_2 . Probably the same as the compound. (Rogstadius, J. pr. 86. 3)

Iron nitrososulphantimonate, $\text{Fe}_2\text{S}(\text{N})$
(Low, C. C. 1865. 948.)

Does not exist, but was impure so rotetranitrososulphide. (Pawel, B. 1)

Iron nitrososulphides.

See Ferrotetranitrososulphydic and Ferroheptanitrososulphide, ammonio

$\text{Fe}_2\text{S}_2\text{H}_2(\text{NO})_4$. (Roussin, C. R. 4
 $\text{Fe}_2\text{S}_2(\text{NO})_4\cdot 2\text{H}_2\text{O}$. (Porczynsky 302.)

$\text{Fe}_2\text{S}_2(\text{NO})_{10}\cdot 4\text{H}_2\text{O}$. (Rosenberg 312.)

The compound to which the above were given was impure, according to 12. 1407 and 1949; 15. 2600), and more or less Na or NH_4 . Pawel contains substance as NH_4 salt of ferroheptasulphydic acid, which see.

$\text{Fe}_2\text{S}_2\text{N}_2\text{O}_8\cdot 1\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2C ether, CHCl_3 , acetone and ethyl. Insol. in benzene and light petroleum. C. R. 1896, 122. 138.)

Iron sodium nitrososulphide, $3\text{Na}_2\cdot 2\text{NO}$.

(Roussin.)

$\text{Na}_2\text{Fe}_2\text{S}_2(\text{NO})_{10}$. (Rosenberg.)

Correct formula is $\text{Na}_2\text{S}_2(\text{NO})_4\text{Fe}$ ferrotetranitrososulphide.

Iron nitrososulphocarbonate, $\text{Fe}_2\text{S}(\text{N})\cdot 3\text{H}_2\text{O}$.

(Low, C. C. 1865. 948.)

Correct formula is $\text{NaS}_2(\text{NO})_4\text{Fe}$ sodium ferroheptanitrososulphide. B. 15. 2600.)

Ferrous oxide, FeO .

Insol. in H_2O . Sol. in acids.

Easily sol. in HCl , and HNO_3 + Aq insol. in H_2SO_4 , even when heated. (Dier, C. R. 74. 531.)

Ferric oxide, Fe_2O_3 .

Attacked by acids with difficulty, so the higher it has been heated. H_2O is the best solvent, in which it is most sol. by long digestion at a gentle heat by boiling. (Fresenius.)

Most easily sol. in 16 pts. of a mix of 8 pts. H_2SO_4 and 3 pts. H_2O . (Mit J. pr. 81. 110.)

bility of Fe₂O₃ in HF + Aq at 25°.

	Time	G. Fe ₂ O ₃ in 10 ccm. of the solution
-HF	4½ hrs.	0.1581
	21¾ "	0.2235
	45¾ "	0.2279
N-HF	2¾ "	0.0579
	8½ "	0.0884
	23½ "	0.1045
	56½ "	0.1162
N-HF	2¼ "	0.0180
	8¼ "	0.0345
	24¾ "	0.0475
	142½ "	0.0534
amts. N-HCl	2¾ "	0.1011
	8½ "	0.1611
	23¾ "	0.1976
	96 "	0.2223
	264 "	0.2297

ussen, Z. anorg. 1905, 44. 414.)

ility of Fe₂O₃ in HCl + Aq at 25°.

	Time	G. Fe ₂ O ₃ in 10 ccm. of the solution
HCl	4¾ hrs.	0.0409
	21½ "	0.1230
	45½ "	0.2125
-HCl	2¾ "	0.0126
	8½ "	0.0188
	23½ "	0.0382
	56½ "	0.0672
-HCl	2¼ "	0.0040
	8½ "	0.0054
	24¾ "	0.0120
	142½ "	0.0306
vol. N-NaF	2¾ "	0.0444
	8½ "	0.0640
	23¾ "	0.0743
	72¼ "	0.0757
	215 "	0.0766

(Deussen, l. c.)

y of Fe₂O₃ in N-oxalic acid at 25°.

Time	G. Fe ₂ O ₃ in 10 ccm. of the solution
1 hrs.	0.0310
1 "	0.0790
2 "	0.1960
1 "	0.2326

(Deussen.)

Absolutely insol. in Br₂ + Aq. (Balard.)
Insol. in hot NH₄Cl + Aq. (Rose.)
Insol. in KOH + Aq. (Chodnew, J. pr. 28. 222.)
Slowly sol. in an aq. solution of calcium hydrogen carbonate. The velocity of the reaction may be much increased by the addition of small amounts of alkali sulphate or CaSO₄. (Rohland, Z. anal. 1909, 48. 629.)
Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)
Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)
Solubility in (calcium succate + sugar) + Aq. 1 l. solution containing 418.6 g. sugar and 34.3 g. CaO dissolves 6.26 g. Fe₂O₃; 296.5 g. sugar and 24.2 g. CaO dissolves 4.71 g. Fe₂O₃; 174.4 g. sugar and 14.1 g. CaO dissolves 3.08 g. Fe₂O₃. (Bodenbender, J. B. 1865. 600.)

Solubility of Fe₂O₃ in sugar solutions. 1 l. of sugar solution of given strength dissolves mg. Fe₂O₃.

% Sugar	mg. Fe ₂ O ₃	
	at 17.5°	at 45°
10	1.4	2.0
30	1.4	...
50	0.8	1.1

(Stolle, Z. Ver. Zuckerind, 1900, 50. 340.)

Calcined.
Solubility of calcined Fe₂O₃ in acids at 25°.

Acid	Time	g. Fe ₂ O ₃ in 10 ccm. of the solution
N-HF	4½ hrs.	0.0889
	43½ "	0.2035
	129½ "	0.2194
N-HCl	4½ "	0.0224
	43½ "	0.1000
	139½ "	0.1910

(Deussen, Z. anorg. 1905, 44. 413.)

See also Ferric hydroxide.
Min. Hematite. Rather easily sol. in HCl + Aq, but not readily sol. in other acids.

Metairon oxide.
See Ferric hydroxides.

Ferroferric oxide, 6FeO, Fe₂O₃.
FeO, Fe₂O₃ = Fe₃O₄. With insufficient HCl + Aq for complete solution, FeO is dissolved and Fe₂O₃ left. (Berzelius.)
Insol. in HNO₃ + Aq at the ordinary temperature. (Millon.)
Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Solubility of Fe_2O_3 in sugar solutions. 1 l. of sugar solution of given strength dissolves mg. Fe_2O_3 .

% Sugar	mg. Fe_2O_3		
	at 17.5°	at 45°	at 75°
10	10.3	10.3	12.4
30	12.4	10.3	12.4
50	14.5	10.3	14.5

(Stolle, Z. Ver. Zuckerind. 1900, 50, 340.)

Min. Magnetite. Insol. in HNO_3 , but sol. in hot $\text{HCl} + \text{Aq}$.

Iron sesquioxide zinc oxide, $\text{Fe}_2\text{O}_3 \cdot \text{ZnO}$.

See Ferrite, zinc.

Ferric oxybromide.

Basic ferric bromides containing three equivalents, or less, of base to one of acid may be obtained dissolved in H_2O . (Ordway, Am. J. Sci. (2) 26, 202.)

The most basic soluble compound obtained by three months' digestion of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ with $\text{Fe}_2\text{Br}_6 + \text{Aq}$, is $\text{Fe}_2\text{Br}_6 \cdot 14\text{Fe}_2\text{O}_3$. (Béchamp.)

Ferric oxychlorides.

(a) Soluble. $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ dissolves in $\text{Fe}_2\text{Cl}_6 + \text{Aq}$. By digesting until the acid reaction of the chloride has disappeared a solution of $\text{Fe}_2\text{Cl}_6 \cdot 2\text{Fe}_2\text{O}_3$ is obtained. (Pettenkofer, Rept. (2) 41, 289.)

By digesting for several days in the cold, $\text{Fe}_2\text{Cl}_6 \cdot 5\text{Fe}_2\text{O}_3$ is obtained, and still more basic compounds by further addition of $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$. When the solution contains $\text{Fe}_2\text{Cl}_6 \cdot 12\text{Fe}_2\text{O}_3$ it gelatinizes, but still dissolves completely in H_2O . The most basic soluble compound is $\text{Fe}_2\text{Cl}_6 \cdot 20\text{Fe}_2\text{O}_3$. (Béchamp, A. ch. (3) 57, 296.)

If the digestion is carried on several weeks, a solution containing $\text{Fe}_2\text{Cl}_6 \cdot 23\text{Fe}_2\text{O}_3$ is obtained; this can be boiled and diluted without pptn., but $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$ is precipitated by the addition of very many salts. (Ordway, Sill. Am. J. (2) 26, 197.)

Solutions containing 10 or less molecules Fe_2O_3 to 1 mol. Fe_2Cl_6 can be dried without the oxychloride becoming insoluble. (Ordway.)

The above solutions do not become cloudy by boiling or diluting. (Phillips.)

A very dil. solution of $\text{Fe}_2\text{Cl}_6 \cdot 10\text{Fe}_2\text{O}_3$ remains clear after protracted boiling, and may be boiled without decomp. even when $\text{Fe}_2\text{Cl}_6 \cdot 20\text{Fe}_2\text{O}_3$ is present. (Béchamp.)

HNO_3 , and $\text{HCl} + \text{Aq}$ form precipitates in the above solutions, which are sol. on addition of more H_2O . $\text{H}_2\text{SO}_4 + \text{Aq}$ forms a precipitate insol. in H_2O . (Béchamp.)

$\text{Fe}_2\text{Cl}_6 \cdot 9\text{Fe}_2\text{O}_3$ is easily sol. in H_2O , weak alcohol, and glycerine; but solutions are pptd. by small amts. of H_2SO_4 , M_2SO_4 , citric or

tartaric acids, or a few drops of $\text{HCl} + \text{HNO}_3 + \text{Aq}$. (Jeannel, C. R. 48, 790.)

Solutions containing 5 mols. Fe_2O_3 to Fe_2Cl_6 are completely precipitated by 1 Na_2SO_4 , MgSO_4 , KNO_3 , NaNO_3 , ZnCl_2 , KCl , NaCl , NH_4Cl , CaCl_2 , MgCl_2 , KBr , or KSCN . (Béchamp.)

$\text{Ba}(\text{NO}_3)_2$ does not precipitate sol. less than 18–20 Fe_2O_3 to 1 Fe_2Cl_6 .

$\text{Pb}(\text{NO}_3)_2$ or $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2$ do not taste solutions containing the cor. $\text{Fe}_2\text{Cl}_6 \cdot 12\text{Fe}_2\text{O}_3$, but a mixture of 1 salts causes complete precipitation.

Solution has been obtained contain Fe_2O_3 to 1 Fe_2Cl_6 , probably owing to a tion of soluble colloidal Fe_2O_3 . (Maj la Source, C. R. 90, 1352.)

Solubility determinations in the Fe_2O_3 , HCl and H_2O , show that at definite basic chloride is formed, but 1 stable solid phase is one of a series solutions containing Fe_2O_3 , HCl an (Cameron, J. phys. Chem. 1907, 11, 6)

(β) Insoluble. $\text{Fe}_2\text{Cl}_6 \cdot 6\text{Fe}_2\text{O}_3 + 9\text{H}_2\text{O}$

(1) By exposing $\text{Fe}_2\text{Cl}_6 + \text{Aq}$ to air. in H_2O : sl. sol. in $\text{HCl} + \text{Aq}$. (Wittat

(2) From $\text{Fe}_2\text{Cl}_6 + \text{Aq}$ and HNO_3 in H_2O , and sl. sol. in $\text{HCl} + \text{Aq}$. (Bé $2\text{Fe}_2\text{Cl}_6 \cdot 25\text{Fe}_2\text{O}_3 + 41\text{H}_2\text{O}$. Insol. (Béchamp.)

$\text{Fe}_2\text{Cl}_6 \cdot 2\text{Fe}_2\text{O}_3 + 3\text{H}_2\text{O}$. Decomp. 1 with residue of Fe_2O_3 ; sl. sol. in dil (Rousseau, C. R. 110, 1032.)

$\text{Fe}_2\text{Cl}_6 \cdot 3\text{Fe}_2\text{O}_3$. As above. (R C. R. 113, 542.)

Ferric oxyfluoride, $3\text{Fe}_2\text{O}_3 \cdot 2\text{FeF}_6 + 4\text{H}_2\text{O}$ Ppt. (Scheurer-Kestner.)

Ferric oxysulphide, $\text{Fe}_2\text{O}_3 \cdot 3\text{Fe}_2\text{S}_3$. (Rammelsberg.)

Iron phosphide, FeP .

Very slowly (Freese), not (Hvoslef, 99) sol. in hot $\text{HCl} + \text{Aq}$. Still more 1 dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Freese.)

Slowly sol. in $\text{HNO}_3 + \text{Aq}$, and on in aqua regia. (Struve.)

Insol. in ammonium citrate + Aq ; in HCl . (Dennis, J. Am. Chem. So 16, 483.)

Fe_3P . Slowly but completely sol. or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in hot conc. in HNO_3 , and in aqua regia. (Freese 132, 225.)

Insol. in all acids except in a m e HNO_3 , and HF . (Maronneau, C. I 130, 657.)

Fe_2P_2 . Very slowly sol. in hot conc Aq . 0.1 g. dissolves by 4 days' heat $\text{HCl} + \text{Aq}$. 0.3 g. dissolves in hot conc in 1 $\frac{1}{2}$ hours; 0.4 g. in 2 hours in HNO Quite easily sol. in aqua regia on w (Freese)

Fe_2P_4 . Insol. in HCl , HNO_3 , a e regia. Sol. in potassium hypobromi

r, Bull. Soc. 1896, (3) 15.

slowly sol. in boiling HCl + ol. in HNO₃ or aqua regia. 1860. 77.)

reese, Pogg. 132. 225.)

. in aqua regia. Sol. in fused (er.)

ly insol. in dil. acids; rapidly r aqua regia; decomp. by conc. +Aq. (Schneider, J. B. 1886.

iron phosphides described the as been established for only Fe₃P.

n conc. HCl.

in hot aqua regia. Insol. in Le Chatelier, C. R. 1909, 149.

Fe₃Se.

l by HNO₃ or acetic acid. Sl. conc. HCl. Readily attacked

. Sol. in HF. (Vigouroux, l. 829.)

. Sol. in HCl, HNO₃, or Insol. in alkalies, or (NH₄)₂S

J. Pharm. (4) 9. 173.) in dil. HCl, or HNO₃ + Aq with

Se. Sol. in conc. HNO₃ + Aq. 211.)

decomp. by fuming HNO₃. 1, C. R. 1900, 130. 1711).

decomp. by fuming HNO₃. 1, C. R. 1900, 130. 1711.)

l. in conc. HCl; decomp. by (Fonzes-Diacon, C. R. 1900,

e₃Si.

l. in HCl + Aq; easily sol. even l. (Hahn, A. 129. 57.)

easily sol. in conc. HCl and idily sol. in HF. (Moissan, l. 623.)

. in hot HCl + Aq only when vdered. (Hahn.)

ttacked by conc. HF or H₂SO₄.

HF. (de Chalmot, Am. Ch. J.

estioned by Jouve, (Bull. Soc. 93).

in HF and in fused KNO₃ and Chalmot, J. Am. Chem. Soc.

ide, Fe₂S.

ids with decomposition. (Arf- l. 72.)

le, FeS.

dil. acids, with evolution of ut separation of S, except with

+xH₂O. Sl. sol. in H₂O, especially if hot. (Berzelius.)

1 l. H₂O dissolves 70.1 x 10⁻⁴ moles FeS at 18°. (Weigel, Z. phys. Ch. 1907, 58. 294.)

Very violently decomp., even by dil. acids. Sol. in H₂SO₄ + Aq. Insol. in H₂S, or (NH₄)₂S + Aq. Sl. sol. in Na₂S, or K₂S + Aq. Sol. in Na₂S or K₂S + Aq. (de Koninck, Z. angew. Ch. 1891. 204.)

Insol. in NH₄NO₃, or NH₄Cl + Aq. (Brett.)

Not completely pptd. in presence of Na citrate. (Spiller.)

Contrary to assertion of Persoz, it can be nearly completely pptd. in presence of Na₄P₂O₇ by (NH₄)₂S + Aq. (Rose, Pogg. 76. 18.)

Sol. in alkali sulpho-molybdates, -tungstates, -vanadates, -arsenates, -antimonates, and -stannates. (Storch, B. 16. 2015.)

Sol. in KCN + Aq.

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 828.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Solubility of FeS in sugar solutions. 1 l. sugar of given strength dissolves mg. FeS.

% Sugar	mg. FeS		
	at 17.5°	at 45°	at 75°
10	3.8	3.8	5.3
30	7.1	9.1	7.2
50	9.9	19.8	9.1

(Stolle, Z. Ver. Zuckerind. 1900, 50. 300.)

Colloidal.—A very dilute solution has been obtained which coagulated very readily. (Winssinger, Bull. Soc. (2) 49. 452.)

Ferric sulphide, Fe₂S₃.

Decomp. by dil. HCl, or H₂SO₄ + Aq with evolution of H₂S, leaving a residue of FeS₂.

+1½H₂O. Sol. in NH₄OH + Aq, also in alcoholic ammonia. Sl. sol. in (NH₄)₂S + very dil. Na₂S₂O₃ + Aq. (Phipson, C. N. 30. 139.)

Iron disulphide, FeS₂.

Insol. in dil. HCl, or H₂SO₄ + Aq. Decomp. by HNO₃ or aqua regia with separation of S. Insol. in a 10% solution of alkali sulphide.

Min. *Pyrite, Marcasite*. Sol. in a mixture of Na₂S and NaOH + Aq, Na₂S + Aq, or mixture of Na₂S and NaSH + Aq; insol. in cold NaSH + Aq. Marcasite is more easily sol. in above than pyrite. (Becker, Sill. Am. J. (3) 33. 199.)

Ferroferric sulphide, Fe₃S₄ or Fe₇S₈.

Min. *Pyrrhotite*. Sol. in dil. acids with a residue of S. Extremely slowly sol. in a 10% solution of alkali sulphides. (Terreil, C. R. 69. 1360.)

Iron (ferrous) nickel sulphide, 2FeS , NiS .

Min. *Pentlandite*.

Ferrous phosphorus sulphide, FeS , P_2S_3 .

(Berzelius.)

2FeS , P_2S_3 . Slowly decomp. by H_2O . Insol. in boiling $\text{HCl} + \text{Aq}$; decomp. by aqua regia. (Berzelius, A. 46. 256.)

Iron potassium sulphide (potassium sulphoferrite), $\text{K}_2\text{Fe}_2\text{S}_4 = \text{K}_2\text{S}$, Fe_2S_3 .

Insol. in cold or hot H_2O . Violently attacked by dil. acids. Not decomp. by boiling with alkalis, alkali carbonates, or sulphides + Aq . Decomp. by KCN , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Preis, J. pr. 107. 16.)

K_2S , 2FeS . (Schneider, Pogg. 136. 460.)

Iron silver sulphide (silver sulphoferrite), Ag_2S , Fe_2S_3 .

Not attacked by dil. $\text{HCl} + \text{Aq}$; decomp. by conc. $\text{HCl} + \text{Aq}$. (Schneider.)

$2\text{Ag}_2\text{S}$, FeS_2 . (Schneider, Pogg. 136. 305.)

Ag_2S , 3FeS , FeS_2 . Min. *Sternbergite*. Decomp. by aqua regia.

Iron sodium sulphide (sodium sulphoferrite), $\text{Na}_2\text{Fe}_2\text{S}_4 + 4\text{H}_2\text{O}$.

Insol. in H_2O . Decomp. by very dil. acids. (Schneider, Pogg. 138. 302.)

Iron sulphophosphide, Fe_2PS_3 .

Attacked by acids at 100° . Decomp. by boiling $\text{NaOH} + \text{Aq}$. (Ferrand, A. ch. 1899, (7) 17. 410.)

Ferrous telluride, FeTe .

Insol. in H_2O ; sol. in acids. (Fabre, C. R. 105. 277.)

Kermes.

See Antimony trisulphide.

“Knallplatin” compounds.

See Fulminoplatinum compounds.

Krypton, Kr.

Absorption by H_2O at t° .

t°	Coefficient of absorption det. by two series of experiments	
0	0.1249	0.1166
10	0.0965	0.0877
20	0.0788	0.0670
30	0.0762	0.0597
40	0.0740	0.0561
50	0.0823	0.0610

(Antropoff, Roy. Soc. Proc. 1910, 83. A. 480.)

Lanthanic acid.

Barium metalanthanate, $\text{Ba}(\text{H}_2\text{La}_2\text{O}_7)$.
(Baskerville, J. Am. Chem. Soc. 79.)

Lithium metalanthanate, $\text{LiH}_2\text{La}_2\text{O}_7$.
(Baskerville.)

Potassium metalanthanate, $\text{KH}_2\text{La}_2\text{O}_7$.
15 H_2O .

Decomp. by H_2O . (Baskerville.)

Sodium metalanthanate, $\text{NaH}_2\text{La}_2\text{O}_7$.
4 H_2O .

Almost insol. in H_2O , but decomp. (Baskerville.)

Disodium tetralanthanate, $\text{Na}_2\text{La}_4\text{O}_{14}$.
Insol. in H_2O . (Baskerville.)

Lanthanicotungstic acid.

Ammonium lanthanicotungstate, $2(\text{La}_2\text{O}_3, 16\text{WO}_3 + 16\text{H}_2\text{O})$.

Ppt. Insol. in H_2O . (E. F. Smit Chem. Soc. 1904, 26. 1481.)

Barium lanthanicotungstate, 5BaO , $16\text{WO}_3 + 16\text{H}_2\text{O}$.

Ppt. (E. F. Smith.)

Silver lanthanicotungstate, $5\text{Ag}_2\text{O}$, $16\text{WO}_3 + 4\text{H}_2\text{O}$.

Very insol. in H_2O . (E. F. Smith)

Lanthanum, La.

Slowly decomp. cold, rapidly hot & attacked by cold conc. H_2SO_4 , but not by cold conc. $\text{HNO}_3 + \text{Aq}$. Sol. in acids. (Hillebrand and Norton, P. 633.)

Lanthanum bromide, $\text{LaBr}_3 + 7\text{H}_2\text{O}$.

Easily sol. in H_2O . Not very soluble in alcohol. Insol. in ether. (C. V. A. H. Bih. 2. No. 7.)

Lanthanum nickel bromide, 2LaBr_3 , $18\text{H}_2\text{O}$.

Deliquescent. (Frerichs and S. 191. 355.)

Lanthanum zinc bromide, 2LaBr_3 , $36\text{H}_2\text{O}$.

Very deliquescent. (F. and S.)

Lanthanum carbide, LaC_2 .

Decomp. by H_2O and dil. acids. (Baskerville, B. 1895, 28. 2422.)

Sol. in conc. H_2SO_4 and dil. acids, but not in conc. HNO_3 .

fused oxidizing agents; decomp. by ordinary temps. (Moissan, C. R. 3. 149.)

um chloride, LaCl_3 .

rous. Deliquescent. (Hermann.)
in acetone. (Naumann, B. 1904, 37.)

H_2O . Not deliquescent. (Zschiesche.)
sol. in alcohol. (Hermann.)

um mercuric chloride, $2\text{LaCl}_3, \text{HgCl}_2, \text{H}_2\text{O}$.

deliquescent. Very sol. in H_2O .
ac, Ann. Min. (5) 15. 272.)

um stannic chloride.

lorostannate, lanthanum.

um fluoride, $\text{LaF}_3 + \text{H}_2\text{O}$.

itate. Sl. sol. in $\text{HCl} + \text{Aq}$. (Cleve.)

um hydrogen fluoride, $2\text{LaF}_3, 3\text{HF}$.

itate. (Frerichs and Smith, A. 191.

not exist. (Cleve, B. 11. 910.)

um hydride, La_2H_3 .

ap. by dil. acids. (Winkler, B. 24.

Decomp. by H_2O . Sol. in acids
olution of H_2 . Decomp. by alkalis.
ann, A. 1902, 325. 266.)

um hydroxide, $\text{La}_2\text{O}_3\text{H}_4$.

in H_2O ; easily sol. in acids; insol. in
 $\text{NaOH} + \text{Aq}$.

citric acid. (Baskerville, J. Am.
oc. 1904, 26. 49.)

um zinc iodide, $2\text{LaI}_3, 3\text{ZnI}_2 + 27\text{H}_2\text{O}$.

ol. in H_2O . (Frerichs and Smith, A.
)

um nitride, LaN .

ap. by H_2O with evolution of NH_3 .
mineral acids. Decomp. by alkali.
ann, A. 1902, 325. 275.)

um oxide, La_2O_3 .

sol., even when ignited, in mineral,
ic acids. (Hermann.)

boiling conc. $\text{NH}_4\text{Cl} + \text{Aq}$. (Mos-

cold conc. $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Damour
ille.)

in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Mosander.)
in acetone. (Naumann, B. 1904, 37.

um peroxide, La_2O_6 .

$\text{HCl}, \text{H}_2\text{SO}_4, \text{HNO}_3$, and $\text{HC}_2\text{H}_3\text{O}_2 +$
decomp. (Cleve, Bull. Soc. (2) 43.

$+x\text{H}_2\text{O}$. Unstable. Sol. in dil.
 Aq with decomp. (Melikoff, Z.
109, 21. 71.)

Lanthanum oxybromide, LaOBr .

Ppt. (Frerichs and Smith.)

Lanthanum oxychloride, $3\text{La}_2\text{O}_3, 2\text{LaCl}_3$.

Insol. in H_2O . Difficultly and slowly sol.
in HCl , or $\text{HNO}_3 + \text{Aq}$. (Hermann.)

LaOCl . Boiling H_2O dissolves only traces.
(Frerichs and Smith.)

Lanthanum sulphide, La_2S_3 .

Decomp. by H_2O and acids. (Didier.)

Lanthanum disulphide, LaS_2 .

Decomp. by heat. (Biltz, Z. anorg. 1911,
71. 435.)

Lead, Pb.

Lead, in contact with H_2O and air free from CO_2 ,
gives a solution of PbO which turns litmus blue and
turmeric red, and is turned brown with H_2S .

H_2O which has been boiled does not dissolve Pb if
there is no access of air. When shaken up with air it
dissolves 0.01 to 0.008% PbO in 2 hours. Pure spring
water, containing $1\frac{3}{4}$ grains salts in 2 pounds H_2O and
no CO_2 , when conducted through a lead pipe 150 feet
long, dissolves so much lead that it turns brown with
 H_2S . (Yorke, Phil. Mag. J. 5. 82.)

CO_2 or small amts. of salts prevent the solution of
Pb. 1 vol. H_2O with $\frac{3}{4}$ vol. CO_2 dissolves only a trace
of Pb. Spring H_2O , containing in 10 pounds 1.21
grains NaCl and CaCl_2 , and 6.4 grains CaCO_3 dissolved
in CO_2 , does not dissolve lead. (Yorke.)

If the amt. of salts in solution equals $\frac{1}{1000}$ the amt. of
 H_2O , and especially if they are carbonates, very slight
amts. of Pb are dissolved. (Christison, Phil. Mag. J.
21. 158.)

CaCO_3 dissolved in CO_2 water decreases the solu-
bility of Pb more than any other salt.

Distilled H_2O , quietly standing in a closed flask with
lead and air free from CO_2 , deposits white flocks of
 PbO_2H_2 , and dissolves $\frac{1}{1000}$ pt. PbO . The solution has
an alkaline reaction. (v. Bonsdorff, Pogg, 41. 305.)

Water of 3° hardness does not take up enough Pb to
become injurious. (Clarke, J. B. 1856. 608.)

Soluble carbonates increase the solubility of Pb in
 H_2O (Nevins, C. C. 1851. 608); especially $(\text{NH}_4)_2\text{CO}_3$.
(Böttger.)

Presence of H_2SO_4 decreases the solubility of Pb.
(Horsford, Chem. Gaz. 1849. 247.)

H_2O containing K_2SO_4 takes up only a trace of Pb.
(Wetzlar, Schw. J. 54. 324.)

Presence of sulphates diminishes (Christison), does
not diminish (Graham, Miller, and Hoffmann), the
action of H_2O on Pb.

CaSO_4 protects Pb, but it is attacked by much
 MgSO_4 . (Nevins.)

$\text{NaCl} + \text{Aq}$ dissolves only a trace of Pb.
 $\frac{1}{1000}$ pt. of a chloride in H_2O is not sufficient to pre-
vent the solubility of Pb in H_2O . (Christison.)

Presence of chlorides increases the solubility. (Gra-
ham, Miller, and Hoffmann; Nevins.)

H_2O containing KNO_3 does not corrode Pb.
Nitrates hinder the action of H_2O . (v. Bonsdorff.)

Nitrates increase the action of H_2O . (Graham, Miller,
and Hoffman.) Nitrates have no influence. (Kersting.)

10 lbs. of H_2O dissolved the following amts.
from Pb pipes in 24 hours: if distilled $\text{H}_2\text{O} +$
1% Na_2CO_3 , 0.38 grain Pb; if Duna water,
0.19 grain Pb; if canal water, 0.15 grain Pb; if
distilled $\text{H}_2\text{O} + 1\%$ NH_4NO_3 , 0.15 grain Pb;
if hard well water, 0.04 grain Pb; if distilled
 $\text{H}_2\text{O} + 1\%$ KNO_3 , 0.01 grain Pb. (Kersting,
Dingl. 169. 183.)

200 l. Manchester drinking water dissolved
2.094 g. from 1 sq. metre Pb in 8 weeks; 9 l.
well water dissolved 1.477 g. from 1 sq. metre
Pb in 8 weeks; 11 l. distilled H_2O containing

are dissolved 110.003 g. from 1 sq. metre Pb in 8 weeks; distilled H_2O free from air dissolved 1.829 g. from 1 sq. metre Pb in 8 weeks; sea water dissolved 0.038 g. from 1 sq. metre Pb in 8 weeks. (Calvert and Johnson, C. N. 16. 171.)

A lead pipe taken up in Paris, which had been exposed to action of ordinary H_2O for 200 years, was found perfectly smooth and uncorroded. (Belgrand, C. R. 77. 1055.)

Pb is attacked by all waters, hard or soft; even highly calcareous water dissolves some lead. (Mayençon and Bergeret, C. R. 78. 484.)

Pure distilled H_2O does not act on Pb, but extremely small quantities of NH_3 , HNO_3 , etc. cause an action; but for this action on Pb the presence of air and CO_2 is also required. (Stallman, Dingl. 180. 366.)

100 ccm. distilled H_2O dissolved 3 mg. from 11.8 sq. cm. lead in one week when air without CO_2 was passed through the solution. 8 mg. were dissolved when the air contained CO_2 . (Wagner, Dingl. 221. 260.)

Action of dil. salt solutions on lead. In 500 ccm. of the solutions containing salt, bright sheets of lead of 5600 sq. metres' surface were so suspended that the liquid reached all parts of the metal without hindrance, and the amts. dissolved determined after 24, 48, and 72 hours of action.

Salt	Grammes salt per litre	Dissolved Pb in mg per litre		
		after 24	48	72 hrs
NH_4NO_3	0.020	13.0	...	25
"	0.040	15.0	...	32
"	0.080	15.0
$\{ KNO_3 +$	$\{ 0.020$
$\{ NaNO_3 +$	$\{ 0.050$	2.0	2.0	...
$\{ KNO_3 +$	$\{ 0.040$
$\{ Na_2SO_4 +$	$\{ 0.212$	0.8	1.0	...
$\{ KNO_3 +$	$\{ 0.045$
$\{ K_2CO_3 +$	$\{ 0.308$	0.3
$\{ KNO_3 +$	$\{ 0.070$	0.5
$\{ K_2SO_4 +$	$\{ 0.504$	0.8
$\{ CaSO_4 +$	$\{ 0.252$	0.4
"	0.408	0.4	1.0	...
K_2CO_3	0.310	0.2
"	0.516	0.2
$CuCl_2$	0.250	0.5	0.5	0.5
"	0.510	0.3	...	0.4
Na_2SO_4	0.200	0.8
"	0.400	0.5
$\{ NH_4NO_3 +$	$\{ 0.020$
$\{ CuCl_2 +$	$\{ 0.060$	1.8
$\{ NH_4NO_3 +$	$\{ 0.020$
$\{ K_2CO_3 +$	$\{ 0.100$	0.4
$\{ Na_2SO_4 +$	$\{ 0.200$
$\{ Na_2SO_4 +$	$\{ 0.200$
$\{ K_2CO_3 +$	$\{ 0.040$	0.1
$\{ CaCl_2 +$	$\{ 0.100$
Water from L. Katrine		1.0	1.0	1.5
Distilled water		2.0	2.0	3.0

(Muir, C. N. 25. 294.)

Action of salt solutions on 11.8 sq. cm. Pb in one week while air either with or without CO_2 was passed through the solution.

Solubility of Pb in salt solutions.

100 ccm. solutions containing the given salts dissolve Pb in mg.:—

Salt	g. salt in 100 ccm.	mg. Pb dissolved	
		without CO_2	with CO_2
KCl	0.5	21	12
NaCl	0.5	21	12
NH_4Cl	1.0	12	5
$MgCl_2$	0.83	20	35
K_2SO_4	1.0	0	0
KNO_3	1.0	14	20
Na_2CO_3	1.0	0	...
NaOH	0.923	430	...
$CaO.H_2O$	Saturated	137	...

(Wagner, Dingl. 221. 260.)

Solubility of Pb in salt solutions.

25 sq. cm. were acted upon by a solution containing 0.2 g. salt in a litre for 21 days.

Three series of experiments were carried on. I. In corked flasks. II. In beakers covered with porous paper; diameter of mouth of beaker = 11.5 cm. III. In basins covered with porous paper; diameter of mouth of basin = 14.5 cm. IV. In corked flasks with constant current of air. V. In beakers half filled and covered with porous paper, the lead being suspended so that equal amts. of surface were above and beneath the liquid.

The amts. in mgs. of Pb dissolved were as follows:—

Salt used	I	II.	III.	IV.	V
NH_4NO_3	1.8	4.0	16.0
KNO_3	1.6	0.5	6.0	1.5	...
$CaCl_2$	3.0	2.8	5.5	2.5	3.5
$(NH_4)_2SO_4$	0.7	1.3	16.0	5.0	2.5
K_2CO_3	0.3	0.3	0.7	0.6	0.3
Dist. H_2O	1.5	0.8	4.2	2.0	...

(Muir, Chem. Soc. 36. 600.)

H_2O sat. with CO_2 dissolves 0.012 g. Pb in a litre in 3 days. (Marais, C. R. 77. 1529.)

Action of H_2O charged with CO_2 under 700 mm. pressure on Pb. 3 mg. of Pb were dissolved per litre in 24 hours, and the amt. was not increased by further action. The addition of 100 mg. K_2CO_3 + 20 mg. NH_4NO_3 to a litre prevented all action.

Action of H_2O charged with CO_2 under 6 atmos. pressure on Pb.

14.8 mg. were dissolved per l. in 24 hours, and 24 mg. per l. in 48 hours.

Action of various salt solutions added to above solution of CO_2 were as follows:—

	mg. salt per l.	mg. Pb dissolved	
		after 24 hrs.	after 48 hrs.
. . . .	80	13.2	32.0
. . . .	160	6.0
. . . .	160	32.0	44.0
O ₂ . . .	16	5.0
O ₂ . . .	40	10.0	35.0
and H ₂ O	14.8	24.0

(Muir, C. N. 33. 125.)

corrosion of Pb by ordinary distilled depends upon the presence of CO₂ and O. dissolved CO₂ is double the amt. of the ed O, the action is most energetic. CO₂ is wholly absent and O present, ion is very slight, and when the H₂O is 1½ or more vol. % CO₂ with normal f oxygen, there is no visible corrosion. distilled H₂O containing neither O nor is no action on Pb. In the above cases eater part of the Pb remains in the f a white ppt. or crust on the Pb, but case where O and CO₂ are both present ratio of 1 : 2, very small amts. of Pb solution in a few days; the amt., how- iminishes on standing. As the amt. of creases, the amt. of Pb dissolved in the so increases.

OH alone does not protect Pb from on, but when in combination with CO₂, ion is much diminished.

H₂, and NaOH + Aq attack Pb much ctively in absence of CO₂ and presence

In absence of dissolved O neither l₂ nor NaOH attacks Pb.

CO₂ + Aq in absence of CO₂ attacks Pb y, but NaHCO₃ + Aq has not the slight- ion.

l₂(CO₃)₂ + Aq also has not the slightest on Pb, and the presence of CaCO₃ and holly prevents H₂O attacking Pb.

O₄ + Aq in presence of air forms a crust but no Pb is found in solution, but if eluded there is no visible action. Pres- f CO₂ causes a strong corrosive action.

containing CaSO₄ and CaH₂(CO₃)₂, ot attack Pb.

above reactions are not in the least l by the presence of moderate amts. of m, chlorides, or ammonium, or organic unds; but ammonium salts in excess strong solvent action on Pb. (Muller, (2) 36. 317.)

also an extended report of the action of n Pb made to the Water Committee of rfield, England, in 1886, by Messrs. es, Odling, and Tidy.

y extended researches are published by lley and Frew (Jour. Soc. Chem. Ind. 7. which only the general conclusions can on here.

The action of slaked lime, limestone, sand calcium silicate, mortar, etc., was tested. The results were as follows:—

1. In nearly all cases the corrosion is greater with free exposure to the air than when air is excluded. The difference is especially great in those cases where the greatest action on the lead takes place. Aluminum hydroxide and blue clay form exceptions, and exert a greater corrosive action when air is excluded. In the case of CaCO₃, old mortar, CaSiO₃, or a mixture of CaCO₃ and CaO₂H₂, the exclusion or presence of air makes no appreciable difference.

KNO₃ + Aq shows a peculiar behaviour. In the presence of air it acts nearly as much on the Pb as pure H₂O, but when air is excluded it exerts nearly as much retarding action as CaSiO₃.

2. In the presence of air the action of H₂O on Pb is considerably increased by the presence of NH₄NO₃ or CaO₂H₄; with exclusion of air, by CaSO₄, also by a mixture of CaO₂H₂ and sand. All the other investigated substances, even KNO₃, hinder the action of H₂O on Pb either with or without exclusion of air.

3. CaO₂H₂ + Aq exerts in all cases a much greater corrosive action than pure H₂O, and although this action is diminished by sand yet fresh mortar very quickly destroys lead pipes when in contact therewith. Old mortar, on the other hand, and also CaSiO₃ and CaCO₃, have a protective action.

4. The fact is very important that sand, CaCO₃, old mortar, CaSiO₃, and a mixture of sand and CaCO₃ afford considerable protection to lead against H₂O. A mixture of limestone and sandstone has more effect than the two substances separately.

5. CaSiO₃ totally prevents the corrosive action of KNO₃ and NH₄NO₃, so that the lead is not attacked by solutions of those salts any more than by H₂O containing CaSiO₃ alone. Sand, and a mixture of sand and CaCO₃ have a similar effect, but not to such a degree.

6. The protective influence of CaCO₃ does not appear to depend on the presence of CO₂ and the formation of CaH₂(CO₃)₂.

7. MgCO₃ prevents the corrosion of Pb as much as CaSiO₃. (Carnelley and Frew, Jour. Soc. Chem. Ind. 7. 15.)

Pb in contact with Zn or Fe is protected thereby from the solvent action of H₂O, and in fact the action is nearly null. Sn, on the other hand, increases the action. This is of importance in regard to the use of tin-coated lead pipes.

The presence of Ca salts does not influence the action of the H₂O on Pb, hard or soft H₂O provided it contains CO₂ having a strong corrosive action. Removal of air from H₂O diminishes the solvent action. Simple filtration will remove all Pb from H₂O if suitable filters are used. (Flögel, J. B. 1888. 2645.)

Pure distilled H_2O has strong corrosive action on Pb, which is very much weakened by addition of a solution of $CaCO_3$ in carbonic acid water, but the presence of sulphates increase the action. Pb is not appreciably attacked by H_2O in presence of chlorides alone,

but very strongly when $CaSO_4$ is also present. H_2O containing CO_2 also corrodes Pb. conclusion was drawn that the absence of action of H_2O on Pb in lead pipes is due to presence of traces of $CaH_2(CO_3)_2$. (Barth and Gucci, C. C. 1888. 934.)

Solubility in H_2O containing various solids in solution.

Water used	Pts. of lead per 100,000		
	1	2	3
Water alone, unfiltered	8.19	12.98	8.19
Water alone, filtered	3.00	4.09	2.07
Water containing 0.049 g. NaCl per l., unfiltered	1.36	2.73	0.68
“ “ “ “ “ “ “ “ filtered	0.68	1.50	0.67
Water containing 0.49 g. Na_2SO_4 per l., unfiltered	3.41	6.83	2.05
“ “ “ “ “ “ “ “ filtered	2.05	3.41	1.64
$CaHCO_3$ +Aq containing 0.04 g. CaO as carbonate per l.	2.45	3.14	2.63
$CaHCO_3$ +Aq with NaCl	2.05	3.41	2.35
$CaHCO_3$ +Aq with Na_2SO_4	2.18	3.32	2.05
$CaSO_4$ +Aq containing 0.095 g. CaO as sulphate per l.	6.83	6.83	3.41
$CaSO_4$ +Aq with NaCl	5.46	6.57	3.51
$CaSO_4$ +Aq with Na_2SO_4	4.78	5.87	3.69

Column 1 gives the numbers for distilled water free from air; column 2 for distilled aerated by agitation with air; column 3 for water continuously aerated by passing 1 of air through it per hour; column 4 for distilled water through which 1 litre of air and cc. of CO_2 were passed per hour throughout the experiment. (Antony and Benelli, ch. it. 1896, 26, (2) 97 and 352.)

Almost insol. in cold HCl +Aq, and only sl. attacked when boiling. Completely sol. in HNO_3 +Aq if not too conc., but presence of H_2SO_4 or HCl diminishes the solvent power to a great extent. (Rose.)

Granulated Pb is sl. sol. in conc. HCl +Aq; addition of $PtCl_4$ makes the action very energetic. Dil. HCl +Aq may also be used with $PtCl_4$. (Millon, C. R. 21. 49.)

HCl +Aq of 1.2 sp. gr., with Pb, gives off H at ord. temp., more abundantly when heated. Evolution of H is hastened by placing Cu in contact with the Pb. (Stolba, J. pr. 94. 113.)

Quickly decomp. by hot HCl +Aq, slowly by cold. (Sharples, C. N. 50. 126.)

Scarcely acted upon by boiling conc. HCl +Aq.

Sol. in aqua regia.

HNO_3 +Aq is the best solvent, but Pb is as good as insol. in a mixture of HNO_3 and H_2SO_4 . (Berzelius.)

Not acted upon by very conc. HNO_3 +Aq.

Pb is only sl. attacked by HNO_3 +Aq of any strength below 15° . Above 15° it is most rapidly attacked by a rather weak acid. (Montemartini, Gazz. ch. it. 22. 397.)

Action of H_2SO_4 on Pb.

H_2SO_4 of 1.842 sp. gr. dissolves 201 g. from 1 sq. metre pure lead at ordinary temp. (time?), and H_2SO_4 of 1.705 sp. gr. dissolves only 59 g.

Slight impurities in the lead lessen this

solubility. (Calvert and Johnson, Chem (2) 1. 66.)

Strongly attacked by 99.8% H_2SO_4 at temp. with exclusion of air. (Lunge, 1 261. 131.)

When 0.2 g. pure Pb was heated with ccm. H_2SO_4 of $66^\circ B$. there was no appreciable action below 175° . At $230-250^\circ$ all the Pb was suddenly converted into $PbSO_4$, and dissolved. (Bauer, B. 8. 210.)

Lead is slowly attacked by pure cold H_2SO_4 +Aq (99.78% H_2SO_4). Lead vessels which held the H_2SO_4 were gradually destroyed by long standing. (Napier and Lock, C. N. 42. 314.)

H_2SO_4 +Aq (20%) does not evolve H under the same circumstances. (Stolba.)

Sol. in $HC_2H_3O_2$ +Aq when in contact with the air.

Strong NH_4OH +Aq does not dissolve litharge; but lead immersed in NH_4OH 3 days gives an ammonia solution containing 0.0139% lead. (Endemann, Am. Chem. 1897, 19. 892.)

Somewhat sol. in $NaCl$ +Aq. (Reich, Dingl. 172. 155.)

$NaCl$ +Aq attacks Pb at high temp. (Lunge, l. c.)

Action of $KClO_3$. $KClO_3$ +Aq (10% $KClO_3$) oxidised 64.31 g. Pb from 1 sq. metre surface by boiling 7 hours; $KClO_3$ +Aq (20% $KClO_3$) oxidised 151.12 g. under same conditions; and $Ca(ClO_3)_2$, $CaCl_2$ +Aq ($20^\circ B$)

; and $\text{Ca}(\text{ClO}_3)_2$, $\text{CaCl}_2 + \text{Aq}$ (20° ne), obtained by passing Cl_2 through $\text{H}_2 + \text{Aq}$, oxidised 437.70 g. (Lunge and Jeler, Jour. Soc. Chem. Ind. 4. 31.)
sol. in liquid NH_3 . (Gore, Am. Ch. J. 20. 828.)

l. in a solution of K in liquid NH_3 . us, J. Am. Chem. Soc. 1907, 29. 1562.)

ccm. oleic acid dissolves 0.0592 g. Pb in lys. (Gates, J. phys. Chem. 1911, 15.)

solubility of Pb in petroleum.

b.-pt. is under 230° , only slightest trace dissolved in 4 months; if $230-300^\circ$, 0.0026% months; if over 300° , 0.0244% in 4 months; if over 300° , 0.0244% in 4 ths.

solubility of Pb in commercial oil of turpentine and resin oil.

	Temp.	% Pb dissolved	
		in 8 days	in 14 days
h oil of rprntine .	$15-20^\circ$	sl. trace	0.0722
oil of tur- ntine . .	$15-20$	0.0522	0.1435
h oil of rprntine .	100	0.265	0.715
oil of tur- ntine . .	100	0.982	1.851
h oil of rprntine .	$130-150$	0.938	2.045
oil of tur- ntine . .	$130-150$	1.738	4.083
h resin oil	$15-20$	trace	0.024
"	$15-20$	0.073	0.185
h "	100	0.380	0.880
"	100	1.190	2.711
h "	$130-150$	1.050	2.065
"	$130-150$	2.208	4.740

(Engler and Kneis, Dingl. 263. 193.)

b is strongly attacked by oil of turpentine. l. Chem. 4. 289.)

he fatty oils dissolve Pb in considerable . (Macadam, J. B. 1878. 1169.)

ot attacked by sugar + Aq. (Klein and 5, C. R. 102. 1176.)

d potassium amide.

ae Potassium ammonoplumbite.

d azoimide, basic, PbO , PbN_3 .

sol. in H_2O . (Wöhler, B. 1913, 46. 2054.)

d azoimide, PbN_3 .

sol. in cold H_2O ; much less sol. in boiling than PbCl_2 . 1 l. H_2O dissolves about $\frac{1}{2}$

PbN_3 . Easily sol. in warm $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.
sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Curtius, B. 24.

l.)

Lead bromide, PbBr_2 .

Sl. sol. in cold, more easily in hot H_2O , or in H_2O containing HCl , HNO_3 , or $\text{HC}_2\text{H}_3\text{O}_2$ (Löwig.)

1 l. H_2O dissolves 6 g. PbBr_2 at 10° ; addition of HBr causes a ppt. which redissolves on further addition of HBr . 1000 pts. of a liquid containing 720 pts. HBr dissolve 550 g. PbBr_2 . This solubility increases by heating. (Ditte, C. R. 92. 718.)

1 l. H_2O dissolves 26.28 millimols. PbBr_2 at 25.2° . (von Ende, Z. anorg. 1901, 26. 159.)

Solubility in 100 g. H_2O at t° .

t°	G. PbBr_2
0	0.4554
15	0.7305
25	0.9744
35	1.3220
45	1.7457
55	2.1376
65	2.5736
80	3.3430
95	4.3613
* 100	4.7510

* By extrapolation. (Lichty, J. Am. Chem. Soc. 1903, 25. 474.)

Sl. sol. in H_2O .

8.34×10^{-1} gram. are dissolved in 1 liter of sat. solution at 20° . (Böttger, Z. phys. ch. 1903, 46. 603.)

Solubility of PbBr_2 in $\text{HNO}_3 + \text{Aq}$ at 25.2° .
S = solubility in millimols per litre.

HNO_3 normal	S
0.001	39.11
0.01	39.87
0.051	42.56
0.04 $\text{KNO}_3 +$ 0.01 HNO_3	42.77

(von Ende, Z. anorg. 1901, 26. 162.)

Slowly sol. in cold, easily in warm NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Wittstein.)

Not pptd. in presence of Na citrate. (Spiller.)

Insol. in H_2O containing $\text{Pb}(\text{NO}_3)_2$. (von Ende, Z. anorg. 1901, 26. 159.)

Insol. in benzene. (Franchimont, B. 16. 387.)

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate (Naumann, B. 1910, 43. 314.)

Difficultly sol. in acetone. (Naumann, B. 1904, 37. 4328.)

+ $3\text{H}_2\text{O}$. (Ditte, l. c.)

Lead hydrogen bromide, $5\text{PbBr}_2, 2\text{HBr} + 10\text{H}_2\text{O}$.

Sol. in $\text{HBr} + \text{Aq}$. (Ditte, C. R. 92. 718.)

Lead magnesium bromide, $\text{PbBr}_2, 2\text{MgBr}_2 + 16\text{H}_2\text{O}$.

Very deliquescent. Decomp. immediately by H_2O or alcohol. (Otto and Drewes, Arch. Pharm. 229. 585.)

Lead potassium bromide (potassium bromoplumbite, $\text{PbBr}_2, \text{KBr} + \text{H}_2\text{O}$.

(Remsen and Herty, Am. Ch. J. 14. 124.)
+ H_2O . (Wells, Sill. Am. J. 145. 129.)

$\text{PbBr}_2, 2\text{KBr}$. Sol. in a little H_2O without decomp., but decomp. by an excess with separation of PbBr_2 . (Lowig.)
+ H_2O . (Wells, Sill. Am. J. 145. 129.)
 $2\text{PbBr}_2, \text{KBr}$. (Wells.)

Lead potassium perbromide, $\text{K}_2\text{Pb}_2\text{Br}_8 + 4\text{H}_2\text{O}$.

Decomp. by H_2O and alcohol. (Wells, Z. anorg. 4. 340.)

Lead rubidium bromide, $\text{PbBr}_2, 2\text{RbBr} + \frac{1}{2}\text{H}_2\text{O}$.

(Wells, Sill. Am. J. 146. 34.)
 $2\text{PbBr}_2, \text{RbBr}$. (Wells.)

Lead sodium bromide.

Decomp. by H_2O . (Lowig.)

Lead bromochloride, $\text{PbBrCl} = \text{PbBr}_2, \text{PbCl}_2$.

Can be recrystallised from H_2O without decomp. (Iles, C. N. 43. 216.)
 $3\text{PbCl}_2, \text{PbBr}_2$. Sol. in H_2O with decomp. Sol. in HCl and in HBr . Insol. in cold alcohol; sl. sol. in boiling alcohol. (Thomas, C. R. 1899, 128. 1235.)

Lead bromiodide, $\text{PbBrI} = \text{PbBr}_2, \text{PbI}_2$.

Decomp. by H_2O . Cryst. from a solution of PbI_2 in HBr . (Grissom and Thorp, Am. Ch. J. 10. 229.)
 $3\text{PbBr}_2, \text{PbI}_2$. Decomp. by H_2O . (Thomas C. R. 1899, 128. 1236.)
 $6\text{PbBr}_2, \text{PbI}_2$. (G. and T.)

Lead bromosulphide, $\text{PbBr}_2, \text{PbS}$.

Properties as chlorosulphide. (Parmentier.)

Lead chloride, PbCl_2 .

Slowly sol. in 135 pts. H_2O at 12.5° , and in a much smaller quantity of hot H_2O . (Bischof.)
Sol. in 30 pts. cold, and 22 pts. hot H_2O . (Wittstein.)
Sol. in 30 pts. H_2O at 18.75° . (Abl.)
100 pts. H_2O dissolve 4.59 pts. PbCl_2 at 15.5° . (Ure's Dict.)
100 pts. H_2O dissolve 0.9712 pt. PbCl_2 at 20° . (Formánek, C. C. 1887. 270.)
100 pts. H_2O dissolve 0.946 pt. PbCl_2 at 17.7° . (Bell, Chem. Soc. (2) 6. 355.)
Sol. in 105.2 pts. H_2O at 16.5° . (Bell, C. N. 16. 69.)

100 pts. H_2O dissolve 0.8 pt. PbCl_2 at 1.18 pts. at 20° ; 1.7 pts. at 40° ; 2.1 pt. at 55° ; 3.1 pts. at 80° . (Ditte, C. R. 92. 718.)
1 l. H_2O dissolves 38.80 millimols. PbCl_2 at 25.2° . (von Ende, Z. anorg. 1901, 26. 14.)
 9.61×10^{-1} gram are dissolved in 1 lit sat. solution at 20° . (Böttger, Z. phys. 1903, 46. 603.)

Solubility in H_2O .

100 g. H_2O dissolve g. PbCl_2 at t° .

t°	G. PbCl_2
0	0.6728
15	0.9090
25	1.0842
35	1.3244
45	1.5673
55	1.8263
65	2.1265
80	2.6224
95	3.1654
* 100	3.3420

* By extrapolation.

(Lichty, J. Am. Chem. Soc. 1903, 25. 4)

33.6 millimols. Pb are dissolved in 1 l. H_2O at 18° . (Pleissner, C. C. 1907, II. 1.)
1 l. H_2O dissolves 77.76 milliequivalents of PbCl_2 at 25° . Sp. gr. of the solution 25° 1.0069. (Harkins and Winninghoff, J. Chem. Soc. 1911, 33. 1816.)

0.0388 mol. mg. PbCl_2 are sol. in 1 l. (Kernot and Pomilio, Soc. R. Napoli, (3), XVII, 353.)

A colloidal modification is sol. in hot H_2O to give cryst. modification. (Van de V. Ch. Z. 1893, 17. 1908.)

Solubility in H_2O is not much increased by the addition of acids. (Fresenius.)

Sol. in conc. $\text{HCl} + \text{Aq}$, from which pptd. by H_2O , but less sol. in dil. HCl than in H_2O . (Berzelius.)

Sol. in 1636 pts. H_2O containing 1 pt. PbCl_2 . (Bischof.)

Sat. solution of PbCl_2 in $\text{HCl} + \text{Aq}$ of sp. gr. contains 2.566% PbCl_2 at 16.5° .

Solubility in $\text{HCl} + \text{Aq}$. 100 pts. liquid containing pts. HCl of 1.1162 sp. gr. i. e. 100 pts. H_2O dissolve pts. PbCl_2 at 17.7° .

Pts. HCl	Pts. PbCl_2	Pts. HCl	Pts. PbCl_2	Pts. HCl	Pts. PbCl_2
1	0.347	8	0.099	50	0.000
2	0.201	9	0.096	60	0.000
3	0.165	10	0.093	70	0.000
4	0.145	15	0.090	80	0.000
5	0.131	20	0.111	90	0.000
6	0.107	30	0.151	100	0.000
7	0.100	40	0.216

(Bell, Chem. Soc. 21. 350.)

Solubility of PbCl_2 in HCl .

°C.	Amount PbCl_2 dissolved in 1000 pts. of liquid				
	At 0°	At 20°	At 40°	At 55°	At 80°
	8.0	11.8	17.0	21.0	31.0
	2.8	3.0	4.6	6.5	12.4
	1.2	1.4	3.2	5.5	12.0
	2.4	4.8	7.2	9.8	19.8
	4.7	6.2	10.4	12.9	23.8
	11.9	14.1	19.0	24.0	38.0
	29.8	30.0

(Ditte, C. R. 92, 718.)

Solubility in $\text{HCl} + \text{Aq}$ at 0°. $\frac{\text{PbCl}_2}{2} = \frac{1}{2}$ mols.Solubility in mgs. in 10 cc. solution; HCl = mols. HCl in ditto.

$\frac{1}{2}$	HCl	$\frac{\text{PbCl}_2}{2}$	HCl
2	0.	0.072	5.8
2	0.35	0.088	11.7
35	0.675	0.100	29.5
1	1.125	0.209	46.7
35	1.6	0.95	73.5
99	2.3	1.5	89.0
90	3.4	1.9	96.0
8	4.5	3.01	111.5

It is seen that very little $\text{HCl} + \text{Aq}$ is sufficient to diminish solubility very considerably, but on further addition of $\text{HCl} + \text{Aq}$, the solubility is nearly constant, and increases very much when large amounts of $\text{HCl} + \text{Aq}$ are present. (Engel, A. ch. (6) 17, 359.)

Solubility of PbCl_2 in $\text{HCl} + \text{Aq}$ at 25°

Cl	G. PbCl_2 per l.	G. HCl per l.	G. PbCl_2 per l.
	10.79	3	5.0
5	9.0	6	3.1
	7.6	10	1.8
	6.0		

(Noyes, Z. phys. Ch. 1892, 9, 623.)

Solubility of PbCl_2 in $\text{HCl} + \text{Aq}$ at 25.20°. Solubility in millimols per litre.

Normal	S	HCl normal	S
100	38.80	0.3714	6.35
109	38.66	0.5142	5.37
122	38.20	0.7386	4.73
130	37.94	1.026	4.41
145	37.35	1.538	4.61
161	35.80	2.051	5.18
174	34.99	2.564	6.25
181	33.75	3.085	7.78
186	31.46	3.718	8.16
192	29.33	5.0	19.38
192	25.46	7.5	65.86
210	17.12	10.0	141.35
250	10.12	12.05	164.3

(von Ende, Z. anorg. 1901, 26, 148.)

Solubility of PbCl_2 in HCl at 18°.

HCl Normality	G. PbCl_2 per l.
0	9.34
0.0001	9.305
0.0002	9.300
0.0005	9.243
0.00102	9.200
0.0102	8.504

(Pleissner, Arb. Kais. Gesundheitsamt. 1907, 26, 384.)

Sol. in hot, insol. in cold conc. H_2SO_4 . (Hayes.)Sol. in dil. $\text{HNO}_3 + \text{Aq}$, from which it is pptd. by $\text{HCl} + \text{Aq}$. (Gladstone.)Easily and completely decomp. by hot $\text{HNO}_3 + \text{Aq}$. (Wurts.)Solubility of PbCl_2 in $\text{HNO}_3 + \text{Aq}$ at 25.2°.

S = solubility in millimols per litre.

HNO_3 normal	S
0.001	38.87
0.01	39.71
0.051	42.92
0.04 $\text{KNO}_3 +$	
0.01 HNO_3	43.36

(von Ende, Z. anorg. 1901, 26, 162.)

Solubility of PbCl_2 in $\text{NH}_4\text{Cl} + \text{Aq}$ at 25.20°.

S = solubility in millimols per litre.

NH_4Cl normal	S
0.25	9.47
0.50	7.11
1.0	4.35

(von Ende, Z. anorg. 1901, 26, 152.)

Solubility of $\text{PbCl}_2 + \text{NH}_4\text{Cl}$ at 22°.

G. equiv per l. H_2O NH_4Cl	G. equiv. per 100 cc. H_2O PbCl_2	G. equiv per l. H_2O NH_4Cl	G. equiv per 100 cc. H_2O PbCl_2
0.0	7.49×10^{-4}	1.0	0.758×10^{-4}
0.1	3.10	1.2	0.707
0.2	1.916	1.5	0.671
0.3	1.508	2.0	0.695
0.4	1.348	2.5	0.812
0.5	1.263	3.0	0.968
0.55	1.189	4.0	1.502
0.6	1.092	5.0	2.338
0.65	1.012	6.0	3.580
0.7	0.956	7.0	5.628
0.8	0.837	7.29*	6.46
0.9	0.793		

* Saturated.

(Brönsted, Cong. Appl. Chem. 1909, Sec. X, 110.)

Solubility in $\text{NH}_4\text{Cl} + \text{Aq}$ at t° .			
t°	G. PbCl_2 in 100 g. of the solution	G. NH_4Cl in 100 g. of the solution	Solid phase
17°	0.89	0.0	PbCl_2
	0.21	0.96	
	0.16	1.43	
	0.14	2.40	
	0.076	3.48	
	0.078	4.23	eutectic-pt.
	0.078	4.93	
	0.098	12.36	$2\text{PbCl}_2, \text{NH}_4\text{Cl}$
	0.34	22.33	
	0.64	26.49	
	0.52	26.68	eutectic-pt.
	0.33	26.91	
	0.30	27.03	NH_4Cl
	0.0	27.14	
50°	1.69	0.0	PbCl_2
	1.08	0.51	
	0.67	1.45	
	0.58	2.45	
	0.48	4.86	
	0.49	12.45	eutectic pt.
	0.71	19.42	
	1.76	27.16	$2\text{PbCl}_2, \text{NH}_4\text{Cl}$
	3.31	31.90	
	3.96	33.56	
	2.65	33.62	eutectic pt.
	1.62	33.88	
	0.32	34.14	NH_4Cl
	0.0	34.25	
100°	3.10	0.0	PbCl_2
	2.02	1.32	
	1.85	5.33	
	1.80	6.01	
	1.76	8.59	
	1.98	13.19	eutectic pt.
	4.54	26.08	
	8.32	32.64	$2\text{PbCl}_2, \text{NH}_4\text{Cl}$
	11.40	36.29	
	12.67	37.62	
	12.50	38.14	eutectic pt.
	11.60	38.32	
	10.70	38.66	$\text{PbCl}_2, 2\text{NH}_4\text{Cl}$
	9.88	40.22	
	9.26	41.90	
	4.21	42.91	eutectic pt.
	3.06	43.20	
	1.61	43.42	NH_4Cl
	0.0	43.51	

These results show that the double salt $\text{PbCl}_2, 2\text{NH}_4\text{Cl}$ can only exist in aqueous solution at temperature above 70° .
(Demassieux, C. R. 1913, 156. 894.)

Much more sol. in $\text{HgCl}_2 + \text{Aq}$ than in

Grammes HgCl_2 in 100 ccm.	Grammes PbCl_2 dissolved	After subtracting amt. dissolved by H_2O alone	Calcn. no. gramms 100 g
0	0.9712
4	1.8972	0.9350	23
2	1.4874	0.5208	26
1	1.2272	0.2600	26
0.5	1.0808	0.1134	22
0.25	1.0192	0.0500	20
0.125	0.9926	0.0226	18

(Formánek, C. C. 1887. 270.)

Solubility of PbCl_2 in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$

G. equiv. per l.	
$\text{Pb}(\text{NO}_3)_2$	PbCl_2
0.0	0.0777
0.2	0.0832

(Noyes, Z. phys. Ch. 1892, 9. 623.)

Solubility in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at 25°
C = concentration of $\text{Pb}(\text{NO}_3)_2$ in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ expressed in milliequivalents per l.
 d_1 = Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at 25° .
S = Solubility of PbCl_2 in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ expressed in milliequivalents per l.
 d_2 = Sp. gr. $25^\circ/4^\circ$ of $\text{PbCl}_2 + \text{Pb}(\text{NO}_3)_2 + \text{Aq}$.

C	d_1	S	d_2
20.020	1.0008	76.75	1.0
50.063	1.0045	76.64	1.0
99.660	1.0119	77.98	1.0

(Harkins and Winninghof, J. Am. Chem. 1911, 33. 1816.)

Solubility of PbCl_2 in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$

G. $\text{Pb}(\text{NO}_3)_2$ per l.	% PbCl_2
0	1.09
3.31	1.10
6.62	1.06
33.12	1.11
82.80	1.29

(Armstrong and Eyre, Proc. Roy. Soc. (A) 88. 234.)

Solubility of PbCl_2 in $\text{KCl} + \text{Aq}$ at 25° .
S = Solubility in millimols per litre.

KCl normal	S	KCl normal	
0.0000	38.80	0.0999	16
0.001	38.32	0.5006	7
0.0025	37.85	0.7018	7
0.0049	37.02	0.9991	4
0.0049	37.02	0.9991	4
0.0099	35.28	1.5018	4
0.0200	32.16	2.0024	5
0.0599	22.62	3.0036	9

(von Ende, Z. anorg. 1901, 26. 151.)

ty of PbCl₂+KCl in H₂O at 20°. Values = g. equivalents.

In 1000 g. H ₂ O	Solid phase	
	PbCl ₂	KCl
28.0 4.57	PbCl ₂ , KCl. 1/2H ₂ O + KCl	
18 23.42 4.18	PbCl ₂ , KCl. 1/2H ₂ O	
15 21.50 3.96		
11 19.85 3.73		
7 18.66 3.50		
6 17.48 3.33		
17 16.17 3.03		
15 16.06 3.01		
10 15.80 2.93		
36 14.92 2.87		
35 15.63 2.86		
30 15.03 2.78	2PbCl ₂ , KCl	
29 14.30 2.77		
24 13.70 2.70		
20 12.72 2.64		
29 14.35 2.62		
14 12.47 2.55		
10 12.13 2.49		
04 11.60 2.41		
65 10.96 2.31		
84 10.42 2.20		
75 8.92 1.79	PbCl ₂	
26 8.72 1.744		
04 8.56 1.570		
24 8.29 1.472		
24 8.11 1.348		
23 8.13 1.347		
27 7.98 1.231		
22 8.01 1.225		
59 8.00 1.152		
22 8.10 1.107		
88 8.13 1.068	PbCl ₂	
30 8.28 1.000		
80 8.38 0.943		
21 8.79 0.875		
83 8.96 0.833		
19 9.36 0.761		
39 10.03 0.672		
75 11.18 0.602		
23 12.85 0.545		
03 12.88 0.523		
83 12.85 0.502	PbCl ₂	
75 13.04 0.497		
58 12.95 0.475		
75 13.65 0.387		
99 14.88 0.306	PbCl ₂	
95 19.33 0.199		

A study of the equilibrium between lead chloride and sodium chloride in aqueous solution at 13°, 50° and 100° shows that at none of these temp. do these chlorides form a double salt. (Demassieux, C. R. 1914, 158. 702.)

Solubility in salts + Aq at 25°.

Salt used	Concentration of the salt. Equivalents per liter	Solubility of PbCl ₂ Equivalents per liter
None	0	0.07770
HCl	0.05	0.04786
"	0.1	0.03243
"	0.2	0.01927
KCl	0.05	0.0482
"	0.1	0.0341
"	0.2	0.0219
MgCl ₂	0.05	0.0503
"	0.1	0.0350
CaCl ₂	0.05	0.0503
"	0.1	0.0355
"	0.2	0.0219
MnCl ₂	0.05	0.0501
"	0.1	0.0349
"	0.2	0.0217
ZnCl ₂	0.2	0.0220
CdCl ₂	0.05	0.0601
"	0.1	0.0481
"	0.2	0.0355

(Noyes, Z. phys. Ch. 1892, 9. 623.)

Sol. in KOH + Aq. (Rose.)
Less sol. in dil. salt solutions than in H₂O, especially CaCl₂ + Aq; sol. in 534 pts. H₂O containing CaCl₂. (Bischof.)
More sol. in Na₂S₂O₃ + Aq than in H₂O, but not as sol. as AgCl. (Herschell, 1819.)
More sol. in NaC₂H₃O₂ + Aq than in H₂O. (Anthon.)
Easily sol. in NH₄NO₃ + Aq.
Sl. sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 828.)
Insol. in conc. alcohol. (Wittstein.) Insol. in 94% alcohol; very sl. sol. in cold or hot 76% alcohol.
Solubility in alcohol at 25°.
Alcohol = g. mol. alcohol in 1 l. of solvent.
PbCl₂ = g. mol. PbCl₂ in 1 l. of solution.
Alcohol 4 2 1 1/2 1/4
PbCl₂ 0.0172 0.0257 0.0298 0.0330 0.0338
Alcohol 1/4 0
PbCl₂ 0.0367 0.0388
(Kernot and Pomilio, Soc. R. Napoli, (3) 17. 353.)

ted, Z. phys. Ch. 1912, 80. 208.)

s sol. in 120 pts. pure H₂O, but on % NaCl 437 pts. are required to tion. When PbCl₂ is digested with 1 + Aq, 1 pt. dissolves in 129 pts. of

Insol. in benzene. (Franchimont, B. 16. 387.)
Insol. in CS₂. (Arctowski, Z. anorg. 1894, 6. 257.)
Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)
Insol. in methyl acetate (Naumann, B.

1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in methylal. (Eidmann, C. C. 1899, II; 1014.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Glycerine dissolves 1.995% PbCl_2 .

1 pt. glycerine + 1 pt. H_2O dissolves 1.32% PbCl_2 .

1 pt. glycerine + 3 pts. H_2O dissolves 1.0365% PbCl_2 .

Glycerine containing 87.5% H_2O dissolves 0.91% PbCl_2 . (Piesse, B. 7. 599.)

Solubility of PbCl_2 in mannite + Aq at 25°.

Mannite = g. mol. mannite in 1 l. of solvent.

PbCl_2 = g. mol. PbCl_2 in 1 l. of solution.

Mannite	$\frac{1}{2}$	$\frac{1}{4}$	$\frac{1}{8}$	$\frac{1}{16}$	$\frac{1}{32}$
PbCl_2	0.0408	0.0403	0.0394	0.0384	0.0385

Mannite	$\frac{1}{64}$	0
PbCl_2	0.0377	0.0388

(Kernot and Pomilio, Soc. R. Napoli, (3) 17. 353.)

Min. *Cotunnite*.

Lead tetrachloride, PbCl_4 .

Sol. in H_2O with subsequent decomp. (Rivot, Beudant, and Daguin, Ann. Min. (5) 4. 239.)

Obtained in a pure state by Friedrich. Sol. in a little cold H_2O , but is decomp. by warming or diluting. Miscible with conc. HCl + Aq; not attacked by conc. H_2SO_4 even on warming. (Friedrich, W. A. B. 102, 2b. 534.)

Lead tetrachloride with MCl .

See Chloroplumbate, M.

Lead magnesium chloride, $\text{PbCl}_2, 2\text{MgCl}_2 + 13\text{H}_2\text{O}$.

Deliquescent. Decomp. by H_2O . (Otto and Drewes, Arch. Pharm. 228. 495.)

Lead potassium chloride (potassium chloroplumbite), $\text{PbCl}_2, \text{KCl}$.

(Remsen and Herty, Am. Ch. J. 14. 125.)

Contains $\frac{1}{2}$ H_2O . (Wells, Sill. Am. J. 145. 130.)

See also Demassieux, $\text{PbCl}_2 + \text{KCl}$ under PbCl_2 .

$2\text{PbCl}_2, \text{KCl}$. (Wells.)

See also Demassieux as above.

Lead rhodium chloride.

See Chlororhodite, lead.

Lead rubidium chloride, $\text{PbCl}_2, 2\text{RbCl} + \frac{1}{2}\text{H}_2\text{O}$.

(Wells, Sill. Am. J. 146. 34.)

$2\text{PbCl}_2, \text{RbCl}$. (Wells.)

Lead sodium chloride.

Decomp. by H_2O .

Lead sodium tetrachloride, $2\text{PbCl}_4, 9$

Very sol. in H_2O . (Sobrero and ch. (3) 29. 165.)

See also Chloroplumbate, lead.

Lead thalious chloride, $\text{PbCl}_2, 3\text{TlCl}$

Sl. sol. in cold, more in hot H_2O .

Z. phys. Ch. 9. 622.)

$\text{PbCl}_2, \text{TlCl}$. Ppt. (Ephraim, 2 1909, 61. 245.)

Lead chloride ammonia, $2\text{PbCl}_2, 3\text{NH}_3$

(Rose, Pogg. 20. 157.)

Lead tetrachloride ammonia, $\text{PbCl}_4, 4$

Pptd. from chloroform solution. thews, J. Am. Chem. Soc. 1898, 20. 8

$\text{PbCl}_4, 2\text{NH}_3$. Fumes in the air. by H_2O . (Matthews.)

Lead chloride arsenate, $3\text{Pb}_2(\text{AsO}_4)_3$

See Arsenate chloride, lead.

Lead chloride borate, $\text{Pb}(\text{BO}_2)_2, \text{H}_2\text{O}$.

See Borate chloride, lead.

Lead chloride carbonate.

See Carbonate chloride, lead.

Lead chloride chlorite.

See Chlorite chloride, lead.

Lead chloride with fluoride and iodide.

See Lead chlorofluoride and Lead iodide.

Lead chloride phosphate.

See Phosphate chloride, lead.

Lead chloride phosphite, $\text{PbCl}_2, \text{Pb}_2$

Ppt. (Berzelius.)

Does not exist. (Rose.)

Lead chloride sulphate.

See Sulphate chloride, lead.

Lead chloride sulphide, $\text{PbCl}_2, 3\text{PbS}$

See Lead chlorosulphide.

Lead chlorofluoride, PbClF .

Sl. sol. in H_2O without decomp. sol. in HNO_3 + Aq. (Berzelius.)

Solubility in H_2O .

100 g. H_2O dissolve 0.0211 g. F 0°; 0.0370 g. at 25°; 0.1081 g. at 100°. Z. anorg. Ch. 1911, 70. 174.)

Solubility in HCl + Aq at 25°.

Solution of PbClF in HCl + Aq at 0.0535 g. equiv. per l. contains (PbClF in 100 cc. of solvent.

Solution of PbClF in HCl + Aq at

solubility in $\text{PbCl}_2 + \text{Aq.}$

G. equiv. per l. PbCl_2	G. PbClF in 100 cc. of solvent
0.0100	0.0020
0.0195	0.0016
0.0495	0.0002
0.00996	0.0030
0.0196	0.0008
0.0392	0.0005

Solubility in $\text{KOH} + \text{Aq}$, according to Ditte (C. R. **94**. 130). When $\text{KOH} + \text{Aq}$ is gradu-

ally added to lead hydroxide suspended in H₂O, the lead hydroxide is at first dissolved proportional to the amount of KOH, until the strength reaches 200 g. KOH to 1 litre H₂O. The solubility then diminishes and increases again until 400 g. KOH are dissolved in 1 litre H₂O. The amorphous lead hydroxide is then converted into crystalline 2PbO(PbO₂H₂). By further addition of KOH the solubility is suddenly decreased, and then increases again. (Ditte.)

Sol. in triethyl toluenyl ammonium hydrate + Aq.

Sol. in sorbine + Aq. (Pelouze.)

Sol. in acetates + Aq. (Mercer.)

Sol. in Ca, Ba, Sr, K, or Na succrate + Aq.

Not pptd. in presence of Na citrate + Aq. (Spiller.)

See also under Lead, and Lead oxide.

Lead perhydroxide, PbO₂, H₂O.

See Lead peroxide.

Lead imide, PbNH.

Decomp. by H₂O and dilute acids. (Franklin, Z. anorg. 1905, 46. 27.)

Lead iodide, PbI₂.

Sol. in 187 pts. boiling H₂O. (Berthemot.)

Sol. in 1235 pts. H₂O at ord. temp., and 194 pts. at 100°. (Denot, J. pr. 1. 425.)

Sol. in 2400 pts. H₂O at 18.75°. (Abl.)

Sat. PbI₂ + Aq at 20° contains 0.0017 pt.; at 27°, 0.002 pt.; at 100°, 0.0039 pt. PbI₂. (Lassaigne, J. chim. med. 7. 364.)

1 l. H₂O dissolves 0.6 g. PbI₂ at 10°. (Ditte, C. R. 92. 718.)

1 l. H₂O dissolves 1.58 millimols PbI₂ at 25.2°. (Von Ende, Z. anorg. 1901, 26. 159.)

0.47 × 10⁻¹ gram are dissolved in 1 litre of sat. solution at 20°. (Böttger, Z. phys. Ch. 1903, 46. 603.)

Solubility in 100 g. H₂O at t°.

t°	G. PbI ₂ .
0	0.0442
15	0.0613
25	0.0764
35	0.1042
45	0.1453
55	0.1755
65	0.2183
80	0.3023
95	0.3960
* 100	0.4360

* By extrapolation.

(Lichty, J. Am. Chem. Soc. 1903, 25. 474.)

0.0013 g. mol. PbI₂ are dissolved H₂O at 20°. (Fedotieff, Z. anorg. 19178.)

Not more sol. in HC₂H₃O₂ + Aq t H₂O, contrary to Henry. (Denot, l.)

Pptd. from aqueous solution by littl Aq, but redissolved by the addition of (Ditte, C. R. 92. 718.)

Insol. in cold, sol. in hot HCl + A decomp.

Solubility of PbI₂ in HNO₃ + Aq at 2 S = Solubility in millimols. per litre.

HNO ₃ normal	S
0.001	38.87
0.01	39.06
0.051	39.45
0.04 KNO ₃ +	
0.01 HNO ₃	39.45

(von Ende, Z. anorg. 1901, 26. 16)

Sol. in KOH + Aq.

Sol. in conc. KI, NaI, BaI₂, SrI₂, Ca MgI₂ + Aq, from which it is pptd. b (Berthemot.)

Very sol. in KI + Aq, 2 mola. PbI₂ dissolved for 1 mol. KI. (Boullay.)

Sol. in NH₄I + Aq. Easily sol. in Na₂ Aq. (Werner, C. N. 53. 51.)

Not pptd. in presence of Na citrate. ler.)

Solubility in sat. I₂ + Aq at 20° = 0.0 mol. per l. Solid phase PbI₂ + I₂. (Fe Z. anorg. 1911, 73. 178.)

Very easily sol. in liquid NH₃. (Fr Am. Ch. J. 1898, 20. 828.)

Sl. sol. in alcohol. (Henry.) Deco boiling ether. (Vogel.)

100 g. formic acid dissolve 0.25 g. at (Aschan, Ch. Ztg. 1913, 37. 1117.)

Insol. in CS₂. (Arctowski, Z. anorg 6. 257.)

Sl. sol. in benzonitrile. (Naumai 1914, 47. 1369.)

Difficultly sol. in methyl acetate. mann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumai 1910, 43. 314.)

Insol. in acetone. (Naumann, B. 37. 4329.)

0.02 pts. are sol. in 100 pts. acetone ; 0.02 pts. are sol. in 100 pts. amyl alcoh

133.5. 0.50 pts. are sol. in 100 pts. aniline s

1.10 pts. are sol. in 100 pts. aniline at (Laszczynski, B. 1894, 27. 2287.)

ubility of PbI_2 in pyridine at t° .

G. PbI_2 per 100 g. pyridine	Solid phase
0.166	$\text{PbI}_2, \text{C}_5\text{H}_5\text{N}$
0.175	"
0.186	"
0.200	"
0.215	"
0.225	$\text{PbI}_2, \text{C}_5\text{H}_5\text{N} + \text{PbI}_2, 2\text{C}_5\text{H}_5\text{N}$
0.208	$\text{PbI}_2, 2\text{C}_5\text{H}_5\text{N}$
0.188	"
0.190	"
0.228	"
0.290	"
0.340	"
0.370	"
0.410	"
0.445	"

ise, J. phys. Ch. 1912, **16**, 273.)

Iodogen iodide, $\text{PbH}_2\text{I}_4 = \text{PbI}_2, 2\text{HI}$.

H_2O dissolves out HI . Sol. in hot m which crystallizes PbI_2 . (Guyot, med. **12**. 247.)

H_2O . Decomp. by H_2O . (Berthelot, **1024**.)

Lithium iodide, $\text{PbI}_2, \text{LiI} + 5\text{H}_2\text{O}$.

1 mol. H_2O at 95° and loses another H_2O at 100° . (Bogorodski, C. C. **1894**,

$2\text{LiI} + 6\text{H}_2\text{O}$. Sl. sol. in H_2O . (Mosnier, R. **1895**, **120**. 446.)

Magnesium iodide, $\text{PbI}_2, 2\text{MgI}_2$.

Decomp. by H_2O and by alcohol. (Mosnier, ch. **1897**, (7) **12**. 402.)

H_2O . Very hygroscopic. Decomp. completely by H_2O . (Otto and Drewes, Pharm. **229**. 180.)

Nickel iodide, $\text{PbNi}_2\text{I}_6 + 3\text{H}_2\text{O}$.

Decomp. by H_2O . (Mosnier, A. ch. **1897**, **11**.)

Potassium iodide (Potassium iodoplumb-), PbI_2, KI .

Stable. Completely decomp. by H_2O . Sol. upon by cold, but completely decomp. by hot alcohol. (Boullay, A. ch. **366**.)

H_2O . The only salt that could be prepared by Remsen and Herty (Am. Ch. J. **1897**, **19**. 11.)

2KI . Sl. sol. in boiling chloroform; insol. in strong $\text{KI} + \text{Aq}$, insol. in alcohol. (C. N. **1898**, **77**. 191.)

H_2O . Decomp. by H_2O . (Berthelot, **5** **29**. 289.)

Does not exist. (R. and H.)

H_2O . (Ditte, C. R. **92**. 134.) Does not exist. (R. and H.)

$\text{PbI}_2, 4\text{KI}$. Decomp. by H_2O ; insol. in alcohol. (Boullay.) Does not exist. (R. and H.)

$3\text{PbI}_2, 4\text{KI} + 6\text{H}_2\text{O}$. (Berthelot, l. c.) Does not exist. (R. and H.)

Lead potassium periodide, $\text{K}_2\text{Pb}_2\text{I}_8 + 4\text{H}_2\text{O}$.

Decomp. by H_2O or alcohol. (Wells, Z. anorg. **4**. 346.)

Lead rubidium iodide, $\text{PbI}_2, \text{RbI} + 2\text{H}_2\text{O}$.

(Wells, Sill. Am. J. **146**. 34.)

Lead silver iodide, $\text{PbI}_2, 2\text{AgI}$.

(Ruff and Geisel, B. **1905**, **38**. 2663.)

Lead silver iodide ammonia, $\text{PbI}_2, 2\text{AgI}, 5\text{NH}_3$.

(Ruff and Geisel, B. **1905**, **38**. 2663.)

Lead sodium iodide, PbI_2, NaI .

Decomp. by H_2O . (Poggiale, C. R. **20**. 1180.)

$+x\text{H}_2\text{O}$. (Remsen and Herty, Am. Ch. J. **14**. 124.)

$\text{PbI}_2, 2\text{NaI} + 6\text{H}_2\text{O}$. Sl. sol. in H_2O . (Moisnier, C. R. **1895**, **120**. 445.)

Lead iodide ammonia, $\text{PbI}_2, 2\text{NH}_3$.

Decomp. by H_2O . (Rammelsberg, Pogg. **48**. 166.)

Lead iodide carbonate.

See Carbonate iodide, lead.

Lead iodosulphide, $\text{PbS}, 4\text{PbI}_2$.

Decomp. by light, heat, acids and alkalies. (Lenher, J. Am. Chem. Soc. **1895**, **17**. 512.)

Sol. in conc. HI ; insol. in dil. $\text{HI} + \text{Aq}$. (Lenher, J. Am. Chem. Soc. **1901**, **23**. 681.)

$\text{IPbS}, \text{PbS}, \text{PbI}_2$. Ppt. (Hofmann, B. **1904**, **37**. 251.)

Lead suboxide, Pb_2O .

Decomp. by H_2O into PbO, H_2 .

Decomp. by dil. $\text{H}_2\text{SO}_4, \text{HCl}, \text{HNO}_3, \text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, or alkalies, into PbO , which dissolves, and Pb , which dissolves or not, according to the reagent. Sol. in dil. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$.

Lead monoxide (Litharge), PbO .

Sol. in 7000 pts. H_2O . (Horsford.)

Pure PbO is insol. in H_2O . (Brandecke, Repert. **53**. 155; Siebold, Repert, **53**. 174; Herbergen, Repert. **55**. 55.) Sl. sol. in H_2O . (Yorke, Phil. Mag. (3) **5**. 82.)

0.31 millimoles Pb are dissolved in 1 liter H_2O at 18° . (Pleissner, C. C. **1907**, II. 1056.)

1.71×10^{-2} g. are dissolved in 1 litre of sat. solution at 20° . (Böttger, Z. phys. Ch. **1903**, **46**. 603.)

Easily sol. in acids.

Sol. in KOH, or NaOH + Aq; also in $\text{CaO}_2\text{H}_2 + \text{Aq}$.
Sol. in boiling $\text{Cu}(\text{NO}_3)_2 + \text{Aq}$ with pptn. of Aq.
Sol. in CaCl_2 , and $\text{SrCl}_2 + \text{Aq}$. (André, C. R. 104. 359.)
Sol. in $\text{MgCl}_2 + \text{Aq}$. (Voigt, Ch. Ztg. 13. 695.)
Sol. in boiling $\text{Cu}(\text{NO}_3)_2 + \text{Aq}$ with pptn. of CuO .
Partially sol. in $\text{Cd}(\text{NO}_3)_2$, and $\text{Mn}(\text{NO}_3)_2 + \text{Aq}$ with pptn. of CdO and MnO respectively.
Not acted upon by Mg, Ag, Co, Ni, or Ce nitrates + Aq. (Persoz.)
Very sol. in $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$. (Rochleder.)
Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)
Insol. in acetone. (Eidmann, C. C. 1899, II, 1014.)
When finely pulverised, sol. in cane sugar + Aq, but less than Pb_3O_4 . (Peschier.)
Sl. sol. in glycerine. Readily sol. in glucose + Aq. (Persoz.)
Sol. in volatile oils. (Schweitzer.)
Yellow modification.

Solubility in H_2O at 22° .

	Solubility in g.-equiv. per litre
1. Yellow PbO , obtained by boiling lead hydroxide with 10% NaOH	1.03×10^{-4}
2. Yellow PbO , obtained by heating 1 at 630°	1.05×10^{-4}
3. Yellow PbO , obtained by heating at 740° red PbO , formed by boiling lead hydroxide with conc. NaOH	1.00×10^{-4}
4. Yellow PbO obtained by heating pure, commercial, yellow-brown PbO at 620° .	1.09×10^{-4}

(Ruer, Z. anorg. 1906, 50. 273.)

Red modification. Obtained by boiling lead hydroxide with conc. NaOH + Aq.
Solubility in H_2O at $22^\circ = 0.56 \times 10^{-4}$ g. equiv. per l. (Ruer, Z. anorg. 1906, 50. 273.)
Yellow-brown modification. Solubility in H_2O at $22^\circ = 1.10 \times 10^{-4}$ g. equiv. per litre. (Ruer, Z. anorg. 1906, 50. 273.)
See also **Lead**.
Min. *Massicot*.

Lead oxide (Red lead), Pb_3O_4 .

Insol. in H_2O .
Converted by acids into PbO_2 and salts of monoxide.
Sol. in a large amt. of glacial acetic acid. (Berzelius.) Insol. in acetic acid. (Schonbein, J. pr. 74. 325.)

Solution in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ may de or not according to concentration
When treated with an excess of HC Aq of 8°B , Pb_3O_4 is quickly dissol the solution soon deposits PbO_2 ; composition is facilitated by dilutio if Pb_3O_4 is treated with a large e glacial $\text{HC}_2\text{H}_3\text{O}_2$, it dissolves, and t tion is permanent if atmospheric ai cluded, and temp. does not rise ab (Jacquelain, J. pr. 53. 152.)
Insol in acetone. (Eidmann, C. (II, 1014; Naumann, B. 1904, 37. 43
Easily sol. in cane sugar + Aq. (Pe Min. *Minium*.

Lead sesquioxide, Pb_2O_3 .

Insol. in H_2O or in KOH + Aq.
Decomp. by strong acids into Pb corresponding salt of monoxide.

Lead peroxide, PbO_2 .

Insol. in H_2O . Sol. in acids, also i alkali hydroxides + Aq. The soluti acids are very unstable, except wha centrated and kept at a low temperatu
Decomp. by cold HCl, HCN, HI HI + Aq. Not attacked by other aci cold, but decomp. thereby when hot. in moderately conc. HNO_3 , H_2S $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

There are two forms of PbO_2 , the phous and the crystalline.
1 l. of very conc. H_2SO_4 dissolves 1 mols. crystalline PbO_2 .

Solubility of amorphous PbO_2 in H_2S at 22° .

99.5 millimols. PbO_2 are dissolved i acid containing 1720 g. H_2SO_4 .
4 millimols. PbO_2 are dissolved in acid containing 1097 g. H_2SO_4 .
 v = moles H_2SO_4 per mole of H_2O .
 c = millimols. PbO_2 dissolved in 1 li

v	c	v	
0.32	0.82	0.20	
0.30	0.4	0.15	
0.25	$7 \cdot 10^{-2}$	0.10	

(Dolezalek and Finckli, Z. anorg. 1 323-5.)

Decomp. by $\text{NH}_4\text{OH} + \text{Aq}$. Sol. i KOH, or NaOH + Aq.
Sol. with decomp. in $\text{Hg}_2(\text{NO}_3)_2$ (Levol.)
Insol. in acetone. (Eidmann, C. (II, 1014; Naumann, B. 1904, 37. 294
Min. *Plattnerite*.

Lead manganese peroxide, PbO_2 , 4B

Ppt. (Gibbs and Parkmann, Sill. (2) 39. 58.)

romide, PbBr_2 , PbO .

H_2O .
and $3\text{H}_2\text{O}$. (André, C. R. 96.

$\text{bBr}_2 + 2\text{H}_2\text{O}$. Ppt. (Strömholm, 1904, 38. 436.)

hloride, 2PbCl_2 , $\text{PbO} + 2\text{H}_2\text{O}$.
C. R. 96. 435.)

PbO . Absolutely insol. in hot or
(André, A. ch. (6) 3. 108.)
atlockite.

Sol. in hot $\text{NaOH} + \text{Aq}$. (André.)
lilmols. Pb are dissolved in 1 litre
(Pleissner, C. C. 1907, II. 1055.)

2PbO . Insol. in H_2O . Sol. in dil.
1 (about 110 g. in 1 l.) (Ditte, C. R.

endipite. Easily sol. in $\text{HNO}_3 + \text{Aq}$.
(André, A. ch. (6) 3. 111.)

3PbO . Insol. in H_2O . (Döber-

O. Ppt. (Strömholm, Z. anorg. 435.)

0.10 millimols. Pb are dissolved
 H_2O at 18° . (Pleissner, C. C. 1907,

Ppt. (André, C. R. 104. 359.)

Nearly insol. in H_2O . Sl. sol.
+ Aq . (Vauquelin.)

5PbO . (Döbereiner.)

$\text{PbCl}_2 + 2\text{H}_2\text{O}$. Ppt. (Strömholm, 1904, 38. 434.)

7PbO . *Cassel-yellow*.

ntium oxychloride, 2PbO , $\text{SrCl}_2 +$
C. R. 104. 359.)

hloride, iodide, PbCl_2 , PbI_2 , 4PbO .
Schwarzenbergite. Sol. in dil.
aq.

odide, PbI_2 , PbO .

boiling H_2O or $\text{KI} + \text{Aq}$. (Brandes, 9.)

O. (Ditte, C. R. 92. 145.)

PbO . Insol. in H_2O . (Denot, J. D. 1.)

$\text{PbO} + 2\text{H}_2\text{O}$. Ppt. (Kühn, C. C.)

PbO . Insol. in H_2O . (Denot.)

(Ditte, C. R. 92. 145.)

$\text{PbI}_2 + 2\text{H}_2\text{O}$. Ppt. (Strömholm, 1904, 38. 437.)

$\text{PbI}_2 + 2\text{H}_2\text{O}$. (Strömholm, Z. anorg. 437.)

eriodide, PbO , PbI_2I_2 .

p. by boiling H_2O . Sol. in dil.
+ Aq . (Gröger, W. A. B. 100, 2b.

Lead phosphide, PbP_2 .

Decomp. by H_2O and dil. acids. (Bossuet, C. R. 1913, 157. 721.)

Lead phosphoselenide, PbSe , P_2Se .

Insol. in H_2O or $\text{HCl} + \text{Aq}$. Sol. in $\text{HNO}_3 +$
 Aq .

Insol. in cold, slowly decomp. by hot al-
kalies + Aq . (Hahn, J. pr. (2) 93. 436.)

2PbSe , P_2Se_3 . Insol. in H_2O , HCl , or
 $\text{HNO}_3 + \text{Aq}$. Slowly sol. in red fuming
 HNO_3 . (Hahn.)

2PbSe , P_2Se_3 . Decomp. by fuming HNO_3 .
(Hahn.)

Lead selenide, PbSe .

Cold $\text{HNO}_3 + \text{Aq}$ dissolves Pb with separa-
tion of Se, which dissolves on warming.
(Little, A. 112. 212.)

Min. *Clausthalite*. Sol. in $\text{HNO}_3 + \text{Aq}$
with separation of Se, when warmed.

Lead mercury selenide, $(\text{Pb}, \text{Hg})\text{Se}$.

Min. *Lehrbachite*.

Lead sulphide, PbS .

Very sl. sol. in H_2O .

1 l. H_2O dissolves 3.6×10^{-3} moles. PbS at
 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

1 l. H_2O dissolves 3×10^{-4} g. PbS at 25° .
(Hevesy, Z. anorg. 1913, 82. 328.)

Insol. in dilute acids; alkalies, and alkali
sulphides + Aq . Decomp. with solution in
moderately dil. $\text{HNO}_3 + \text{Aq}$. With conc.
 HNO_3 or aqua regia, PbSO_4 is formed. Sol.
in hot conc. $\text{HCl} + \text{Aq}$.

1 l. H_2O sat. with H_2S dissolves 1.5×10^{-4} g.
 PbS at 25° . (Hevesy, Z. anorg. 1913, 82.
328.)

Insol. in NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett.)
Somewhat sol. in $\text{H}_2\text{S} + \text{Aq}$ when heated
therewith in a sealed tube. (Senarmont, A.
ch. (3) 32. 168.)

Insol. in potassium thiocarbonate + Aq .
(Rosenblatt, Z. anal. 26. 15.)

Sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Waller, J. Anal.
Ch. 5. 646.)

Insol. in liquid NH_3 . (Franklin, Am. Ch.
J. 1898, 20. 828.)

Min. *Galena*, *Galenite*.

Lead polysulphide, PbS_x .

Ppt.; insol. in alkali sulphides; decomp.
by conc. HNO_3 . (Bodroux, C. R. 1900, 130.
1398.)

Lead platinum sulphide.

See Sulphoplatinate, lead.

Lead sulphide mercuric chloride, 3PbS ,
 4HgCl_2 .

Decomp. by H_2O . (Levallois, C. R. 96.
1666.)

Lead sulphobromide, chloride, or iodide.

See Lead bromosulphide, etc.

Lead disulphodiimide, PbN_2S_2 , NH_3 .

Ppt.

Very stable in the air or in a vacuum.

Sol. in no solvent without decomp.

When rapidly heated it explodes very violently at 140° . (Ruff, B. 1904, 37. 1581.)

Lead telluride, PbTe .

Insol. in H_2O . Sol. in cold $\text{HNO}_3 + \text{Aq}$. (Rose, Pogg. 18. 68.)

Min. *Altaite*. Easily sol. in $\text{HNO}_3 + \text{Aq}$.

"Leucone."

Wöhler (A. 127. 268) gives this substance the formula $\text{H}_{10}\text{Si}_3\text{O}_{10}$, but it is identical with silicoformic anhydride, $\text{Si}_3\text{H}_2\text{O}_3$, which see.

Lime.

Quicklime, CaO . See Calcium oxide.

Slaked lime, $\text{CaO} \cdot \text{H}_2\text{O}$. See Calcium hydroxide.

Lithium, Li .

Decomposes H_2O .

Easily sol. in dil. acids. Slowly attacked by conc. H_2SO_4 , rapidly by conc. $\text{HNO}_3 + \text{Aq}$.

Insol. in hydrocarbons. Sol. in liquid NH_3 , but not so easily as K.

Sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 820.)

1 gram atom dissolves:

at	0°	in	3.93	mol.	liquid	NH_3 .
"	-25°	"	3.93	"	"	"
"	-50°	"	3.93	"	"	"
"	-80°	"	3.93	"	"	"

(Ruff, B. 1906, 39. 840.)

Sol. in ethylamine. Insol. in propylamine and in secondary and tertiary amines. (Kraus, J. Am. Chem. Soc. 1907, 29. 1561.)

Lithium amalgam, LiHg_8 .

LiHg_8 is obtained at all temp. up to 100° . Can be cryst. from Hg without decomp. at any temp. below 100° . (Kerp, Z. anorg. 1900, 25. 68.)

Lithium amide, LiNH_2 .

(Ruff, B. 1911, 44. 505.)

Decomp. slowly in the air.

Slowly decomp. by cold, rapidly by hot H_2O .

Slowly decomp. by HCl .

Slowly decomp. by cold, rapidly by hot abs. alcohol. (Titherley, Chem. Soc. 1894, 65. 518.)

Trilithium amide, Li_3NH_2 .

Hydroscopic; decomp. by H_2O . (Dafert, M. 1910, 31. 994.)

Lithium ammonia, Li, NH_3 .

Decomp. by H_2O at ordinary temp. in liquid NH_3 . (Moissan, C. R. 1868, 689.)

$\text{Li}, 3\text{NH}_3$. (Moissan, C. R. 1901, 131. 1000.)

Trilithium ammonium, Li_3NH_4 .

Very hydroscopic, decomp. by H_2O . (Dafert, M. 1910, 31. 992.)

Lithium antimonide, Li_3Sb .

Decomp. by H_2O with evolution of H_2 . Sol. in liquid NH_3 . (Lebeau, C. R. 1885, 285.)

Lithium arsenide, Li_3As .

Decomp. by H_2O ; decomp. violent in fuming HNO_3 . (Lebeau, C. R. 1899, 128. 1000.)

Lithium azoimide, LiN_3 .

Deliquescent. Stable in aq. solution. 36.12 pts. are sol. in 100 pts. H_2O at 16° .
62.07 " " " " 100 " H_2O
66.41 " " " " 100 " H_2O
20.26 " " " " 100 " abs.

Insol. in ether. (Curtius, J. pr. 58. 277.)

+ H_2O . Very hydroscopic; decomposes in air. Very sol. in H_2O and alcohol. Z. anorg. 1898, 17. 18.)

Lithium bromide, LiBr .

Deliquescent.

100 pts. H_2O dissolve at:

0°	34°	59°	82°	103°
143	196	222	244	270 pts.

Sp. gr. of $\text{LiBr} + \text{Aq}$ at 19.5° contains:
0 10 15 20 25 30
1.035 1.072 1.113 1.156 1.204 1.250

35 40 45 50 55% LiBr
1.309 1.368 1.432 1.500 1.580

(Kremers, Pogg. 103. 65; 104. 133; Z. anal. 8. 285.)

Temp. of maximum density of $\text{LiBr} + \text{Aq}$ g.-mol. LiBr in 1000 g.

1.921°	0.294
0.881°	0.438

(de Coppet, C. R. 1900, 131. 1000.)

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Sol. in benzonitrile. (Naumann, 47. 1369.)

Solubility in glycol at $14.7^\circ - 37.1^\circ$. Coninck, Belg. Acad. Bull. 1906, 35. 1000.)

Sol. in acetone. (Naumann, B. 4328; Eidmann, C. C. 1899, II. 1014)

Sol. in methyl acetate (Naumann, 42. 3789); ethyl acetate. (Naumann, 37. 3601.)

tly sol. in ethyl acetate. (Nau-1910, 43. 314.)
100 g. H₂O dissolve 209 g. LiBr (Bogorodski, C. C. 1894, II. 514.)
) (Bogorodski.)
) 100 g. H₂O dissolve 80 g. LiBr and 122 g. at -10°. (Bogorodski.)

nolybdenyl bromide, LiBr, MoOBr₂,
[₂O.
ygroscopic. (Weinland and Knöll, 1905, 44. 111.)

romide ammonia.
H₂. Sol. in H₂O with decomp.
NH₃. " " " " "
NH₃. " " " " "
NH₃. " " " " "
nefoi, C. R. 1900, 130. 1395.)

arbide, LiC₂.
n conc. acids.
used oxidizing agents; decomp. H₂O
d. (Moissan, C. R. 1896, 122. 363.)

ubchloride, Li₂Cl.
p. by H₂O. (Guntz, C. R. 1895,

hloride, LiCl.
deliquescent. Most deliquescent salt
Berzelius. Very sol. in H₂O. Sol.
ts. H₂O at 15°. (Gerlach.)
H₂O dissolve at:
65° 80° 69° 140° 160°
104.2 115 129 139 145 pts. LiCl.
(Gerlach, Z. anal. 8. 281.)

of LiCl + Aq at 15° containing:
5 10 15 20% LiCl,
030 1.058 1.086 1.117
30 35 40% LiCl.
1.182 1.219 1.256
Gerlach, Z. anal. 8. 281.)

of LiCl + Aq at 18° containing:
10 20 30 40% LiCl.
.0563 1.115 1.181 1.255
ohrausch, W. Ann. 1879. 1.)

p. gr. of LiCl + Aq at 25°.

ation of LiCl + Aq.	Sp. gr.
l-normal	1.0243
r "	1.0129
r "	1.0062
r "	1.0030

er, Z. phys. Ch. 1890, 5. 38.)

Sp. gr. of LiCl + Aq.	
g. LiCl in 1000 g. of solution	Sp. gr. 16°/16°
0	1.000000
2.3923	1.001405
6.2360	1.003647
10.1093	1.005921

(Dijken, Z. phys. Ch. 1897, 24. 109.)

Sp. gr. of LiCl + Aq at 0°.

% LiCl	4.26	12.18	22.2	32.5	41.4	43.2
Sp. gr.	1.026	1.073	1.133	1.203	1.267	1.282

(Lemoine, C. R. 1897, 125. 603.)

Sp. gr. of LiCl + Aq at 20°.

Normality of LiCl + Aq	% LiCl	Sp. gr.
10.35	35.97	1.2230
7.17	26.40	1.1550
5.57	21.10	1.1215
2.98	11.83	1.0691
1.06	4.37	1.0232

(Forchheimer, Z. phys. Ch. 1900, 34. 25.)

Sp. gr. 20°/4° of a normal solution of LiCl = 1.022375. (Haigh, J. Am. Chem. Soc. 1912, 34. 1151.)

Sp. gr. of dil. LiCl + Aq at 20.004°.

Conc. = g. equiv. LiCl per l. at 20.004°.

Sp. gr. compared with H₂O at 20.004 = 1.

Conc.	Sp. gr.
0.0000	1.000,000,0
0.0001	1.000,002,5
0.0002	1.000,005,0
0.0005	1.000,012,6
0.0010	1.000,025,3
0.0020	1.000,050,5
0.0050	1.000,125,8
0.0100	1.000,251,0

(Lamb and Lee, J. Am. Chem. Soc. 1913, 35. 1688.)

B.-pt. of LiCl + Aq.

% LiCl	B.-pt.	% LiCl	B.-pt
3.38	101°	16.66	107°
6.54	102	19.35	109
13.04	105	21.8	111

(Skinner, Chem. Soc. 61. 341.)

Sat. LiCl + Aq boils at 171°. (Kremers.)

B.-pt. of LiCl + Aq. P = pts. LiCl to 100 pts. H₂O.

B.-pt.	P	B.-pt.	P	B.-pt.	P
101°	3.5	124°	48.5	147°	87.5
102	7	125	50	148	90
103	10	126	51.5	149	92.5
104	12.5	127	53	150	95
105	15	128	54.5	151	97.5
106	17.5	129	56	152	100
107	20	130	57.5	153	102.5
108	22	131	59	154	105
109	24	132	60.5	155	107.5
110	26	133	62	156	110.5
111	28	134	63.5	157	113.5
112	30	135	65	158	116.5
113	32	136	66.5	158.5	117.96
114	33.5	137	68	159	119.5
115	35	138	69.75	160	122.5
116	36.5	139	71.5	161	125.5
117	38	140	73.25	162	128.5
118	39.5	141	75	163	131.5
119	41	142	77	164	135
120	42.5	143	79	165	138.5
121	44	144	81	166	142.5
122	45.5	145	83	167	146.5
123	47	146	85	168	151

(Gerlach, Z. anal. 26. 437.)

Solubility of LiCl in HCl + Aq at 0°.

Mg. mol. per 10 cc. of solution		G. per 10 cc. of solution		Sp. gr. of solution
LiCl	HCl	LiCl	HCl	
120	0.0	51.0	0.0	1.255
97.5	22.5	41.4	8.2	1.243
67.0	66.0	28.5	24.1	1.249
58.0	81.0	24.6	29.5	1.251

(Engel, A. ch. (6) 13. 385.)

See also LiCl + H₂O.

Sl. sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 828.)

Sol. in absolute alcohol, ether, and alcohol-ether.

B.-pt. of alcoholic solution of LiCl.

% LiCl	B.-pt.	% LiCl	B.-pt.
2.4	78.43° + 0.70°	9.93	78.43° + 5.55°
5.39	" + 2.15	15.94	" + 11.75
8.01	" + 4.18

(Skinner.)

Solubility in alcohol + Aq decreases to 40% alcohol + 60% H₂O. Curve shows minimum at 30% H₂O to 70% alcohol. The sat. pure H₂O and pure alcohol solutions cooled to 0° deposit LiCl, H₂O and LiCl, 4C₂H₅O respectively. (Pina de Rubies, C. A. 1914. 743, 3006.)

Solubility in ethyl alcohol.

Temp. 1.06° 5.07° 13.0° 25.0° 40.6°
% LiCl 14 14 13 14 15
(Lemoine, C. R. 1897, 125. 605.)

100 g. ethyl alcohol dissolve at:

0° 5° 10° 15° 17°
14.42 15.04 16.77 18.79 20.31 g

The solid phase from 0°-17° is LiCl, 4C

20° 30° 40° 50° 60°
24.28 25.10 25.38 24.40 23.46 g.

The solid phase from 20°-60° is
(Turner and Bissett, Chem. Soc. 191
1907.)

Solubility in ethyl alcohol + Aq at

100 g. of the solution contain		
G. C ₂ H ₅ OH	G. H ₂ O	G.
0	55.10	44
5.96	51.52	42
11.07	48.73	40
17.46	43.90	38
18.56	43.70	37
22.16	41.17	36
26.29	39.51	34
28.97	37.42	33
29.27	36.89	33
30.10	36.64	33
30.51	35.67	33
32.79	34.95	32
38.40	31.58	30
49.27	24.67	26
50.32	24.04	25
53.50	20.94	25
58.15	18.47	23
59.78	17.46	22
63.09	14.83	22
70.24	8.66	21
70.70	8.26	21
70.74	7.78	21
79.26	0	20

The solid phase in the mixtures w
richest in alcohol is LiCl; in the ot
tures the solid phase is LiCl + H₂O.

(Pina de Rubies, C. C. 1915, I. 1

Sp. gr. of LiCl in ethyl alcoh
2C₂H₅O).

% salt	Sp. gr. at 14°	Sp. g
0	0.797	0.
5.2	0.839	0.
10.1	0.871	0.
14.6	0.903	0.

(Lemoine, C. R. 1897, 125. 60

Solubility in methyl alcohol at t°.

t°	% LiCl in sat. solution
1	26
23	27
50	30

Lemoine, C. R. 1897, 125. 604.)

Solubility of LiCl in methyl alcohol (LiCl,

	5.2	14.5	22.1
at 21.5°	0.836	0.910	0.974
" 0°	0.854	0.926	0.988

Lemoine, C. R. 1897, 125. 604.)

g. are sol. in 100 g. propyl alcohol.
ap, Z. phys. Ch. 1894, 14. 276.)
n 15 pts. fusel oil. (Gooch, Am. Ch. J.

. methyl alcohol dissolve 42.36 g. LiCl

; ethyl alcohol dissolve 25.83 g. LiCl

. propyl alcohol dissolve 16.22 g. LiCl

. isoamyl alcohol dissolve 9.03 g. LiCl

er and Bissett, Chem. Soc. 1913, 103.
1909.)

Solubility of fused LiCl in alcohols at 25°.

solvent	% LiCl
	45.0
lcohol	2.475
alcohol	3.720
lcohol	9.56
lcohol	8.26
cohol	4.20
ne	4.14
(at 53°C)	1.89

and Mott, J. phys. Chem. 1904, 8.
158.)

ts. pyridine dissolve 7.78 pts. LiCl at
26 pts. LiCl at 100°. (Laszczynski,
, 27. 2288.)

le in anhydrous pyridine, 97% pyri-
aq, 95% pyridine+Aq, and 93%
e+Aq. (Kahlenberg, J. Am. Chem.
18, 30. 1107.)

ultly sol. in methyl acetate (Nau-
B. 1909, 42. 3789); ethyl acetate.
ann, B. 1910, 43. 314.)

n ethyl acetate. (Naumann, B. 1904,
1.)

ility in glycol at 15°=11%. (de
k, Belg. Acad. Bull. 1905, 359.)

. in benzonitrile. (Naumann, B.
7. 1370.)

n ethylamine. (Shinn, J. phys. Chem.
1. 538.)

n acetone. (Eidmann, C. C. 1899, II.

Solubility of LiCl in acetone.

Pts. sol. in 100 pts. acetone	t°
4.60	0
4.40	12
4.11	25
3.76	46
3.12	53
2.14	58

(Laszczynski, B. 1894, 27. 2287.)

+H₂O. 13.536 millimols. are contained
in 1 l. sat. solution at 25°. (Herz, Z. anorg.
1912, 73. 274.)

Solubility in HCl+Aq at 25°.

Millimols HCl in 10 ccm.	Millimols LiCl in 10 ccm.
	135.36
6.30	134.14
10.53	126.52
17.64	122.58

(Herz, Z. anorg. 1912, 73. 274.)

+2H₂O. Sol. in acetone. (Krug and
M'Elroy, J. Anal. Ch. 6. 184.)

+3H₂O. (Bogorodski, C. C. 1894, II,
514.)

Lithium gold chloride.

See Chloraurate, lithium.

Lithium manganous chloride, LiCl, MnCl₂+
3H₂O.

Decomp. by H₂O; stable only in excess of
LiCl. (Chassevant, A. ch. (6) 30. 10.)

Lithium mercuric chloride, 2LiCl, HgCl₂.

Very deliquescent and sol. in H₂O. (Harth,
Z. anorg. 1897, 14. 323.)

Lithium nickel chloride, LiCl, NiCl₂+3H₂O.

Deliquescent. Sol. in H₂O and alcohol.
(Chassevant.)

Lithium thallic chloride, 3LiCl, TiCl₃+
8H₂O.

Very deliquescent. Sol. in H₂O. (Pratt,
Am. J. Sci. 1895, (3) 49. 404.)

Lithium stannic chloride.

See Chlorostannate, lithium.

Lithium uranous chloride, Li₂UCl₄.

As K salt. (Aloy, Bull. Soc. 1899, (3) 21.
264.)

Lithium uranium chloride, UCl₄, 2LiCl.

Very hygroscopic; sol. in H₂O with decomp.
Sol. in acetic acid. Decomp. by alcohol.
(Aloy, Bull. Soc. 1899, (3) 21. 264.)

Lithium zinc chloride, $\text{LiCl}, \text{ZnCl}_2 + 3\text{H}_2\text{O}$.

Extremely deliquescent. (Ephraim, Z. anorg. 1908, 59. 68.)

$3\text{LiCl}, \text{ZnCl}_2 + 10\text{H}_2\text{O}$ (?). Not obtained in solid state. (Ephraim, Z. anorg. 1908, 59. 69.)

Lithium chloride ammonia.

LiCl, NH_3 .

$\text{LiCl}, 2\text{NH}_3$.

$\text{LiCl}, 3\text{NH}_3$.

$\text{LiCl}, 4\text{NH}_3$.

Above salts are all decomp. by H_2O . (Bonnet, C. R. 1898, 127. 367-369.)

Lithium chloriodide, $\text{LiCl}_2\text{I} + 4\text{H}_2\text{O}$.

Deliquescent. (Wells and Wheeler, Sill. Am. J. 144. 42.)

Lithium fluoride, LiF .

Very difficultly sol. in H_2O . (Berzelius, Pogg. 1. 17.)

Two crystalline forms. Only very sl. sol. in H_2O . Very sl. decomp. by H_2O at red heat. (Poulenc, Bull. Soc. 1894, (3) 11. 17.)

Sp. gr. of solution sat. at $18^\circ = 1.003$ and contains 0.27% LiF . (Mylius, B. 1897, 30. 1718.)

Sol. in 800 pts. H_2O , and the presence of NH_4F and NH_3 decreases solubility to 1 : 3500. (Carnot, Bull. Soc. 1889, (3) 1. 250.)

Two crystalline forms are very sl. sol. in HCl ; easily sol. in HNO_3 . (Poulenc, Bull. Soc. 1894, (3) 11. 17.)

Two crystalline forms are insol. in 95% alcohol. (Poulenc.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Lithium hydrogen fluoride, LiHF_2 .

Difficultly sol. in H_2O , but more easily than LiF . (Berzelius.)

Lithium silicon fluoride.

See Fluosilicate, lithium.

Lithium stannic fluoride.

See Fluostannate, lithium.

Lithium tantalum fluoride.

See Fluotantalate, lithium.

Lithium uranyl fluoride, $\text{UO}_2\text{F}_2, 4\text{LiF}$.

(Ditte.)

Lithium zirconium fluoride.

See Fluozirconate, lithium.

Lithium hydride, LiH .

Not deliquescent. Decomp. by H_2 . (Guntz, C. R. 1896, 123. 997.)

Lithium hydrosulphide, LiSH (?).

Deliquescent. Sol. in H_2O and alcohol. (Berzelius, Pogg. 6. 439.)

Lithium hydroxide, LiOH .

Not so deliquescent as NaOH , and apparently not more sol. in hot than cold H_2O . (Gmelin, Gilb. 62. 399.)

Not deliquescent. (Arfvedson, A. ch. 12. 82.)

The solubility of LiOH in H_2O can be expressed by $y = 6.6750 + 0.00346t + 0.00001t^2$, where y = the percentage of Li_2O in a saturated solution. (Dittmar, Jour. Soc. Chem. Ind. 7. 730.)

Solubility of LiOH in H_2O at t° .

t°	G. per 100 g. Solution		G. LiOH per 100 g. H_2O
	Li_2O	LiOH	
0	6.67	10.64	12.7
10	6.74	10.80	12.7
20	6.86	10.99	12.8
25	6.95	11.14	12.9
30	7.05	11.27	12.9
40	7.29	11.68	13.0
50	7.56	12.12	13.3
60	7.96	12.76	13.8
80	8.87	14.21	15.3
100	10.02	16.05	17.5

(Seidell's Solubilities, 1st Ed. 174.)

A sat. aq. solution contains 7.99% LiOH . (Schreinemakers, C. C. 1906, II. 1486.)

Sp. gr. of $\text{LiOH} + \text{Aq}$ at 18° containing:

1.25 2.5 5 7.5% LiOH .
1.0132 1.0276 1.0547 1.0804

(Kohlrausch, W. Ann. 1879. 1.)

Solubility in $\text{Li}_2\text{SbS}_4 + \text{Aq}$ at 30° .

% LiOH	% Li_2SbS_4	Solid Phase
11.4	0	$\text{LiOH}, \text{H}_2\text{O}$
9.1	8.3	"
2.3	29.9	"
2.1	48.3	"
2.1	52.1	$\text{LiOH}, \text{H}_2\text{O} + \text{Li}_2\text{SbS}_4, 10\text{H}_2\text{O}$
1.4	51.8	$\text{Li}_2\text{SbS}_4, 10\text{H}_2\text{O}$
0	51.3	"

(Donk, Chem. Weekbl. 1908, 5. 529, 629, 767.)

Sl. sol. in alcohol; insol. in alcohol-ether. (Mayer.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3801.)

acetone. (Naumann, B. 1904, **37**. Naumann, C. C. **1899**, II. 1014.)
 Iso with H_2O , and $\frac{1}{2}H_2O$. (Göttig, 2.)

ide, Li_2NH .

at high temp. Insol. in toluene, ether and ethyl acetate. Decomp. Ruff, B. 1911, **44**. 506.)

ide, LiI .

scent.

solubility in 100 pts. H_2O at:

0° 59° 75° 80° 99° 120°
 79 200 263 435 476 588 pts. LiI .

of $LiI + Aq$ at 19.5° containing:

15 20 25 30% LiI ,
 79 1.124 1.172 1.224 1.280

45 50 55 60% LiI .
 14 1.489 1.575 1.670 1.777

ra, Pogg. **104**. 133; **111**. 60: Gerlach, Z. anal. **8**. 295.)

of $LiI + Aq$ at 18° containing:

10 15 20 25% LiI .
 .0756 1.1180 1.1643 1.2138

ohlrausch, W. Ann. **1879**. 1.)

maximum g.-mol. LiI dissolved
 $LiI + Aq$ in 1000 g. H_2O .

16° 0.1795

39° 0.4666

Coppet, C. R. 1900, **131**. 178.)

sol. in liquid NH_3 . (Franklin, Am. **18**, **20**. 828.)

ethyl alcohol dissolve 343.4 g. LiI

ethyl alcohol dissolve 250.8 g. LiI

propyl alcohol dissolve 47.52 g. LiI

isoamyl alcohol dissolve 112.50 g.

in case of propyl alcohol the solid
 25° is LiI , $4C_2H_5O$. (Turner and
 Chem. Soc. 1913, **103**. 1909.)

solubility in organic solvents at t°.

by wt. of LiI in 100 ccm. of the
 on.

of liters which at the saturation
 d in solution 1 mol. LiI .

at	t°	C	L
thane	25°	45.86	0.292
	25°	2.519	5.32
	0°	1.219	10.98

len, Z. phys. Ch. 1906, **55**. 718.)

ity in glycol at 15.3° = 28%. (de
 Belg. Acad. Bull. **1905**, 359.)

Sol. in benzonitrile. (Naumann, B. 1914, **47**. 1369.)

Sol. in methyl acetate. (Naumann, B. 1909, **42**. 3789.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, **43**. 314.)

Sol. in acetone. (Eidmann, C. C. **1899**, II. 1014; Naumann, B. 1904, **37**. 4328.)

+ H_2O . Mpt. below 200°. (Bogorodsky, C. C. **1897**, II. 175.)

+2 H_2O . Mpt. 86–88°. (Bogorodsky.)

+3 H_2O . Mpt. 75°. (Bogorodsky.)

Sol. in absolute alcohol without decomp. (Thirsoff, Chem. Soc. 1894, **66**. (2) 234.)

The composition of the hydrates formed by LiI at different dilutions is calculated from determinations of the lowering of the fr.-pt. produced by LiI and of the conductivity and sp. gr. of $LiI + Aq$. (Jones, Am. Ch. J. 1905, **34**. 301.)

Lithium mercuric iodide, $2LiI, HgI_2 + 6H_2O$.

Very deliquescent. Decomp. by H_2O . Very sol. without decomp. in alcohols, glycerine, acetone, fuming formic acid, acetic acid, ethyl acetate, ethyl oxalate, etc. Less sol. in nitrobenzene. Insol. in benzene and methyl iodide. (Duboin, C. R. 1905, **141**. 1017.)

+8 H_2O . Decomp. by H_2O . Very sol. in alcohols, glycerine, acetone, fuming formic acid, acetic acid, ethyl acetate, etc. without decomp. Sl. sol. in nitrobenzene. Insol. in benzene and methyl iodide. (Duboin, l. c.)
 +9 H_2O . Hygroscopic. Sol. in alcohol and acetone without decomp. (Dobroserdoff, C. C. **1901**, I. 664.)

Lithium nitride, Li_3N .

Sol. in H_2O with decomp. (Ouvrard, C. R. **114**. 120.)

Very hygroscopic. (Dafert, M. 1910, **31**. 987.)

Lithium oxide, Li_2O .

Slowly sol. in H_2O to form $LiOH$.

See Lithium hydroxide.

Lithium peroxide, Li_2O_2 .

(de Forcrand, C. R. 1900, **130**. 1467.)

Lithium hydrogen peroxide, $Li_2O_2, H_2O_2 + 3H_2O$.

Sol. in H_2O . Insol. in alcohol. (de Forcrand, C. R. 1900, **130**. 1466.)

Lithium selenide, Li_2Se .

Sol. in H_2O . (Fabre, C. R. **103**. 269.)

+9 H_2O . Sol. in H_2O . (Fabre.)

Lithium silicide, Li_4Si_3 .

Decomp. by H_2O and by dil. acids. Decomp. by aqueous solutions of alkalis with evolution of H_2 . (Moissan, C. R. 1902, **134**. 1083.)

Lithium monosulphide, Li_2S .

More sol. in H_2O or alcohol than LiOH .

Luteochromium bromide, $\text{Cr}(\text{NH}_3)_6\text{Br}_3$.

Less sol. in H_2O than the chloride. (Jørgensen, J. pr. (2) 30. 1.)

— **bromoplatinate**, $[\text{Cr}(\text{NH}_3)_6]_2(\text{PtBr}_6)_3 + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O . Insol. in alcohol. (Jørgensen.)

— **chloride**, $\text{Cr}(\text{NH}_3)_6\text{Cl}_3 + \text{H}_2\text{O}$.

Efflorescent, and very sol. in H_2O . (Jørgensen.)

— **chloroplatinate**.

(a) $[\text{Cr}(\text{NH}_3)_6]_2(\text{PtCl}_6)_3 + 6\text{H}_2\text{O}$. Nearly completely insol. in H_2O . (Jørgensen.)

(b) $\text{Cr}(\text{NH}_3)_6\text{Cl}(\text{PtCl}_6) + 2\frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O into above; insol. in alcohol. (Jørgensen.)

(c) $[\text{Cr}(\text{NH}_3)_6]_2\text{Cl}_4(\text{PtCl}_6) + 2\text{H}_2\text{O}$. Decomp. by H_2O into (a). (Jørgensen.)

— **mercuric chloride**, $\text{Cr}(\text{NH}_3)_6\text{Cl}_3, \text{HgCl}_2$.

Decomp. by H_2O ; sl. sol. in dil. $\text{HCl} + \text{Aq}$; insol. in alcohol.

$\text{Cr}(\text{NH}_3)_6\text{Cl}_3, 3\text{HgCl}_2 + 2\text{H}_2\text{O}$. Decomp. by dil. $\text{HCl} + \text{Aq}$ into above salt. (Jørgensen.)

— **chromicyanide**, $\text{Cr}(\text{NH}_3)_6\text{Cr}(\text{CN})_6$.

Precipitate.

— **cobalticyanide**, $\text{Cr}(\text{NH}_3)_6\text{Co}(\text{CN})_6$.

Nearly insol. in H_2O or in conc. $\text{HCl} + \text{Aq}$. (Jørgensen.)

— **ferrocyanide**, $\text{Cr}(\text{NH}_3)_6\text{Fe}(\text{CN})_6$.

Very sl. sol. in cold H_2O or dil. acids. (Jørgensen.)

— **iodide**, $\text{Cr}(\text{NH}_3)_6\text{I}_3$.

Sl. sol. in H_2O . (Jørgensen, l. c.)

— **iodosulphate**, $\text{Cr}(\text{NH}_3)_6\text{SO}_4\text{I}$.

Sol. in H_2O ; nearly insol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$ or alcohol. (Jørgensen.)

— **nitrate**, $\text{Cr}(\text{NH}_3)_6(\text{NO}_3)_3$.

Sol. in 35–40 pts. H_2O . Insol. in cold dil. $\text{HNO}_3 + \text{Aq}$ or alcohol. Can be crystallised out of H_2O containing a little HNO_3 . (Jørgensen, J. pr. (2) 30. 1.)

— **nitrate chloroplatinate**,

$\text{Cr}(\text{NH}_3)_6(\text{NO}_3)\text{PtCl}_6 + \text{H}_2\text{O}$.

Insol. in H_2O . Sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Jørgensen.)

— **nitratosulphate**, $\text{Cr}(\text{NH}_3)_6(\text{NO}_3)\text{SO}_4$.

Sol. in H_2O ; insol. in alcohol. (Jørgensen.)

Luteochromium oxalate, $[\text{Cr}(\text{NH}_3)_6]_2(\text{C}_2\text{O}_4)_3 + 4\text{H}_2\text{O}$.

Nearly insol. in cold H_2O . (Jørgensen.)

— **orthophosphate**, $\text{Cr}(\text{NH}_3)_6\text{PO}_4 + \text{H}_2\text{O}$.

Sl. sol. in H_2O ; easily sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Jørgensen.)

— **sodium pyrophosphate**,

$\text{Cr}(\text{NH}_3)_6(\text{NaP}_2\text{O}_7) + 11\frac{1}{2}\text{H}_2\text{O}$.

Nearly insol. in cold H_2O ; wholly sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Jørgensen.)

— **sulphate**, $[\text{Cr}(\text{NH}_3)_6]_2(\text{SO}_4)_3 + 5\text{H}_2\text{O}$.

Quite sol. in H_2O ; insol. in alcohol. (Jørgensen.)

— **sulphate chloroplatinate**,

$[\text{Cr}(\text{NH}_3)_6(\text{SO}_4)]_2\text{PtCl}_6$.

Nearly insol. in H_2O . (Jørgensen.)

Luteocobalt diamine chromium cyanide.

See Diamine chromium luteocobalt cyanide.

Luteocobaltic bromide, $\text{Co}(\text{NH}_3)_6\text{Br}_3$.

Sol. in H_2O . Precipitated from H_2O solution by dil. $\text{HBr} + \text{Aq}$. (J. pr. (2) 35. 417.)

— **bromopermanganate**,

$\text{Co}(\text{NH}_3)_6\text{Br}_3(\text{MnO}_4)_2$.

Easily sol. in H_2O . (Klobb, A. c. 5.)

— **bromoplatinate**, $\text{Co}(\text{NH}_3)_6\text{Br}_3, \text{H}_2\text{O}$.

Sl. sol. in H_2O ; can be recrystallised from hot H_2O containing HBr . (Jørgensen.)

— **bromosulphate**, $\text{Co}(\text{NH}_3)_6\text{Br}_3(\text{SO}_4)_2$.

Nearly insol. in H_2O . Very sl. sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Jørgensen.)

— **carbonate**, $[\text{Co}(\text{NH}_3)_6]_2(\text{CO}_3)_3$.

Efflorescent; easily sol. in H_2O .

$[\text{Co}(\text{NH}_3)_6]_2(\text{CO}_3)_3, \text{H}_2\text{CO}_3 + 5\text{H}_2\text{O}$. Sol. in H_2O than the neutral salt and Genth.)

— **chloride**, $\text{Co}(\text{NH}_3)_6\text{Cl}_3$.

Sol. in 17.09 pts. H_2O at 10.5° ; at 11.4° ; 16.48 pts. at 12° ; and more in hot H_2O . (F. Rose.)

100 pts. H_2O dissolve 4.26 pts. at 12.74° at 46.6° . (Kurnakoff, J. 24. 629.)

Not appreciably sol. in conc. HCl . (Jørgensen.)

Insol. in alcohol or solutions of HCl and chlorides. (Gibbs and Genth.)

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$.

solution is pptd. by alcohol, min-
alkali chlorides.

mercuric chloride, $\text{Co}(\text{NH}_3)_6\text{Cl}_3$,
 $1\frac{1}{2}\text{H}_2\text{O}$.

H_2O . (Krok, 1870.)
allizing from hot H_2O containing
rted into—

Cl_3 , $3\text{HgCl}_2 + \text{H}_2\text{O}$. Very sl. sol.
(Jørgensen.)

Cl_3 , $2\text{HgCl}_2 + \frac{1}{2}\text{H}_2\text{O}$. Sol. in hot
which it crystallizes on cooling.
d conc. $\text{HCl} + \text{Aq}$, and is pptd.
lution by HCl or alcohol. (Car-

xist. (Jørgensen.)

More easily sol. in cold H_2O and
ts than the preceding comp.
Berlin, 1861.)

xist. (Jørgensen.)

s chloride, $2\text{Co}(\text{NH}_3)_6\text{Cl}_3$,
 $10\text{H}_2\text{O}$.

ate, $\text{Co}(\text{NH}_3)_6\text{Cl}_3$, AuCl_3 .

. in cold, more easily in hot H_2O
 Cl . (Gibbs and Genth, Sill. Am. J.
)

late, $[\text{Co}(\text{NH}_3)_6\text{Cl}_2]_2\text{I}_4\text{O}_{11} + \text{H}_2\text{O}$.

lite, $\text{Co}(\text{NH}_3)_6$, IrCl_6 .

boiling H_2O or dil. $\text{HCl} + \text{Aq}$.

late, $2\text{Co}(\text{NH}_3)_6\text{Cl}_3$, 3IrCl_4 .
 2O . (Gibbs.)

hromate, $\text{Co}(\text{NH}_3)_6\text{CrO}_4\text{Cl} +$

). (Klobb, Bull. Soc. 1901, (3)

ioride, $\text{Co}(\text{NH}_3)_6\text{Cl}_2\text{F}$.

anorg. 1905, 43. 339.)

lladite, $2\text{Co}(\text{NH}_3)_6\text{Cl}_3$, 3PdCl_2 .

in dil. $\text{HCl} + \text{Aq}$. (Gibbs, Sill.
. 58.)

rchlorate, $\text{Co}(\text{NH}_3)_6\text{Cl}(\text{ClO}_4)_2$.

h, Gazz. ch. it. 1901, 31. (2)

rmanganate,

$6\text{Cl}_2(\text{MnO}_4)$.

rySTALLIZED from H_2O . (Klobb,
f.)

rmanganate ammonium chlor-
 $[\text{Co}(\text{NH}_3)_6\text{Cl}_2(\text{MnO}_4)]$, NH_4Cl .

$n \text{H}_2\text{O}$. (Klobb.)

**Luteocobaltic chloropermanganate potassium
chloride**, $\text{Co}(\text{NH}_3)_6\text{Cl}_2(\text{MnO}_4)$, KCl .

Very easily sol. in H_2O , with decomp. into
constituents; sol. in $\text{KCl} + \text{Aq}$. (Klobb.)

— **chloropermanganate sodium chloride**,
 $\text{Co}(\text{NH}_3)_6\text{Cl}_2(\text{MnO}_4)$, NaCl .

Very sol. in H_2O . (Klobb.)

— **chloroplatinate**, $2\text{Co}(\text{NH}_3)_6\text{Cl}_3$, $3\text{PtCl}_4 +$
 $6\text{H}_2\text{O}$.

Can be recrystallized from much hot H_2O .
(Gibbs and Genth.)

$+ 21\text{H}_2\text{O}$. (Gibbs and Genth.)

$\text{Co}(\text{NH}_3)_6\text{Cl}_3$, $\text{PtCl}_4 + \frac{1}{2}\text{H}_2\text{O}$. Very sl. sol.
in cold, decomp. by hot H_2O into—

$2\text{Co}(\text{NH}_3)_6\text{Cl}_3$, $\text{PtCl}_4 + 2\text{H}_2\text{O}$. By recrystallizing from hot H_2O containing HCl this salt is converted into the above salt. (Jørgensen.)

— **chlororhodite**.

Nearly insol. in boiling H_2O or dil. acids.
Sol. in conc. $\text{HCl} + \text{Aq}$. (Gibbs, Sill. Am. J.
(2) 37. 57.)

— **chlororuthenate**, $2\text{Co}(\text{NH}_3)_6\text{Cl}_3$, 3RuCl_4 .

Sol. in dil. acids. (Gibbs.)

— **chloroselenate**, $\text{Co}(\text{NH}_3)_6\text{ClSeO}_4 + 3\text{H}_2\text{O}$.

Decomp. by H_2O . (Klobb, Bull. Soc.
1901, (3) 25. 1029.)

— **chlorosulphate**, $\text{Co}(\text{NH}_3)_6\text{Cl}(\text{SO}_4)$.

Sol. in H_2O .

$+ 3\text{H}_2\text{O}$. Only sl. sol. in cold H_2O . (Klobb,
Bull. Soc. 1901, (3) 25. 1025.)

— **ammonium chlorosulphate**,

$[\text{Co}(\text{NH}_3)_6]_4\text{Cl}_2(\text{SO}_4)_5$, $3(\text{NH}_4)_2\text{SO}_4 +$
 $6\text{H}_2\text{O}$.

Decomp. by H_2O . (Klobb, Bull. Soc.
1901, (3) 25. 1027.)

— **chlorosulphate chloroplatinate**,

$2\text{Co}(\text{NH}_3)_6\text{Cl}(\text{SO}_4)$, PtCl_4 .

Very sl. sol. in cold pure H_2O . Can be re-
crystallized out of H_2O containing HCl .
(Krok.)

— **chlorosulphate mercuric chloride**,

$\text{Co}(\text{NH}_3)_6\text{Cl}(\text{SO}_4)$, HgCl_2 .

Scarcely sol. in pure H_2O , but can be crys-
tallized from warm acidified H_2O . (Krok.)

— **chlorosulphite**, $\text{Co}(\text{NH}_3)_6(\text{SO}_3)\text{Cl} +$
 $3\text{H}_2\text{O}$.

Sol. in H_2O . (Vortmann and Magdeburg,
B. 22. 2637.)

— **chromate**, $[\text{Co}(\text{NH}_3)_6]_2(\text{CrO}_4)_3 + 5\text{H}_2\text{O}$.

Ppt. Sol. in hot H_2O .

$[\text{Co}(\text{NH}_3)_6]_2(\text{Cr}_2\text{O}_7)_3 + 5\text{H}_2\text{O}$. Moderately
sol. in hot H_2O .

Luteocobaltic chromicyanide,
 $\text{Co}(\text{NH}_3)_6\text{Cr}(\text{CN})_6$.

Ppt. (Braun.)

— **cobalticyanide,** $\text{Co}(\text{NH}_3)_6\text{Co}(\text{CN})_6$.
 Ppt.

— **dithionate, basic,**
 $4[\text{Co}(\text{NH}_3)_6(\text{S}_2\text{O}_8)(\text{OH})], \text{Co}_2(\text{S}_2\text{O}_8)_2\text{O}$.
 Sol. in H_2O and dil. alcohol.

— **ferricyanide,** $\text{Co}(\text{NH}_3)_6\text{Fe}(\text{CN})_6 + \frac{1}{2}\text{H}_2\text{O}$.
 Insol. in H_2O . (Braun.)

— **fluoride,** $\text{Co}(\text{NH}_3)_6\text{F}_3$.
 Sl. sol. in cold H_2O . Nearly insol. in acids.
 (Böhm, Z. anorg. 1905, 43. 340.)

— **hydrogen fluoride,** $\text{Co}(\text{NH}_3)_6\text{H}_2\text{F}_4$.
 Sl. sol. in H_2O . Decomp. by hot H_2O .
 (Miolati and Rossi, Real. Ac. Linc. 1896, (5) 5. II, 185.)

— **hydrogen boron fluoride,**
 $\text{Co}(\text{NH}_3)_6\text{F}_3, 3\text{BF}_3, \text{HF}$.
 Cryst. from H_2O acidified with HF . (Miolati and Rossi.)

— **molybdenyl fluoride,**
 $\text{Co}(\text{NH}_3)_6\text{F}_3, 2 \text{MoO}_2\text{F}_2$.
 Cryst. from H_2O containing HF . (Miolati and Rossi.)

— **silicon fluoride,**
 $\text{Co}(\text{NH}_3)_6\text{F}_3, 2\text{SiF}_4$. (Miolati and Rossi.)

— **titanium hydrogen fluoride,**
 $2\text{Co}(\text{NH}_3)_6\text{F}_3, 3\text{TiF}_4, 2\text{HF}$. (Miolati and Rossi.)

— **tungstyl fluoride,**
 $\text{Co}(\text{NH}_3)_6\text{F}_3, 2\text{WO}_2\text{F}_2$. (Miolati and Rossi.)

— **uranyl fluoride,**
 $\text{Co}(\text{NH}_3)_6\text{F}_3, \text{UO}_2\text{F}_2$.
 Can be cryst. from H_2O containing HF .
 (Miolati and Rossi.)

— **vanadyl fluoride,**
 $2\text{Co}(\text{NH}_3)_6\text{F}_3, 5\text{VO}_2\text{F}_2, 7\text{HF}$.
 Ppt. (Miolati and Rossi.)

— **fluoride nitrate,**
 $\text{Co}(\text{NH}_3)_6\text{F}(\text{NO}_3)_2$.
 (Bohm, Z. anorg. 1905, 43. 336.)

— **hydroxide,** $\text{Co}(\text{NH}_3)_6(\text{OH})_3$.
 Known only in aqueous solution.

Luteocobaltic mercuric hydroxychlorid
 $\text{CoN}_6\text{H}_{14}(\text{HgCl})_2(\text{HgOH})\text{Cl}_2$.

Ppt. Easily decomp. (Vortman Morgulis, B. 22. 2644.)

$\text{CoN}_6\text{H}_{14}(\text{HgOH})_2\text{Cl}_2$. Ppt. (V. and B.)
 $\text{CoN}_6\text{H}_{14}(\text{HgOH})_2\text{Cl}_2$. Ppt. (V. and B.)

— **iodide,** $\text{Co}(\text{NH}_3)_6\text{I}_3$.

Insol. in cold, but moderately sol. H_2O .

According to Jörgensen, contains HN has the formula $\text{Co}_2(\text{NH}_3)_{12}\text{I}_6(\text{NO}_3)_2$.

— **iodosulphate,** $\text{Co}(\text{NH}_3)_6\text{I}(\text{SO}_4)$.

Can be recrystallized from hot H_2O sol. in warm, nearly insol. in cold (Krok, B. 4. 711.)

— **mercuriodide,** $\text{Co}_2\text{N}_{12}\text{H}_{22}(\text{HgI})_2\text{I}$.
 Ppt. (Vortmann and Borsbach.)
 $\text{CoN}_6\text{H}_{14}(\text{HgI})_2\text{I}_2$. Ppt. (V. and B.)

— **mercuriodide, basic,**
 $\text{CoN}_6\text{H}_{14}(\text{HgOH})_2\text{I}_2(\text{OH})$.
 Insol. in H_2O . Sl. sol. in H_2O .
 mann and Borsbach, B. 23. 2804.)

— **nitrate,** $\text{Co}(\text{NH}_3)_6(\text{NO}_3)_3$.

Sol. in H_2O . Can be recrystallized boiling H_2O . Sol. in about 60 pts insol. in conc. $\text{HNO}_3 + \text{Aq}$. (Jörgen pr. (2), 35. 417.)

Almost insol. in acids. (Rogojski, (3), 41. 454.)

Insol. in NH_4OH , HCl , and HNO_3 decomp. by $\text{H}_2\text{SO}_4 + \text{Aq}$. (Gibbs and $\text{Co}(\text{NH}_3)_6(\text{NO}_3)_3$, HNO_3 . Decom H_2O or dil. alcohol. (Jörgensen, J. pr. 63.)

— **nitrate chloroplatinate,**
 $\text{Co}(\text{NH}_3)_6(\text{NO}_3)\text{Cl}_2, \text{PtCl}_4 + \text{H}_2\text{O}$.
 Not decomp. by H_2O . (Jörgensen

— **nitratosulphate,** $\text{Co}(\text{NH}_3)_6(\text{NO}_3)_2$
 Sol. in H_2O . (Jörgensen.)

— **nitrite cobaltic nitrite,**
 $\text{Co}_2(\text{NH}_3)_{12}(\text{NO}_3)_6, \text{Co}_2(\text{NO}_3)_6 = \text{Co}(\text{NH}_3)_6(\text{NO}_3)_2\text{Co}$.
 Nearly insol. in H_2O . (Jörgensen.)
 Much less sol. in H_2O than the corresponding roseo salt. (Gibbs.)

— **diamine cobaltic nitrite,**
 $\text{Co}(\text{NH}_3)_6[\text{Co}(\text{NH}_3)_2(\text{NO}_3)_2]_2$.
 Ppt. (Gibbs.)
 $= \text{Co}(\text{NH}_3)_6[(\text{NO}_3)_2(\text{NH}_3)_2\text{Co}(\text{NO}_3)_2]$
 Nearly insol. in cold, sl. sol. in boiling (Jörgensen, Z. anorg. 5. 179.)

— **oxalate,** $[\text{Co}(\text{NH}_3)_6]_2(\text{C}_2\text{O}_4)_3 + 4\text{H}$
 Insol. in hot or cold H_2O . Easily $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$.

obaltic oxalate chloraurate,
 $2\text{Co}(\text{NH}_3)_6(\text{C}_2\text{O}_4)\text{Cl}, \text{AuCl}_3 + 4\text{H}_2\text{O}$.
 ly sol. in hot H_2O . (Gibbs.)

erchlorate, $\text{Co}(\text{NH}_3)_6(\text{ClO}_4)_3$.
 losevich, *Gazz. ch. it.* 1901, 31, (2)

ermanganate, $[\text{Co}(\text{NH}_3)_6]_2(\text{MnO}_4)_3$.
 ly insol. in H_2O . 100 pts. H_2O at 0°
 e only 0.072 pt. salt. Moderately
 hot H_2O . (Klobb, *A. ch.* (6) 12. 5.)

ersulphate sulphate,
 $[\text{Co}(\text{NH}_3)_6]_2\text{S}_2\text{O}_8(\text{SO}_4)_2$.
 h less sol. in H_2O than the sulphate.
 in 641 pts. H_2O at 18.8° and in 632
 20° . Not easily sol. even in boiling
 (Jørgensen, *Z. anorg.* 1898, 17. 459.)

orthophosphate, $\text{Co}(\text{NH}_3)_6(\text{PO}_4) +$
 $\frac{1}{2}\text{H}_2\text{O}$.
 l. in cold H_2O . Easily sol. in dil. acids.
 sen.)
 $[\text{Co}(\text{NH}_3)_6]_2(\text{PO}_4)(\text{PO}_4\text{H})_3 + 5\frac{1}{2}\text{H}_2\text{O}$ (?).
 Braun.)
 $[\text{Co}(\text{NH}_3)_6]_2(\text{PO}_4\text{H})_3 + 4\text{H}_2\text{O}$. Ppt. Easily
 very dil. $\text{HCl} + \text{Aq}$. (Jørgensen.)

etaphosphate.

pyrophosphate, $[\text{Co}(\text{NH}_3)_6]_2\text{P}_2\text{O}_7 +$
 $\frac{1}{2}\text{H}_2\text{O}$.
 bs, *Am. Acad. Proc.* 11. 29); or
 $[\text{Co}(\text{NH}_3)_6]_2\text{P}_2\text{O}_7(\text{ONa})_2$ (Vortmann, *B.* 11.
 or $\text{Co}(\text{NH}_3)_6(\text{P}_2\text{O}_7\text{Na}) + 11\frac{1}{2}\text{H}_2\text{O}$.
 sen, *J. pr.* (2) 35. 438.)
 nearly insol. in H_2O . With H_2O at
 s decomp. into—
 $[\text{Co}(\text{NH}_3)_6]_4(\text{P}_2\text{O}_7)_3 + 20\text{H}_2\text{O}$. Less easily
 n the preceding salt.

rophosphate, acid, $\text{Co}(\text{NH}_3)_6(\text{P}_2\text{O}_7\text{H})$.
 lly insol. in H_2O . Somewhat sol. in
 $\frac{1}{2}\text{H}_2\text{O} + \text{Aq}$. Easily sol. in $\text{HCl} + \text{Aq}$
 sen.)

dium pyrophosphate,
 $(\text{NH}_3)_6(\text{P}_2\text{O}_7\text{Na}) + 11\frac{1}{2}\text{H}_2\text{O}$.
 Not wholly insol. in cold H_2O .
 p. by hot H_2O . Less sol. in $\text{NH}_4\text{OH} +$
 n in H_2O . (Jørgensen.)
 $[\text{Co}(\text{NH}_3)_6]_4(\text{P}_2\text{O}_7)_3, 2\text{Co}(\text{NH}_3)_6(\text{NaP}_2\text{O}_7)$
 $\frac{1}{2}\text{O}$. As above. (Jørgensen.)

lenate, $[\text{Co}(\text{NH}_3)_6]_2(\text{SeO}_4)_3 + 5\text{H}_2\text{O}$.
 sol. in H_2O . (Klobb, *Bull. Soc.*
) 25. 1028.)

ydrogen selenate,
 $[\text{Co}(\text{NH}_3)_6]_2\text{H}(\text{SeO}_4)_3 + 2\frac{1}{2}\text{H}_2\text{O}$.
 lecomp. by H_2O . (Klobb.)

Luteocobaltic ammonium selenate,
 $[\text{Co}(\text{NH}_3)_6]_2(\text{SeO}_4)_3, (\text{NH}_4)_2\text{SeO}_4 +$
 $4\text{H}_2\text{O}$.

Very sol. in H_2O .
 $[\text{Co}(\text{NH}_3)_6]_2(\text{SeO}_4)_3, (\text{NH}_4)_2\text{SeO}_4 + 8\text{H}_2\text{O}$.
 Very sol. in H_2O . (Klobb.)

— **sulphocyanide**, $\text{Co}(\text{NH}_3)_6(\text{SCN})_3$.
 Decomp. by hot H_2O . (Miolati, *Z. anorg.*
 1900, 23. 241.)

— **mercuric sulphocyanide**,
 $\text{Co}(\text{NH}_3)_6(\text{SCN})_3, 2\text{Hg}(\text{SCN})_2$.
 Decomp. by H_2O .
 Cryst. from dil. $\text{NH}_4\text{SCN} + \text{Aq}$. (Miolati.)

— **platinum sulphocyanide**,
 $[\text{Pt}(\text{SCN})_4]_2[\text{Co}(\text{NH}_3)_6(\text{SCN})_3]_2$.
 Decomp. by hot H_2O . (Miolati.)

— **silver sulphocyanide**,
 $\text{Co}(\text{NH}_3)_6(\text{SCN})_3, 2\text{AgSCN}$.
 Decomp. by hot H_2O . (Miolati.)

— **sulphate**, $[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_4)_3 + 5\text{H}_2\text{O}$.
 Sl. sol. in cold, more easily in hot H_2O .
 $+ 6\text{H}_2\text{O}$. (Krok, *B.* 4. 711.)

— **hydrogen sulphate**,
 $\text{Co}(\text{NH}_3)_6\text{H}(\text{SO}_4)_3$.
 Decomp. by alcohol to sulphate. (Jör-
 gensen, *Z. anorg.* 1898, 17. 458.)

$4[(\text{Co}6\text{NH}_3)_2(\text{SO}_4)_3], 5\text{H}_2\text{SO}_4 + 10\text{H}_2\text{O}$.
 Very sol. in H_2O with decomp. into the normal
 sulphate.

When pulverized it seems to dissociate
 slowly in contact with abs. alcohol. (Klobb,
Bull. Soc. 1901, (3) 25. 1025.)

— **ammonium sulphate**,
 $[\text{Co}_6(\text{NH}_3)_6]_2(\text{SO}_4)_3, (\text{NH}_4)_2\text{SO}_4 + 8\text{H}_2\text{O}$.
 Sol. in H_2O with decomp. (Klobb.)

— **cerium sulphate**, $[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_4)_3,$
 $\text{Ce}_2(\text{SO}_4)_3 + 1\frac{1}{2}\text{H}_2\text{O}$.
 Very sl. sol. in cold, and practically insol.
 in boiling H_2O . Sol. in acids. (Gibbs, *Am.*
Ch. J. 15. 560.)
 $[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_4)_3, 3\text{Ce}(\text{SO}_4)_2 + \text{H}_2\text{O}$. As
 above. (Wing, *Sill. Am. J.* (2) 49. 363.)

— **lanthanum sulphate**, $[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_4)_3,$
 $\text{La}_2(\text{SO}_4)_3 + \text{H}_2\text{O}$.
 Sl. sol. in H_2O . (Wing.)

— **thallic sulphate**, $[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_4)_3,$
 $\text{Tl}_2\text{O}(\text{SO}_4)_3 + 5\text{H}_2\text{O}$.
 Decomp. by cold H_2O . (Gibbs.)

— **sulphate bromaurate**,
 $\text{Co}(\text{NH}_3)_6(\text{SO}_4)(\text{AuBr}_4)$.
 Very sl. sol. in H_2O with apparent decomp.
 Insol. in alcohol. (Jørgensen.)

Luteocobaltic sulphate chloraurate,Sl. sol. in H_2O . (Jörgensen.)

— **cobaltic sulphite**, $[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_3)_3$, $\text{Co}_2(\text{SO}_3)_3 + \text{H}_2\text{O} =$ **dichrocobaltic sulphite**, $[\text{Co}(\text{NH}_3)_3]_2(\text{SO}_3)_3 + 2\text{H}_2\text{O}$, which see.

$[\text{Co}(\text{NH}_3)_6]_2(\text{SO}_3)_3$, $2\text{Co}_2(\text{SO}_3)_3 + 15\text{H}_2\text{O} =$ **diamine cobaltic sulphite**, $[\text{Co}(\text{NH}_3)_3]_2(\text{SO}_3)_3 + 5\text{H}_2\text{O}$, which see.

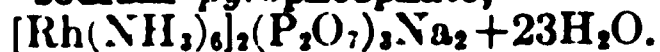
Luteorhodium bromide, $\text{Rh}(\text{NH}_3)_6\text{Br}_3$.Less sol. in H_2O than the chloride. (Jörgensen, J. pr. (2) 44. 51.)— **chloride**, $\text{Rh}(\text{NH}_3)_6\text{Cl}_3$.Sol. in 7 to 8 pts. H_2O at 8° . (J.)+ H_2O . Extremely efflorescent. (J.)— **rhodium chloride**, $\text{Rh}(\text{NH}_3)_6\text{Cl}_3$, RhCl_3 .Sol. in H_2O . (Jörgensen, Z. anorg. 5. 174.)— **chloroplatinate**, $2\text{Rh}(\text{NH}_3)_6\text{Cl}_3$, $3\text{PtCl}_4 + 6\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in warm $\text{HCl} + \text{Aq}$. (J.) $\text{Rh}(\text{NH}_3)_6\text{Cl}_3$, $\text{PtCl}_4 + \frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O into chloride and above salt. (J.)

— **nitrate**, $\text{Rh}(\text{NH}_3)_6(\text{NO}_3)_3$.

Sol. in 48 to 49 pts. H_2O at ord. temp. $\text{HNO}_3 + \text{Aq}$ diluted with 5 vols. H_2O ppt. the salt completely from aqueous solution. (Jörgensen, J. pr. (2) 44. 51.)

$\text{Rh}(\text{NH}_3)_6(\text{NO}_3)_3$, HNO_3 . Decomp. by H_2O or dil. alcohol. (Jörgensen, J. pr. (2), 44. 63.)

— **orthophosphate**, $\text{Rh}(\text{NH}_3)_6\text{PO}_4 + 4\text{H}_2\text{O}$.Sl. sol. in cold H_2O . (J.)— **sodium pyrophosphate**,Nearly wholly insol. in H_2O . Wholly insol. in $\text{NH}_4\text{OH} + \text{Aq}$. (J.)— **sulphate**, $[\text{Rh}(\text{NH}_3)_6]_2(\text{SO}_4)_3 + 5\text{H}_2\text{O}$.Sol. in 43 pts. H_2O at 20° . (J.)**Magnesium, Mg.**

Does not decomp. H_2O at ord. temp., but decomp. slowly at 100° . H_2O containing acids dissolves Mg easily. Sol. in cold dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Difficultly sol. in cold $\text{H}_2\text{SO}_4 + \text{Aq}$. (Bunsen.) Cold nitrosulphuric acid does not attack. (Bunsen.) Cold $\text{NH}_4\text{OH} + \text{Aq}$, $\text{KOH} + \text{Aq}$, or $\text{NaOH} + \text{Aq}$ do not attack. (Maak, Phipps.) Sol. in NH_4Cl or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Wöhler.)

Very rapidly sol. in $\text{K}_2\text{S}_2\text{O}_8 + \text{Aq}$ with violent evolution of gas. (Levi, Gazz. ch. it. 1908, 38. (1))

Attacked by aqueous solution of NH_4Cl , NaCl , LiCl , CuCl_2 , CdCl_2 , PbCl_2 , HgCl_2 , FeCl_3 , CrCl_3 , PtCl_4 , CuSO_4 , ZnSO_4 , FeSO_4 and MnSO_4 .

Sl. attacked by hot 30% $\text{CaCl}_2 + \text{Aq}$, by 30% $\text{MgCl}_2 + \text{Aq}$, $\text{BaCl}_2 + \text{Aq}$ and Aq . (Tommasi, Bull. Soc. 1899, (3), 1)

Not attacked by $\text{NH}_4\text{F} + \text{Aq}$, very by solutions of BaCl_2 , CaCl_2 , KCl , K_2F , NaNO_3 , $\text{Na}_2\text{S}_2\text{O}_3$ and Na_2HPO_4 . S of $\text{NaC}_2\text{H}_3\text{O}_2$, $\text{Na}_2\text{B}_4\text{O}_7$, alum and alum attack vigorously. Solution $(\text{NH}_4)_2\text{CO}_3$, NH_4Cl , $(\text{NH}_4)_2\text{C}_2\text{O}_4$, and Na_2CO_3 attack even more vigorously. (Mourmour, C. R. 1900, 130. 140.)

Insol. in liquid NH_3 . (Gore, Am. 1898, 20. 828.)

Insol. in liquid HF . (Franklin, Z. 1905, 46. 2.)

Somewhat sol. in liquid NH_3 , if metallic surface is in contact with the solvent. (Kraus, J. Am. Chem. Soc. 29. 1561.)

$\frac{1}{2}\text{ccm.}$ oleic acid dissolves 0.6 Mg in 6 days. (Gates, J. phys. Chem. 15. 143.)

Magnesium arsenide, Mg_3As_2 .

Decomp. on air. (Parkinson, Chem. 5. 127.)

Magnesium azoimide, $\text{Mg}(\text{N}_3)_2$.Decomp. by hot H_2O . (Curtius, 1898, (2) 58. 292.)**Magnesium boride, Mg_2B_3 .**Sol. in $\text{HCl} + \text{Aq}$. (Winkler, B. 23)**Magnesium bromide, MgBr_2 .**Deliquescent. Very sol. in H_2O with evolution of heat.Sat. $\text{MgBr}_2 + \text{Aq}$ contains at:

—18°	+17°	48°	62°	97°
52	58	60.9	62.5	65.8%

(Étard, A. ch. 1894, (7), 2. 54)

See also $\text{MgBr}_2 + 6\text{H}_2\text{O}$.Sp. gr. of $\text{MgBr}_2 + \text{Aq}$ at 19.5° contains:

5	10	15	20	25 %
1.043	1.087	1.137	1.191	1.247

30	35	40	45	50 %
1.31	1.377	1.451	1.535	1.625

(Kremers, Pogg. 108. 118, calculated by Gerlach, Z. anal. 8. 285.)

 $\text{MgBr}_2 + \text{Aq}$ is sl. decomp. by evap.

Solubility of MgBr_2 in alcohols.

Br_2 forms with methyl alcohol a complex, $\text{MgBr}_2 \cdot 6\text{CH}_3\text{OH}$.

Solubility of $\text{MgBr}_2 \cdot 6\text{CH}_3\text{OH}$ in CH_3OH at t° .

% by weight of $\text{MgBr}_2 \cdot 6\text{CH}_3\text{OH}$	t°	% by weight of $\text{MgBr}_2 \cdot 6\text{CH}_3\text{OH}$
42.6	130	63.6
44.6	140	66.8
46.7	150	70.2
48.9	160	74.0
51.4	170	78.5
55.5	180	84.5
58.0	185	88.0
60.7	190 mpt.	100

(Menschutkin, Z. anorg. 1907, 52. 11.)

Br_2 forms with ethyl alcohol a complex, $\text{MgBr}_2 \cdot 6\text{C}_2\text{H}_5\text{OH}$.

Solubility of $\text{MgBr}_2 \cdot 6\text{C}_2\text{H}_5\text{OH}$ in $\text{C}_2\text{H}_5\text{OH}$ at t° .

% by weight of $\text{MgBr}_2 \cdot 6\text{C}_2\text{H}_5\text{OH}$	t°	% by weight of $\text{MgBr}_2 \cdot 6\text{C}_2\text{H}_5\text{OH}$
17.2	80	73.8
24.9	85	76.2
32.7	90	78.7
40.3	95	82.3
47.8	100	86.7
55.1	103	90.0
62.2	106	94.4
68.8	108.5 mpt.	100
71.4		

(Menschutkin.)

Br_2 forms with propyl alcohol a complex, $\text{MgBr}_2 \cdot 6\text{C}_3\text{H}_7\text{OH}$.

Solubility of $\text{MgBr}_2 \cdot 6\text{C}_3\text{H}_7\text{OH}$ in $\text{C}_3\text{H}_7\text{OH}$ at t° .

% by weight of $\text{MgBr}_2 \cdot 6\text{C}_3\text{H}_7\text{OH}$	t°	% by weight of $\text{MgBr}_2 \cdot 6\text{C}_3\text{H}_7\text{OH}$
77.9	43	93.0
81.5	46	94.3
85.1	48	95.8
89.5	50	97.8
92.0	52 mpt.	100

(Menschutkin.)

Br_2 forms with isobutylalcohol a complex, $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_4\text{H}_9\text{OH}$.

Solubility of $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_4\text{H}_9\text{OH}$ in $\text{C}_4\text{H}_9\text{OH}$ at t° .

t°	% by weight of $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_4\text{H}_9\text{OH}$	t°	% by weight of $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_4\text{H}_9\text{OH}$
0	55.8	60	82.4
10	60.5	65	84.2
20	65.2	71	88.0
30	69.8	75	92.0
40	74.3	77	94.6
50	78.5	80 mpt.	100

(Menschutkin.)

MgBr_2 forms with isoamylalcohol a complex, $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_5\text{H}_{11}\text{OH}$.

Solubility of $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_5\text{H}_{11}\text{OH}$ in $\text{C}_5\text{H}_{11}\text{OH}$ at t° .

t°	% by weight of $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_5\text{H}_{11}\text{OH}$	t°	% by weight of $\text{MgBr}_2 \cdot 6(\text{iso})\text{C}_5\text{H}_{11}\text{OH}$
0	70.2	38	88.7
10	75.6	40	90.0
20	80.2	42	92.0
30	84.5	44	94.2
35	86.7	46 mpt.	100

(Menschutkin.)

Solubility in ether at t° .

t°	% MgBr_2	% $\text{MgBr}_2 \cdot 2\text{C}_4\text{H}_{10}\text{O}$
-8	0.6	1.08
0	0.8	1.44
+10	1.27	2.3
14	1.64	2.95
16	1.93	3.48
18	2.3	4.14
20	2.7	4.86
22 mpt.	3.22	5.80

(Menschutkin, Z. anorg. 1906, 49, 36.)

Solubility of $\text{MgBr}_2 \cdot 2\text{C}_4\text{H}_{10}\text{O}$ in ether at t° .

"Lower solution" = the melted $\text{MgBr}_2 \cdot 2\text{C}_4\text{H}_{10}\text{O}$ which does not mix with the ether above.

t°	Composition of lower solution		Composition of the upper layer	
	% MgBr_2	% $\text{MgBr}_2 \cdot 2\text{C}_4\text{H}_{10}\text{O}$	% MgBr_2	% $\text{MgBr}_2 \cdot 2\text{C}_4\text{H}_{10}\text{O}$
-10	42.0	75.7	1.8	3.2
0	41.0	73.9	2.3	4.1
+10	40.1	72.2	2.8	5.0
20	39.3	70.8	3.3	5.9
30	38.7	69.8	3.8	6.8
40	38.2	68.8	4.3	7.7
50	37.8	68.0	4.7	8.5
60	37.6	67.7	5.1	9.2
70	37.6	67.7	5.4	9.7
80	37.8	68.0	5.6	10.0
90	38.1	68.6	5.7	10.2

(Menschutkin.)

Solubility of MgBr_2 in formic acid.
 MgBr_2 forms with formic acid a complex, $\text{MgBr}_2 \cdot 6\text{HCOOH}$.

Solubility of $\text{MgBr}_2 \cdot 6\text{HCOOH}$ in HCOOH at t° .

t°	% by wt. $\text{MgBr}_2 \cdot 6\text{HCOOH}$
0	49.8
20	57.5
40	65.1
60	73.1
70	78.1
80	86.0
86	95.0
88 mpt.	100

(Menschutkin, Z. anorg. 1907, 54. 90.)

Solubility of MgBr_2 in acetic acid.
 MgBr_2 forms with acetic acid a complex, $\text{MgBr}_2 \cdot 6\text{CH}_3\text{COOH}$.

Solubility of $\text{MgBr}_2 \cdot 6\text{CH}_3\text{COOH}$ in CH_3COOH at t° .

t°	% by wt. $\text{MgBr}_2 \cdot 6\text{CH}_3\text{COOH}$
17	0.3
30	1.5
50	4.5
60	7.9
70	16.2
80	38.5
85	49.5
90	57.7
100	71.8
105	80.0
110	89.5
112 mpt.	100.0

(Menschutkin.)

Solubility of MgBr_2 in acetic anhydride.
 MgBr_2 forms with acetic anhydride a complex, $\text{MgBr}_2 \cdot 6(\text{CH}_3\text{CO})_2\text{O}$.

Solubility of $\text{MgBr}_2 \cdot 6(\text{CH}_3\text{CO})_2\text{O}$ in $(\text{CH}_3\text{CO})_2\text{O}$ at t° .

t°	% by wt. $\text{MgBr}_2 \cdot 6(\text{CH}_3\text{CO})_2\text{O}$
0	26.4
30	30.0
60	37.7
90	44.5
120	57.8
130	69.8
135	85.0
136–137 mpt.	100

(Menschutkin, Z. anorg. 1909, 61. 112.)

Solubility of MgBr_2 in acetone.
 MgBr_2 forms with acetone a complex, $\text{MgBr}_2 \cdot 3\text{CH}_3\text{COCH}_3$.

Solubility of $\text{MgBr}_2 \cdot 3\text{CH}_3\text{COCH}_3$ in CH_3COCH_3 at t° .

t°	% by wt. $\text{MgBr}_2 \cdot 3\text{CH}_3\text{COCH}_3$
0	0.2
30	0.8
60	1.45
70	2.0
72	3.7
73	5.5
74	14.0
75	50.0
76	71.6
80	83.3
84	89.8
88	95.2
92 mpt.	100

(Menschutkin, Z. anorg. 1907, 53. 3)

Solubility of MgBr_2 in acetamide.
 MgBr_2 forms with acetamide a complex, $\text{MgBr}_2 \cdot 6\text{CH}_3\text{CONH}_2$.

Solubility of $\text{MgBr}_2 \cdot 6\text{CH}_3\text{CONH}_2$ in CH_3CONH_2 at t° .

t°	% by wt. $\text{MgBr}_2 \cdot 6\text{CH}_3\text{CONH}_2$
50.5	56.0
70	57.8
90	60.5
110	65.0
130	71.5
150	80.0
160	85.5
165	90.0
169 mpt.	100.0

(Menschutkin, Z. anorg. 1909, 61. 1)

Solubility of MgBr_2 in acetanilide.
 MgBr_2 forms with acetanilide a complex, $\text{MgBr}_2 \cdot 6\text{CH}_3\text{CONHC}_6\text{H}_5$.

Solubility of $\text{MgBr}_2 \cdot 6\text{CH}_3\text{CONHC}_6\text{H}_5$ in $\text{CH}_3\text{CONHC}_6\text{H}_5$ at t° .

t°	% by wt. $\text{MgBr}_2 \cdot 6\text{CH}_3\text{CONHC}_6\text{H}_5$
107.5	9.0
140	19.3
170	29.6
185	39.0
195	49.0
200	59.5
205	73.2
209 mpt.	100.0

(Menschutkin, Z. anorg. 1909, 61.)

f MgBr_2 in aniline.
ns with aniline three complexes;
 H_5NH_2 ; MgBr_2 , $4\text{C}_6\text{H}_5\text{NH}_2$;
 $6\text{C}_6\text{H}_5\text{NH}_2$.

of these complexes in aniline.
 MgBr_2 , $6\text{C}_6\text{H}_5\text{NH}_2$ in $\text{C}_6\text{H}_5\text{NH}_2$.

	% by weight of MgBr_2 , $6\text{C}_6\text{H}_5\text{NH}_2$
	3.2
	3.9
	5.1
	7.5
	12.8
	18.5
	27.5

MgBr_2 , $4\text{C}_6\text{H}_5\text{NH}_2$ in $\text{C}_6\text{H}_5\text{NH}_2$.

	% by weight of MgBr_2 , $4\text{C}_6\text{H}_5\text{NH}_2$
	24.0
	24.3
	24.9
	26.0
	28.3
	33.5
	45.0
	55.0
	76.3

MgBr_2 , $2\text{C}_6\text{H}_5\text{NH}_2$ in $\text{C}_6\text{H}_5\text{NH}_2$.

	% by weight of MgBr_2 , $2\text{C}_6\text{H}_5\text{NH}_2$
	76.3
	77.3
	78.1
	79.0

kin, Z. anorg. 1907, 52. 159.)

f MgBr_2 in benzaldehyde.
ms with benzaldehyde a com-
 $3\text{C}_6\text{H}_5\text{CHO}$.

y of MgBr_2 , $3\text{C}_6\text{H}_5\text{CHO}$ in
 $\text{C}_6\text{H}_5\text{CHO}$ at t° .

MgBr_2 , CHO	t°	% by wt. MgBr_2 , $3\text{C}_6\text{H}_5\text{CHO}$
7	140	17.8
3	145	37.5
9	146	65.0
5	148	84.5
4	153	93.2
0	159 mpt.	100
5		

tkin, Z. anorg. 1907, 53. 26.)

Solubility of MgBr_2 in methylal.
 MgBr_2 forms with methylal a complex,
 MgBr_2 , $2\text{CH}_2(\text{OCH}_3)_2$.

Solubility of MgBr_2 , $2\text{CH}_2(\text{OCH}_3)_2$ in
 $\text{CH}_2(\text{OCH}_3)_2$ at t° .

t°	% by wt. MgBr_2 , $2\text{CH}_2(\text{OCH}_3)_2$
20	0.3
40	0.45
60	0.6
80	0.75
100	0.9
106	1.1
106	86.2
108	90.8
110	95.4
112 mpt.	100

(Menschutkin, Z. anorg. 1907, 53. 32.)

Solubility of MgBr_2 in dimethylcarbinol.
 MgBr_2 forms with dimethylcarbinol a
complex, MgBr_2 , $4(\text{CH}_3)_2\text{CHOH}$.

Solubility of MgBr_2 , $4(\text{CH}_3)_2\text{CHOH}$ in
 $(\text{CH}_3)_2\text{CHOH}$ at t° .

t°	% by weight of MgBr_2 , $4(\text{CH}_3)_2\text{CHOH}$	t°	% by weight of MgBr_2 , $4(\text{CH}_3)_2\text{CHOH}$
0	40.0	110	62.5
20	42.2	120	67.3
40	45.0	130	74.0
60	48.5	136	83.6
80	53.3	138	90.00
100	59.0	139 mpt.	100

(Menschutkin.)

Solubility of MgBr_2 in trimethylcarbinol.
 MgBr_2 forms with trimethylcarbinol a
complex, MgBr_2 , $4(\text{CH}_3)_3\text{COH}$.

Solubility of MgBr_2 , $4(\text{CH}_3)_3\text{COH}$ in
 $(\text{CH}_3)_3\text{COH}$ at t° .

t°	% by weight of MgBr_2 , $4(\text{CH}_3)_3\text{COH}$	t°	% by weight of MgBr_2 , $4(\text{CH}_3)_3\text{COH}$
24.4	0.06	65	50.5
25	1.0	70	62.5
35	9.5	75	77.0
45	19.1	77.5	85.0
55	32.2	79	91.5
60	40.5	80 mpt.	100

Menschutkin.)

Solubility of MgBr_2 in phenylhydrazine.
 MgBr_2 forms with $\text{C}_6\text{H}_5\text{NHNH}_2$ a complex,
 MgBr_2 , $6\text{C}_6\text{H}_5\text{NHNH}_2$.

Solubility of $\text{MgBr}_2 \cdot 6\text{C}_6\text{H}_5\text{NHNH}_2$ in $\text{C}_6\text{H}_5\text{NHNH}_2$ at t° .	
t°	% by wt. of $\text{MgBr}_2 \cdot 6\text{C}_6\text{H}_5\text{NHNH}_2$
20	3.0
40	7.0
60	16.4
80	33.0
99	54.8
100	54.8
140	60.8
180	68.4
200	73.4
(Menschutkin, Z. anorg. 1907, 52. 162.)	
Solubility of MgBr_2 in urea. MgBr_2 forms with urea a complex, $\text{MgBr}_2 \cdot 6\text{CO}(\text{NH}_2)_2$.	
Solubility of $\text{MgBr}_2 \cdot 6\text{CO}(\text{NH}_2)_2$ in urea at t° .	
t°	% by wt. $\text{MgBr}_2 \cdot 6\text{CO}(\text{NH}_2)_2$
108.5	24.2
115	29.8
120	35.0
125	41.6
127	45.5
130	60.0
(Menschutkin, Z. anorg. 1909, 61. 116.)	
$\text{MgBr}_2 \cdot 6\text{CO}(\text{NH}_2)_2$ decomposes at 130° giving $\text{MgBr}_2 \cdot 4\text{CO}(\text{NH}_2)_2$.	
Solubility of $\text{MgBr}_2 \cdot 4\text{CO}(\text{NH}_2)_2$ in urea at t° .	
t°	% by wt. $\text{MgBr}_2 \cdot 4\text{CO}(\text{NH}_2)_2$
130	58.0
145	60.7
160	67.2
165	71.4
170	83.7
171	96.0
(Menschutkin, Z. anorg. 1909, 61. 116.)	
Solubility of MgBr_2 in urethane. MgBr_2 forms with urethane a complex, $\text{MgBr}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$.	
Solubility of $\text{MgBr}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$ in $\text{NH}_2\text{COOC}_2\text{H}_5$ at t° .	
t°	% by wt. $\text{MgBr}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$
35	43.3
50	45.6
70	51.3
80	56.2
85	59.8
90	66.5
*91.5	75.5

* Mpt. of $\text{MgBr}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$.
(Menschutkin, Z. anorg. 1909, 61. 113.)

$\text{MgBr}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$ decorr $90.5\text{--}91^\circ$ forming $\text{MgBr}_2 \cdot 4\text{NH}_2\text{COOC}_2\text{H}_5$	
Solubility of $\text{MgBr}_2 \cdot 4\text{NH}_2\text{COOC}_2\text{H}_5$ at t° .	
t°	% by wt. $4\text{NH}_2\text{COOC}_2\text{H}_5$
91	69.4
100	73.8
110	80.0
115	84.1
120	90.0
123 mpt.	100.0
(Menschutkin.)	
$+6\text{H}_2\text{O}$.	
Solubility of $\text{MgBr}_2 + 6\text{H}_2\text{O}$ in 1	
t°	% by weight of $\text{MgBr}_2 + 6\text{H}_2\text{O}$
0	76.0
20	78.1
40	80.2
60	82.3
80	84.4
100	86.6
(Menschutkin, Z. anorg. 1907, 52. 162.)	
Sp. gr. of solution sat. at 18° = contains 50.8% MgBr_2 . (Mylus B. 1897, 30. 1718.)	
Sl. sol. in liquid NH_3 . (Franklin J. 1898, 20. 828.)	
Sol. in alcohol. Sol. in acetomann, B. 1904, 37. 4328; Eidmann, 1899, II. 1014.)	
Difficultly sol. in methyl acetomann, B. 1909, 42. 3790.)	
Insol. in ethyl acetate. (Nasch, 1910, 43. 314.)	
$+10\text{H}_2\text{O}$. Sol. in H_2O . (Panfili, Soc. 26. 234.)	
Magnesium manganous bromide, $2\text{MnBr}_2 + 12\text{H}_2\text{O}$.	
Deliquescent. (Saunders, Am. 150.)	
Magnesium mercuric bromide, $\text{MgBr}_2 \cdot 2\text{HgBr}_2$.	
Deliquescent. Not deliquescent.	
Magnesium molybdenyl bromide, $\text{MoOBr}_2 + 7\text{H}_2\text{O}$.	
(Weinland and Knoll, Z. anorg. 112.)	
Magnesium potassium bromide, $2\text{KBr} + 6\text{H}_2\text{O}$.	
Easily sol. in H_2O , from which tallises at 75 to 87° . Alcohol dissolves. (Löwig, Repert. 23. 231.)	

ula is MgBr_2 , $\text{KBr} + 6\text{H}_2\text{O}$. De-
ent. (Lerch, J. pr. (2) 28. 338.)

sium stannic bromide.
bromostannate, magnesium.

sium chloride, MgCl_2 .
quescent. Very sol. in H_2O with
on of heat. The solution decomposes
poration losing HCl , when less than 6
 H_2O are present to 1 mol. MgCl_2 .
eca, C. R. 37. 350.)
ydrous. Sol. in 1.857 pts. H_2O at 15° .
ch.)
1 pt. cold H_2O . (Fourcroy.)
 $\text{MgCl}_2 + \text{Aq}$ at 12.5° contains 64.8% MgCl_2 .
(fratz.)
ta. H_2O at 15.5° dissolve 200 pts. MgCl_2 . (Ure's
pts. H_2O dissolve 52.2 pts. MgCl_2 at 0°
p. gr. of sat. solution = 1.3619 at 15° .
l, Bull. Soc. (2) 47. 318.)
0 mols. H_2O dissolve 108 mols. MgCl_2 ,
C. (Löwenherz, Z. phys. Ch. 1894, 13.

00 mols. $\text{MgCl}_2 + \text{Aq}$ contain at t° :
° 67.5 68.5 68.7 79.5 79.95
 MgCl_2 11.58 11.92 11.71 12.28 12.39
° 116.67 152.6 181-2 186
 MgCl_2 16.2 18.24 23.8 24.1-24.4
Hoff and Meyerhoffer, B. A. B. 1897,
73.)

lubility of MgCl_2 in H_2O at t° .

	% MgCl_2	Solid phase
	11.4	Ice
	16.	"
	19.4	"
5	20.6	Ice + $\text{MgCl}_2 \cdot 12\text{H}_2\text{O}$
	26.7	$\text{MgCl}_2 \cdot 12\text{H}_2\text{O}$
4	30.5	"
3	31.6	$\text{MgCl}_2 \cdot 12\text{H}_2\text{O} +$ $\text{MgCl}_2 \cdot 8\text{H}_2\text{O}$
4	34.3	$\text{MgCl}_2 \cdot 8\text{H}_2\text{O} + \text{MgCl}_2 \cdot$ $6\text{H}_2\text{O}$
	34.6	$\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$
	34.9	"
	35.3	"
	35.6	"
	36.5	"
	37.9	"
	39.8	"
	42.2	"
7	46.1	$\text{MgCl}_2 \cdot 6\text{H}_2\text{O} + \text{MgCl}_2 \cdot$ $4\text{H}_2\text{O}$
6	49.1	$\text{MgCl}_2 \cdot 4\text{H}_2\text{O}$
5	55.8	$\text{MgCl}_2 \cdot 4\text{H}_2\text{O} + \text{MgCl}_2 \cdot$ $2\text{H}_2\text{O}$
	56.1	$\text{MgCl}_2 \cdot 2\text{H}_2\text{O}$

kolt-Börnstein, Tab. 5th Ed. 1912. 480.)
also $\text{MgCl}_2 + 6\text{H}_2\text{O}$.

Sp. gr. of $\text{MgCl}_2 + \text{Aq}$ at 15° .

% MgCl_2	Sp. gr.	% MgCl_2	Sp. gr.	% MgCl_2	Sp. gr.
1	1.0084	13	1.1130	25	1.2274
2	1.0169	14	1.1220	26	1.2378
3	1.0253	15	1.1311	27	1.2482
4	1.0338	16	1.1404	28	1.2586
5	1.0422	17	1.1498	29	1.2690
6	1.0510	18	1.1592	30	1.2794
7	1.0597	19	1.1686	31	1.2903
8	1.0684	20	1.1780	32	1.3012
9	1.0772	21	1.1879	33	1.3121
10	1.0859	22	1.1977	34	1.3230
11	1.0949	23	1.2076	35	1.3340
12	1.1040	24	1.2175

(Gerlach, Z. anal. 8. 281.)

Sp. gr. of $\text{MgCl}_2 + \text{Aq}$ at 18° .

% MgCl_2	Sp. gr.	% MgCl_2	Sp. gr.	% MgCl_2	Sp. gr.
5	1.0416	20	1.1764	34	1.3210
10	1.0859	30	1.2779

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of $\text{MgCl}_2 + \text{Aq}$ at 0° . S = pts. salt in
100 pts. of solution; S_1 = mols. salt in 100
mols. solution.

S	S_1	Sp. gr.
29.2056	7.230	1.2788
20.9293	4.762	1.1927
15.7989	3.423	1.1427
11.3249	2.355	1.1007
6.2008	1.233	1.0545

(Charpy, A. ch. (6) 29.'23.)

Sp. gr. of $\text{MgCl}_2 + \text{Aq}$ at 19.5° .

Pts. MgCl_2 in 100 pts. H_2O	Sp. gr.	Pts. MgCl_2 in 100 pts. H_2O	Sp. gr.
10.7	1.0826	35.3	1.2388
22.0	1.1592	51.5	1.3235

(Kremera, Pogg. 104. 155.)

Sp. gr. of MgCl ₂ +Aq at 14°.					
% MgCl ₂ +6H ₂ O	Sp. gr.	% MgCl ₂ +6H ₂ O	Sp. gr.	% MgCl ₂ +6H ₂ O	Sp. gr.
0	0.9993	17	1.0682	34	1.1407
1	1.0033	18	1.0724	35	1.1451
2	1.0073	19	1.0765	36	1.1495
3	1.0113	20	1.0807	37	1.1540
4	1.0154	21	1.0849	38	1.1584
5	1.0194	22	1.0891	39	1.1628
6	1.0234	23	1.0933	40	1.1673
7	1.0274	24	1.0976	41	1.1718
8	1.0314	25	1.1018	42	1.1763
9	1.0355	26	1.1061	43	1.1809
10	1.0395	27	1.1103	44	1.1855
11	1.0435	28	1.1146	45	1.1901
12	1.0476	29	1.1189	46	1.1948
13	1.0517	30	1.1232	47	1.1995
14	1.0558	31	1.1275	48	1.2042
15	1.0599	32	1.1319
16	1.0641	33	1.1363

(Oudemans, Z. anal. 7. 420.)

Sp. gr. of MgCl ₂ +Aq at 24°.					
% MgCl ₂ +6H ₂ O	Sp. gr.	% MgCl ₂ +6H ₂ O	Sp. gr.	% MgCl ₂ +6H ₂ O	Sp. gr.
2	1.0069	30	1.1062	58	1.2167
4	1.0138	32	1.1137	60	1.2252
6	1.0207	34	1.1212	62	1.2338
8	1.0276	36	1.1288	64	1.2425
10	1.0345	38	1.1364	66	1.2513
12	1.0415	40	1.1441	68	1.2602
14	1.0485	42	1.1519	70	1.2692
16	1.0556	44	1.1598	72	1.2783
18	1.0627	46	1.1677	74	1.2875
20	1.0698	48	1.1756	76	1.2968
22	1.0770	50	1.1836	78	1.3063
24	1.0842	52	1.1918	80	1.3159
26	1.0915	54	1.2000
28	1.0988	56	1.2083

(Gerlach, Z. anal. 8. 283. Calculated from Schiff.)

Sp. gr. of MgCl ₂ +Aq at 25°.	
Concentration of MnCl ₂ +Aq.	Sp. gr.
1-normal	1.1375
1/2 " "	1.0188
1/5 " "	1.0091
1/5 " "	1.0043

(Wagner, Z. phys. Ch. 1890, 5. 38.)

Sp. gr. of MgCl ₂ +Aq.	
1/2 MgCl ₂ g. in 1000 g. of solution	Sp. gr. 16°
0	1.0000
0.4400	1.0003
0.8801	1.0007
1.7780	1.0014
3.4533	1.0028
7.4691	1.0062
14.7187	1.0122
29.6307	1.0246

(Dijken, Z. phys. Ch. 1897, 24.

Sp. gr. of MgCl₂ at 20.1°.
p = per cent strength of solution
served density; w = volume conc.
per cc. ($\frac{pd}{100} = w.$)

p	d	
28.83	1.2569	0.
25.59	1.2241	0.
20.31	1.1735	0.
15.79	1.1324	0.
10.185	1.0833	0.
8.058	1.0650	0.
5.919	1.0473	0.
3.913	1.0304	0.
3.903	1.0240	0.
1.743	1.0126	0.

(Barnes, J. Phys. Chem. 1898, 2

Sp. gr. of MgCl ₂ +Aq at t	
t°	Concentration of MgCl ₂ +Aq
23	1 pt. MgCl ₂ in 8.1874 pts. H ₂ O
24	1 pt. " " 102.1 " "

(Hittorf, Z. phys. Ch. 1902, 20.

Sp. gr. of MgCl₂ at 0°.
G. MgCl₂ in 100 ccm. of solution 6.71
Sp. gr. 1.06

G. MgCl₂ in 100 ccm. of sol. 13.811
Sp. gr. 1.110

(Bremer, C. C. 1902, I. 29

Sp. gr. of MgCl₂+Aq at 20° cont
g. mols. MgCl₂ per l.
M 0.00493 0.007327 0.01
Sp. gr. 1.000344 1.000524 1.000842

M 0.05108 0.07171 0.10
Sp. gr. 1.004224 1.006036 1.008505

M . 0.50 0.75 0.9415
Sp. gr. 1.038496 1.056905 1.069317

(Jones and Pearce, Am. Ch. J. 1907,

$\text{gCl}_2 + \text{Aq}$ containing 10% MgCl_2 boils at 1° ; containing 20% MgCl_2 boils at 106.2° ; containing 30% MgCl_2 boils at 115.6° . (Gerlach, Z. anal. 26. 426.)

$\text{MgCl}_2 + \text{Aq}$ forms a crust at 122.5° , contains 52.9 pts. MgCl_2 to 100 pts. H_2O . (Gerlach, Z. anal. 26. 426.)

t. of $\text{MgCl}_2 + \text{Aq}$. P = pts. MgCl_2 to 100 pts. H_2O .

pt.	P	B.-pt.	P	B.-pt.	P
1°	4.9	111°	34.6	121°	50.8
2	9.2	112	36.6	122	52.2
3	13.2	113	38.4	123	53.6
4	16.7	114	40.2	124	55.0
5	19.9	115	41.8	125	56.4
6	22.5	116	43.4	126	57.7
7	25.0	117	44.9	127	59.0
8	27.5	118	46.4	128	60.3
9	29.9	119	47.9	129	61.6
0	32.3	120	49.4	130	62.9

(Gerlach, Z. anal. 26. 440.)

pt. of $\text{MgCl}_2 + \text{Aq}$ containing % MgCl_2 .

MgCl_2	B.-pt.	% MgCl_2	B.-pt.
1.6	101°	11.6	103°
3.4	102	14.3	104

(Skinner, Chem. Soc. 61. 341.)

l. in $\text{KCl} + \text{Aq}$ at 50° . (Uhlig, C. C. , II. 749.)

in 7 pts. alcohol at 15° . (Bergmann.)
5 " " moderate heat. (B)

pts. alcohol of given sp. gr. dissolve pts. MgCl_2 :

gr.	Pts. MgCl_2	Sp. gr.	Pts. MgCl_2
900	21.25	0.834	36.25
848	23.75	0.817	50.00

(Kirwan.)

$\text{Cl}_2 + 6\text{H}_2\text{O}$ is sol. in 5 pts. alcohol of 0.90 sp. gr. in 2 pts. alcohol of 0.817 sp. gr.

in 0.1828 pt. strong alcohol at 82.5° . (Wenzel.)

-pt. of an alcoholic solution of MgCl_2 .

% MgCl_2	B.-pt.
5.56	$78.43^\circ + 0.73^\circ$
8.53	" +1.34
9.62	" +1.77
13.84	" +3.54

(Skinner, Chem. Soc. 61. 341.)

ven more sol. in acetic ether than CaCl_2 . (C. R. 102. 363.)

l. in boiling amyl alcohol. (Riggs, Sill. J. 144. 103.)

sol. in anhydrous pyridine. Sol. in 95% and 93% pyridine + Aq. (Kahlen- J. Am. Chem. Soc. 1908, 30. 1107.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

+ $2\text{H}_2\text{O}$. Very deliquescent. (Ditte, A. ch. 1881, (5) 22. 560.)

+ $4\text{H}_2\text{O}$. (van't Hoff and Meyerhoffer.)

+ $6\text{H}_2\text{O}$. Deliquescent. Sol. in 0.6 pt. cold, and 0.273 pt. hot H_2O . (Casaseca, l. c.)

Solubility in H_2O at t° .

t°	1000 mols H_2O dissolve mols MgCl_2	100 g. H_2O dissolve g. MgCl_2
3.5	99.6	52.65
25.0	104.5	55.26
50.0	110.6	58.66

(Biltz and Marcus, Z. anorg. 1911, 71. 169.)

Solubility in $\text{KCl} + \text{NH}_4\text{Cl} + \text{Aq}$ at 25° has been studied. (Biltz and Marcus, Z. anorg. 1911, 71. 178.)

When the solid phases are $\text{MgSO}_4 + 6\text{H}_2\text{O}$ and $\text{MgCl}_2 + 6\text{H}_2\text{O}$, 1000 mols. H_2O dissolve 104 mols. MgCl_2 and 14 mols. MgSO_4 at 25° . (Löwenherz, Z. phys. Ch. 1894, 13. 480.)

Solubility of $\text{MgCl}_2 + 6\text{H}_2\text{O}$ in $(\text{NH}_4)\text{MgCl}_2 + 6\text{H}_2\text{O} + \text{Aq}$ at t° .

t°	Per 1000 mols H_2O	
	Mols NH_4Cl	Mols MgCl_2
3.5	0.5	99.5
25.0	0.5	103.8
50.0	0.8	111.2

(Biltz and Marcus, Z. anorg. 1911, 71. 170.)

Solubility data of $\text{MgCl}_2 + \text{KCl} + \text{MgKCl}_2$ are given by van't Hoff and Meyerhoffer. (Z. phys. Ch. 1899, 30. 64.)

+ $8\text{H}_2\text{O}$. Pptd. from an aqueous solution which contains about 10 mols. H_2O to 1 mol. MgCl_2 .

+ $12\text{H}_2\text{O}$. Pptd. from an aq. solution which contains 1 mol. MgCl_2 in about 12.06 mols. of H_2O . (Bogorodsky, C. C. 1899, I. 246.)

Magnesium manganous chloride, $\text{MgCl}_2, 2\text{MnCl}_2 + 12\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O and alcohol. (Saunders, Am. Ch. J. 14. 148.)

$2\text{MgCl}_2, \text{MnCl}_2 + 12\text{H}_2\text{O}$. Ppt. Deliquesces in the air. (Gossner, C. C. 1904, I. 707.)

Magnesium mercuric chloride, $\text{MgCl}_2, \text{HgCl}_2 + 6\text{H}_2\text{O}$.

Very deliquescent. More sol. than the following salt. (v. Bonsdorff, Pogg. 17. 133.)

$\text{MgCl}_2, 3\text{HgCl}_2 + 5\text{H}_2\text{O}$. Sol. in H_2O with-

out decomp. Easily sol. in alcohol. (v. Bonsdorff.)

Magnesium phosphoryl chloride, $\text{MgCl}_2, \text{POCl}_3$.

Deliquescent. Sol. in H_2O with evolution of heat and decomposition. Very sl. sol. in warm POCl_3 . (Casselmann, A. 98. 223.)

Magnesium potassium chloride, $\text{MgCl}_2, 2\text{KCl}+6\text{H}_2\text{O}$.

Deliquescent, forming a solution of MgCl_2 , while KCl remains undissolved. 100 pts. H_2O dissolve 64.5 pts. at 18.75° . 20 pts. salt dissolved in 80 pts. H_2O lower the temp. 1.75° . (Bischof.) Alcohol dissolves out MgCl_2 . Decomp. into the two salts by solution in H_2O . (Marcet.)

A sat. solution in contact with solid KCl and KCl , MgCl_2 , H_2O at 50° contains 79.5 mol. MgCl_2 and 14.9 mol. KCl per 1000 mol. H_2O . A sat. solution in contact with solid MgCl_2 , $6\text{H}_2\text{O}$ and KCl , MgCl_2 , H_2O at 50° contains 111.9 mol. MgCl_2 and 1.2 mol. KCl per 1000 mol. H_2O . (Uhlig, Chem. Soc. 1913, 104. (2) 775; C. B. Miner. 1913, 417.)

Min. *Carnallite*.

Magnesium rubidium chloride, $\text{MgCl}_2, \text{RbCl}+6\text{H}_2\text{O}$.

Not decomp. by a small quantity of H_2O . (Feit and Kubierscky, Ch. Ztg. 16. 335.)

Magnesium sodium chloride, $\text{MgCl}_2, \text{NaCl}+2\text{H}_2\text{O}$.

Sol. in H_2O . (Poggiale.)

Magnesium thallic chloride, $2\text{TlCl}_3, \text{MgCl}_2+6\text{H}_2\text{O}$.

Hygroscopic. Can be cryst. from H_2O . (Gewecke, A. 1909, 366. 224.)

Magnesium stannic chloride.

See Chlorostannate, magnesium.

Magnesium vanadium chloride, $\text{MgCl}_2, \text{VCl}_3+\text{H}_2\text{O}$.

Difficultly sol. in H_2O and alcohol. (Stähler, B. 1904, 37. 4412.)

Magnesium zinc chloride, $\text{MgCl}_2, \text{ZnCl}_2+6\text{H}_2\text{O}$.

Deliquescent; sol. in H_2O . (Warner, C. N. 27. 271.)

Magnesium chloride ammonia, $\text{MgCl}_2, 4\text{NH}_3$.

Easily decomp. (Clark, A. 78. 369.)

Magnesium chloride hydroxylamine, $\text{MgCl}_2, 2\text{NH}_2\text{OH}+2\text{H}_2\text{O}$.

100 g. of solution in H_2O contain 44.4% at 20° . (Antonow, J. Russ. Phys. Chem. Soc. 1905, 37. 478.)

Magnesium fluoride, MgF_2 .

1 l. H_2O dissolves 76 mg. MgF_2 . (Kohlrausch, Z. phys. Ch. 1904, 60. 38) 87 mg. are dissolved in 1 l. of sat. at 18° . (Kohlrausch, Z. phys. Ch. 11. 168.)

Scarcely sol. in acids. (Gay-Lussac, Thénard.) Insol. in excess of HF . precipitated, is sol. in aqueous sol. ammonium and magnesium salts. S. HNO_3+Aq , from which it is precip. alcohol.

Insol. in methyl acetate. (Naur 1909, 42. 3790.)

Min. *Sellaite*.

Magnesium potassium fluoride, MgF_2 .

Decomp. by H_2SO_4 . (Duboin, C. 120. 679.)

$\text{MgF}_2, 2\text{KF}$. Decomp. by H_2SO_4 (Duboin.)

Magnesium sodium fluoride, MgF_2 .

Insol. in H_2O . (Geuther, J. B. 11)

Magnesium stannic fluoride.

See Fluostannate, magnesium.

Magnesium titanium fluoride.

See Fluotitanate, magnesium.

Magnesium zirconium fluoride.

See Fluozirconate, magnesium.

Magnesium hydrosulphide, MgS_2H_2 .

Known only in aqueous solution decomposes on warming. Solution containing 16% MgS_2H_2 has sp. gr. 1.11 (Divers and Shimidzu, Chem. Soc.)

Magnesium hydroxide, MgO_2H_2 .

MgO is sol. in 55,368 pts. H_2O at ordinary also at 100° . (Fresenius, A. 50. 117.)

MgO is sol. in 5142 pts. H_2O at 15.5° (Fy pts. at 15.8° (Henry, J. Pharm. 12. 2); in 79 wan); in 16,000 pts. (Dalton); in 100,000-cold H_2O (Bineau); in 36,000 pts. boiling Ed. Phil. J. 5. 305.)

Calculated from electrical conduct. $\text{MgO}_2\text{H}_2+\text{Aq}$, 1 l. H_2O dissolves MgO_2H_2 at 18° . (Kohlrausch and phys. Ch. 12. 241.)

Calculated from electrical conduct. 1 l. H_2O dissolves 00.076 g. MgO_2H_2 (Dupré and Brutus, Z. angew. Ch. 55.)

Presence of CaO_2H_2 or CaSO_4 decrease the solubility. (Henry.) For the salts of the alkali metals, especially ammonium salts, increase the solubility in conc. Na_2SO_4 , NaNO_3 , NaCl , or Aq . (Karsten.) Sol. in NH_4OH insol. in $\text{KOH}+\text{Aq}$. (Odling.)

Easily sol. in acids. Sol. in a solution of sugar. Boiling alcohol traces.

y of MgO_2H_2 in $NH_4Cl + Aq$ at 29° .

(Normal)	Normality of		G. per l.	
	MgO_2H_2	NH_4Cl	MgO_2H_2	NH_4Cl
35	0.156	0.388	4.55	20.86
8	0.108	0.250	3.15	13.39
35	0.089	0.172	2.60	9.21
8	0.0638	0.106	1.86	5.67
8	0.049	0.0771	1.43	4.13

Muhs, Z. anorg. 1909, 38. 140.)

MgO_2H_2 in $NH_4NO_3 + Aq$ at 29° .

re- d er- [H, re- al)	Normality of		G. per l.	
	MgO_2H_2	NH_4NO_3	MgO_2H_2	NH_4NO_3
08	0.0833	0.1834	2.43	14.69
08	0.0495	0.076	1.45	6.09

(Herz and Muhs.)
y insol. in 16% $NaCl + Aq$ in 0.8 g. $NaOH$. (Maigret, Bull. 331.)

MgO_2H_2 in $NaCl + NaOH + Aq$.

G. MgO per l. of solution with added	
0.8 g. $NaOH$ per l.	4.0 g. $NaOH$ per l.
0.07	0.03
0.045	...
None	None

(Maigret.)
otd. $Mg(OH)_2$ is sol. in $Th(NO_3)_4$ as a colloidal solution. (Halla, 12, 79. 262.)
etone. (Naumann, B. 1904, 37. ann, C. C. 1899, II, 1014.)
Magnesium oxide.
ite. Sol. in cold citric acid + Aq . N. 37. 14.)
 I_2O . (Bender, B. 3. 932.)
iodide, MgI_2 .
rescent.
in H_2O . See $MgI_2 + 6$, and $8H_2O$.
 $MgI_2 + Aq$ at 19.5° containing:
15 20 25 30% MgI_2 ,
1.139 1.194 1.254 1.32
45 50 55 60% MgI_2 .
1.568 1.668 1.78 [1.915
s, Pogg. 111. 62, calculated by
erlach, Z. anal. 8. 285.)
decomp. slightly on evaporation.
in liquid NH_3 . (Franklin, Am. 20. 828.)
cohol, ether, and wood-spirit.

Solubility of MgI_2 in alcohols.
 MgI_2 forms with methyl alcohol a complex, $MgI_2, 6CH_3OH$.

Solubility of $MgI_2, 6CH_3OH$ in CH_3OH at t° .

t°	% by weight of MgI_2, CH_3OH	t°	% by weight of $MgI_2, 6CH_3OH$
0	49.6	120	66.2
20	52.6	140	69.5
40	55.3	160	73.2
60	58.0	180	77.1
80	60.6	200	81.5
100	63.3		

(Menschutkin, Z. anorg. 1907, 52. 15.)

MgI_2 forms with ethyl alcohol a complex, $MgI_2, 6C_2H_5OH$.

Solubility of $MgI_2, 6C_2H_5OH$ in C_2H_5OH at t° .

t°	% by weight of $MgI_2, 6C_2H_5OH$	t°	% by weight of $MgI_2, 6C_2H_5OH$
0	21.9	120	82.7
20	33.2	130	87.2
40	44.4	135	90.0
60	55.3	140	93.3
80	65.5	143	96.0
100	74.7	145	98.0
110	78.8	146.5 mpt.	100

(Menschutkin.)

MgI_2 forms with dimethylcarbinol a complex, $MgI_2, 6(CH_3)_2CHOH$.

Solubility of $MgI_2, 6(CH_3)_2CHOH$ in $(CH_3)_2CHOH$ at t° .

t°	% by weight of $MgI_2, 6(CH_3)_2CHOH$	t°	% by weight of $MgI_2, 6(CH_3)_2CHOH$
10	57.1	110	76.2
30	60.0	120	79.4
50	63.3	130	84.8
70	67.0	136	91.7
90	71.2	138 mpt.	100

(Menschutkin.)

Solubility of MgI_2 in ether at t° .

t°	% MgI_2	% $MgI_2, 2C_4H_{10}O$
5.4	1.45	2.2
11.8	2.43	3.7
15.6	3.46	5.3
18.1	5.4	8.3
20.4	7.55	11.6
22.2	11.28	17.3

(Menschutkin, Z. anorg. 1906, 49. 41.)

t°	% MgI ₂	% MgI ₂ . 2C ₆ H ₁₀ O
	in lower layer	
14.8	35.5	54.4
17.6	35.5	54.4
20	35.8	54.8
28.4	35.5	54.4
33	35.7	54.7
35	35.3	54.1
in upper layer		
18.6	13.57	20.8
23.2	14.4	22.1
24.4	14.6	22.4
32.4	15.82	24.2
in solution when two layers mix		
37.3	19.4	29.3
38.5	22.45	34.4
38.5	26.07	39.9
38.5	29.8	45.7
38	32.8	50.3

(Menschutkin.)

Solubility of MgI₂ in acetic acid.
MgI₂ forms with acetic acid a complex, MgI₂. 6CH₃COOH.

Solubility of MgI₂. 6CH₃COOH in CH₃COOH at t°.

t°	% by wt. MgI ₂ . 6CH ₃ COOH
20	0.6
40	2.0
60	5.0
70	9.5
75	13.0
80	18.5
85	27.1
95	42.0
105	54.5
115	65.0
125	73.8
135	85.0
140	94.0
142 mpt.	100.0

(Menschutkin, Z. anorg. 1907, 54. 93.)

Solubility of MgI₂ in acetone.
MgI₂ forms with acetone a complex, 6CH₃COCH₃.

Solubility of MgI₂. 6CH₃COCH₃ in CH₃COCH₃ at t°.

t°	% by wt. MgI ₂ . 6CH ₃ COCH ₃
0	4.9
30	6.7
50	8.3
60	10.2
70	15.2
80	28.6
85	40.0
90	59.2
95	80.0
100	92.5
105	98.5
106.5 mpt.	100

(Menschutkin, Z. anorg. 1907, 53. 3)

Solubility of MgI₂ in acetal.
MgI₂ forms with acetal a complex, 2CH₃CH(OC₂H₅)₂.

Solubility of MgI₂. 2CH₃CH(OC₂H₅)₂ in CH₃CH(OC₂H₅)₂ at t°.

t°	% by wt. MgI ₂ . 2CH ₃ CH(OC ₂ H ₅) ₂
20	0.15
60	0.45
77	0.6
77	92.0
79	93.7
81	95.5
83	97.3
86 mpt.	100

(Menschutkin, Z. anorg. 1907, 53. 3)

Solubility of MgI₂ in acetamide.
MgI₂ forms with acetamide a complex, 6CH₃CONH₂.

Solubility of MgI₂. 6CH₃CONH₂ in CH₃CONH₂ at t°.

t°	% by wt. of MgI ₂ . 6CH ₃ CONH ₂
49	56.5
80	63.4
110	70.5
130	76.0
150	82.1
160	85.5
170	90.8
175	96.2
177 mpt.	100.0

(Menschutkin, Z. anorg. 1909, 61. 1)

Solubility of MgI_2 in acetonitrile.
 MgI_2 forms with acetonitrile a complex,
 $\text{I}_2, 6\text{CH}_3\text{CN}$.

Solubility of $\text{MgI}_2, 6\text{CH}_3\text{CN}$ in CH_3CN
at t° .

t°	% by wt. $\text{MgI}_2, 6\text{CH}_3\text{CN}$
0	37.2
30	49.8
50	58.2
70	67.9
80	76.5
89	91.3

(Menschutkin, Z. anorg. 1909, 61. 110.)

Solubility of MgI_2 in benzaldehyde.
 MgI_2 forms with benzaldehyde a complex,
 $\text{I}_2, 6\text{C}_6\text{H}_5\text{CHO}$.

Solubility of $\text{MgI}_2, 6\text{C}_6\text{H}_5\text{CHO}$ in $\text{C}_6\text{H}_5\text{CHO}$
at t° .

t°	% by wt. $\text{MgI}_2, 6\text{C}_6\text{H}_5\text{CHO}$
0	3.2
20	3.8
40	5.3
60	7.7
80	11.0
100	18.5
110	26.5
120	40.0
125	53.0
130	74.5
133	86.0
136	94.2
139 mpt.	100

(Menschutkin, Z. anorg. 1907, 53. 28.)

Solubility of MgI_2 in methyl acetate.
 MgI_2 forms with methyl acetate a complex,
 $\text{I}_2, 6\text{CH}_3\text{COOCH}_3$.

Solubility of $\text{MgI}_2, 6\text{CH}_3\text{COOCH}_3$ in
 $\text{CH}_3\text{COOCH}_3$ at t° .

t°	% by wt. $\text{MgI}_2, 6\text{CH}_3\text{COOCH}_3$
0	0.4
30	0.55
60	0.75
90	0.9
100	1.8
103	2.4
103	74.2
110	81.7
120	98.0
121 mpt.	100.0

(Menschutkin, Z. anorg. 1909, 61. 101.)

Solubility of MgI_2 in methyl acetate.
 MgI_2 forms with ethyl acetate a complex,
 $\text{MgI}_2, 6\text{CH}_3\text{COOC}_2\text{H}_5$.

Solubility of $\text{MgI}_2, 6\text{CH}_3\text{COOC}_2\text{H}_5$ in
 $\text{CH}_3\text{COOC}_2\text{H}_5$ at t° .

t°	% by wt. $\text{MgI}_2, 6\text{CH}_3\text{COOC}_2\text{H}_5$
0	3.2
20	4.8
40	8.6
50	13.7
55	21.5
60	38.0
65	63.5
70	90.5
75	97.7
78.5 mpt.	100.0

(Menschutkin.)

Solubility of MgI_2 in ethyl formate.
 MgI_2 forms with ethyl formate a complex,
 $\text{MgI}_2, 6\text{HCOOC}_2\text{H}_5$.

Solubility of $\text{MgI}_2, 6\text{HCOOC}_2\text{H}_5$ in
 HCOOC_2H_5 at t° .

t°	% by wt. $\text{MgI}_2, 6\text{HCOOC}_2\text{H}_5$
0	15.1
10	17.4
20	20.5
30	25
40	31.8
50	44
60	68
70.5 mpt.	100

(Menschutkin.)

Solubility of MgI_2 in isoamylacetate.
 MgI_2 forms with isoamylacetate a complex,
 $\text{MgI}_2, 6\text{CH}_3\text{COO}(\text{iso})\text{C}_5\text{H}_{11}$.

Solubility of $\text{MgI}_2, 6\text{CH}_3\text{COO}(\text{iso})\text{C}_5\text{H}_{11}$ in
 $\text{CH}_3\text{COO}(\text{iso})\text{C}_5\text{H}_{11}$ at t° .

t°	% by wt. $\text{MgI}_2, 6\text{CH}_3\text{COO}(\text{iso})\text{C}_5\text{H}_{11}$
0	7.7
20	11.5
40	20.9
45	25.5
50	33.2
55	47.8
57.5	63.0
60 mpt.	100.0

(Menschutkin.)

Solubility of MgI_2 in isobutyl acetate.
 MgI_2 forms with isobutylacetate a complex,
 $\text{MgI}_2, 6\text{CH}_3\text{COOC}_4\text{H}_9$.

Solubility of MgI_2 , $6\text{CH}_3\text{COO}(\text{iso})\text{C}_4\text{H}_9$ in $\text{CH}_3\text{COO}(\text{iso})\text{C}_4\text{H}_9$.

t°	% by wt. MgI_2 , $6\text{CH}_3\text{COO}(\text{iso})\text{C}_4\text{H}_9$
0	10.5
20	13.6
40	17.6
50	20.4
60	24.9
70	33.7
75	40.5
80	52.0
85	89.0
87.5 mpt.	100.0

(Menschutkin.)

Solubility of MgI_2 in propyl acetate.

MgI_2 forms with propyl acetate a complex, $\text{MgI}_2 \cdot 6\text{CH}_3\text{COOC}_2\text{H}_5$.

Solubility of $\text{MgI}_2 \cdot 6\text{CH}_3\text{COOC}_2\text{H}_5$ in $\text{CH}_3\text{COOC}_2\text{H}_5$ at t° .

t°	% by wt. MgI_2 , $6\text{CH}_3\text{COOC}_2\text{H}_5$
0	4.1
20	5.4
30	6.5
35	7.8
40	19.0
45	46.0
50	72.5
55	88.2
60	96.0
65 mpt.	100.0

(Menschutkin.)

Solubility of MgI_2 in urethane.

MgI_2 forms with urethane a complex, $\text{MgI}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$.

Solubility of $\text{MgI}_2 \cdot 6\text{NH}_2\text{COOC}_2\text{H}_5$ in $\text{NH}_2\text{COOC}_2\text{H}_5$ at t° .

t°	% by wt. MgI_2 , $6\text{NH}_2\text{COOC}_2\text{H}_5$
32	51.8
50	59.4
70	70.7
80	78.8
84	85.0
87 mpt.	100.0

(Menschutkin.)

+ $6\text{H}_2\text{O}$.

Solubility of $\text{MgI}_2 + 6\text{H}_2\text{O}$ in H_2O at t° .

t°	% by weight of $\text{MgI}_2 + 6\text{H}_2\text{O}$	t°	% by weight of $\text{MgI}_2 + 6\text{H}_2\text{O}$
43°	89.8	160°	91.7
80°	90.3	200°	93.4
120°	90.9	215°	94.3

(Menschutkin, Z. anorg. 1907, 52. 156.)

+ $8\text{H}_2\text{O}$. Sp. gr. of solution of MgI_2 sat. at 18° containing 59.7% MgI_2 . (Mylius, B. 1897, 30. 1718.)

Solubility of $\text{MgI}_2 + 8\text{H}_2\text{O}$ in H_2O :

t°	% by weight of MgI_2
0	76.0
20	81.0
40	88.0
43.5	90.8

(Menschutkin.)

+ $10\text{H}_2\text{O}$. Sol. in H_2O . (Panfiloff 1894, II. 610.)

Magnesium mercuric iodide, MgI_2 , H

Known only in solution.

+ $9\text{H}_2\text{O}$. Very deliquescent. (C. R. 1906, 142. 1338.)

Very sol. in ethyl, methyl, propyl isobutyl, amyl, isopropyl and allyl ; ethyl, amyl, propyl and isobutyl ; ethyl cyanide and acetone. Sol. in alcohol. Decomp. by glycerine. Sol. in ethyl benzoate, amyl benzoate, nitroethyl benzoate. Decomp. by ethyl oxalate. Insol. in benzene, ethyl iodide, CHCl_3 , CCl_4 , bromide, monochlor and monobromide (Duboin, A. ch. 1909, (8) 16. 276.)

$\text{MgI}_2 \cdot 2\text{HgI}_2$. Decomp. by H_2O in and above compound, which remains in solution. (Boullay.)

+ $7\text{H}_2\text{O}$. Sat. solution in H_2O at 1° the composition MgI_2 , 1.29 HgI_2 , 11. (Duboin, C. R. 1906, 142. 1338.)

Magnesium potassium iodide, MgI_2 , $6\text{H}_2\text{O}$.

Deliquescent. (Lerch, J. pr. (2) 2)

Very hygroscopic. (de Schulten, B. 1900 (3) 23. 158.)

Magnesium iodide ammonia, MgI_2 , 6

Practically insol. in liquid NH_3 . (F. J. Am. Chem. Soc. 1913, 35. 1459.)

Magnesium nitride, Mg_3N_2 .

Decomp. by moist air or H_2O . Sol. in conc. $\text{HCl} + \text{Aq.}$ or $\text{HNO}_3 + \text{Aq.}$ warm H_2SO_4 . Insol. in alcohol, ethyl or phosphorus oxychloride. (Briegleb, Geuther, A. 123. 236.)

Decomp. by H_2O . (Smits, R. t. 12. 198.)

Easily decomp. H_2O when finely powdered. (Rossel, C. R. 1895, 121. 942.)

Magnesium suboxide (?)

Decomp. H_2O . Sol. in dil. acids. (Pogg. 127. 45.)

Magnesium oxide, MgO .

Sol. in 50,000-100,000 pts. H_2O (Bineau, C. R. 510); in 55,368 pts. cold or hot H_2O (Fresenius, 123); in 100,000-200,000 pts. H_2O (Bunsen); in 7900 pts. H_2O at ord. temp. (Dalton); in 7900 pts. H_2O at ord. temp. (Kirwan); in 5760 pts. H_2O at 1° (Fyfe).

lc. from electrical conductivity of MgO
1 pt. MgO is sol. in 172,000 pts. H₂O
(Dupré, Zeit. angew. Ch. 1903, 16. 55.)
"heavy" MgO is more sol. in H₂O than
"light" MgO. The temp. of preparation
affects the rate of solution, the rate being
increased as the temp. of preparation is
increased. (Anderson, Chem. Soc. 1905, 87.)

Slightly sol. in acids, even in H₂SO₄+Aq.

Solubility in P₂O₅+Aq at 25°.

Composition of the solution		Sp. gr. 25°/25°	Solid phase
MgO g/l.	G. P ₂ O ₅ per l.		
0.07	0.486	...	MgHPO ₄ · 3H ₂ O
0.80	0.732	...	
1.53	1.917	...	
1.38	4.85	...	
1.3	7.35	1.006	
1.3	16.84	1.017	
1.9	38.59	1.042	
1.3	61.21	1.069	
1.9	93.09	1.109	
1.0	130.7	1.144	
1.1	281.8	1.285	
1.1	439.0	...	
1.1	498.4	1.470	
1.1	546.5	...	
1.1	584.0	...	
1.1	623.3	1.595	MgH ₂ (PO ₄) ₂ · xH ₂ O
1.1	625.9	...	
1.1	645.8	...	
1.1	680.7	...	
1.1	779.6	1.626	
1.1	809.6	1.644	
1.1	835.1	1.654	

meron, J. phys. Chem. 1907, 11. 364.)
sol. in NH₄ salts, NaCl, or KCl+Aq.
(minius.)

Solubility in MgCl₂+Aq at 25°.

% MgCl ₂	% MgO as Mg(OH) ₂
2.36	0.00008
4.47	0.00028
6.79	0.00048
9.02	0.00080
13.14	0.00115
15.15	0.00195
17.53	0.00240
18.52	0.00250
22.04	0.00245
23.78	0.00235
25.13	0.00230
26.88	0.00250
28.34	0.00230
29.80	0.00240
30.04	0.00250
34.22	0.0030

Robinson, J. phys. Chem. 1909, 13. 676.)

More sol. in K₂SO₄, and Na₂SO₄+Aq than
in H₂O. (Warrington.)
Insol. in liquid NH₃. (Franklin, Am. Ch.
J. 1898, 20. 828.)
Sol. in methyl alcohol to form a colloidal
solution containing 1.6% MgO. (Neuberg
and Rewald. (Biochem. Z. 1908, 9. 547.)
Insol. in methyl acetate (Naumann, B.
1909, 42. 3790); ethyl acetate. (Naumann,
B. 1904, 37. 3602.)
Insol. in acetone. (Naumann, B. 1904, 37.
4329.)

Insol. in acetone and in methylal. (Eid-
mann, C. C. 1899, II. 1014.)
Solubility in (calcium succinate+sugar)+
Aq.
1 l. solution containing 418.6 g. sugar and
34.3 g. CaO dissolves 0.30 g. MgO; contain-
ing 296.5 g. sugar and 24.2 g. CaO dissolves
0.24 g. MgO; containing 174.4 g. sugar and
14.1 g. CaO dissolves 0.22 g. MgO. (Boden-
bender, J. B. 1865. 600.)
See also Magnesium hydroxide.
Min. Periclase.

Magnesium peroxide, MgO₂.

Sol. in 14,550 pts. H₂O at 20°. (Foregger
and Philipp, J. Soc. Chem. Ind. 1906, 25.
298.)
5MgO, 2MgO₂+3H₂O.
3MgO, 2MgO₂+3H₂O.
2MgO, 2MgO₂+3H₂O.
4MgO, 2MgO₂+3H₂O.
Above salts are decomp. by H₂O.
(Carrasco, Gazz. ch. it. 1909, 39, (1) 47.)

Magnesium oxybromide, MgBr₂, 3MgO+
12H₂O.

Decomp. in the air and also by H₂O, al-
cohol and most reagents. (Tassilly, C. R.
1897, 125. 607.)

Magnesium oxychloride, Mg₂OCl₂+16H₂O.

Easily decomp. by H₂O and alcohol.
(André, A. ch. (6) 3. 80.)
+6H₂O. (André.)
2MgO, HCl, 5H₂O or 3MgO, MgCl₂+
10H₂O. Solubility determinations show that
this salt is the solid phase in equilibrium at
25° with solutions of MgCl₂ and MgO con-
taining from 10-15% MgCl₂. (Robinson, J.
phys. Chem. 1909, 13. 677.)
Mg₂O₃Cl₂+6, 8, 14, or 17H₂O. Decomp.
by H₂O, which dissolves out MgCl₂. (Ben-
der, B. 3. 932.)
Mg₁₁O₁₀Cl₂+14, or 18H₂O. (Krause, A.
165. 38.)
Mg₁₀O₉Cl₂+24H₂O=9MgO, MgCl₂+
24H₂O. H₂O removes all MgCl₂ by long di-
gesting. (Bender, A. 159. 341.)
+10, and 15H₂O. (Bender.)

Magnesium oxysulphide, Mg₂OS.

(Reichel, J. pr. (2) 12. 55.)

Magnesium phosphide, Mg_3P_2 .

Decomp. by H_2O , dil. $HCl + Aq$, or $HNO_3 + Aq$. (Parkinson, Chem. Soc. 5. (2) 125 and 309.)

Insol. in moderately dil. cold $HCl + Aq$, or boiling dil. $H_2SO_4 + Aq$. Difficultly and slowly sol. in aqua regia. (Blunt, Chem. Soc. 3. (2) 106.)

Decomp. by H_2O , HCl , conc. H_2SO_4 , and by HNO_3 . (Gautier, C. R. 1899, 128. 1169.)

Magnesium silicide, Mg_2Si .

Slowly decomp. by warm H_2O . Slowly decomp. by cold, rapidly by hot $NH_4Cl + Aq$. Decomp. by cold dil. $HCl + Aq$. (Geuther, J. pr. 95. 425.)

Mg_2Si . Decomp. by $HCl + Aq$ with residue of Si . (Wöhler, A. 107. 113.)

Slowly decomp. by H_2O at ord. temp. Violently decomp. by HCl . (Lebeau and Bossuet, C. R. 1908, 146. 284.)

Magnesium sulphide, MgS .

Decomp. by H_2O . (Reichel, J. pr. (2) 12. 55.)

Sl. sol. in H_2O with rapid decomp. (Freymy.)

Sol. in acids with decomp.

Anhydrous. Crystalline. Only very sl. sol. in cold H_2O . Sol. in HNO_3 and H_2SO_4 at ord. temp. Sol. in PCl_5 and in chromyl chloride. (Mourlot, C. R. 1898, 127. 182.)

Magnesium polysulphide, MgS_x .

Known only in solution. (Reichel.)

Magnus' green salt.

See Platodiamine chloroplatinite.

Manganese, Mn .

Decomposes H_2O even in the cold, more rapidly when hot. (Regnault.)

Decomposes cold water violently. (Bunsen.)

Sol. in all dil. acids. Slowly sol. in cold H_2SO_4 . (John.)

Insol. in cold, but rapidly sol. in hot H_2SO_4 . Very easily sol. in dil. H_2SO_4 , or $HCl + Aq$, HNO_3 , or $HC_2H_3O_2 + Aq$. (Brunner.)

Pure manganese is unaltered in dry air, even when finely powdered. Slowly attacked by cold, quickly by hot H_2O . Very sl. attacked by cold H_2SO_4 , rapidly on warming; rapidly attacked by cold dil. $H_2SO_4 + Aq$; violently by conc. $HNO_3 + Aq$; and rapidly by dil. HNO_3 , HCl , $HC_2H_3O_2 + Aq$, and also $NaOH + Aq$. Sol. in $NH_4Cl + Aq$. (Prelinger, W. A. B. 102, 2b. 359.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 828.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0276 g. Mn in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Manganese antimonide, $MnSb$.

Sol. in hot aqua regia. (Wedekind, 1907, 40. 1266.)

Manganese azoimide, basic, $Mn(OH)N_2$.

Only sl. sol. in H_2O with decomp. (Cattus, J. pr. 1898, (2) 58. 293.)

Manganese bismuthide, $MnBi$.

Very sensitive towards acids with exception of conc. HCl . (Wedekind, 1911, 44. 2665.)

Manganese boride, MnB .

Attacked by cold H_2O and by acids. (Soneix, C. R. 1904, 139. 1210.)

Easily attacked by HCl , H_2SO_4 , and with evolution of BH_3 . (Wedekind, B. 1 38. 1231.)

MnB_2 . Sol. in acids, with evolution of (Troost and Hautefeuille, A. ch. (5) 9. 61)

Slowly decomp. by H_2O . Sol. in dil. and other dil. acids with evolution of H_2 (Wedekind, B. 1905, 38. 1229.)

Manganous bromide, $MnBr_2$.

Anhydrous. Very deliquescent.

Sat. $MnBr_2 + Aq$ contains at:

—21°	+7°	11°	18°	38°	52°
52.1	56.5	57.0	59.1	62.7	64.2% Mn

64°	76°	89°	97°	105°
68.2	70.1	69.7	69.2	70.2% $MnBr_2$

(Étard, A. ch. 1894, (7) 2. 541.)

Insol. in liquid NH_3 . (Franklin, Am. J. 1898, 20. 828.)

+ H_2O . (Lescœur, A. ch. 1894, (7) 2. 1)

+4 H_2O . More deliquescent than Mn

Melts in crystal water when heated. (Barnett.)

+6 H_2O . (Kusnetsoff, C. C. 1897, II. 1)

Manganous mercuric bromide.

Deliquescent.

Manganous palladium bromide.

See Bromopalladite, manganous.

Manganous stannic bromide.

See Bromostannate, manganous.

Manganese carbide, MnC .

(Brown, J. pr. 17. 492.)

MnC_2 .

MnC_3 . (Troost and Hautefeuille, A. (5) 9. 60.)

Decomp. by H_2O and by dil. acids. (Barnett, C. R. 1896, 122. 422.)

Manganous chloride, $MnCl_2$.

Anhydrous. Deliquescent.

pts. H ₂ O at t° dissolve pts. MnCl ₂ :			
	Pts. MnCl ₂	t°	Pts. MnCl ₂
25 5	62.16	87.5	122.22
	85.72	106.25	123.81
	122.22

r, sat. MnCl ₂ +Aq at t° contains:			
	% MnCl ₂	t°	% MnCl ₂
25 5	38.33	87.5	55.0
	46.15	106.25	55.32
	55.0

(Brandes, Pogg. 22. 263.)

also below under +2H₂O, and +4H₂O.

of MnCl₂+Aq at 15°. a=sp. gr. if %
MnCl₂; b=sp. gr. if % is MnCl₂+
H₂O.

a	b	%	a	b
1.045	1.0285	40	1.443	1.250
1.091	1.057	45	1.514	1.290
1.138	1.086	50	...	1.331
1.189	1.116	55	...	1.375
1.245	1.147	60	...	1.419
1.306	1.180	65	...	1.463
1.372	1.214	70	...	1.508

(Gerlach, Z. anal. 28. 476.)

Sp. gr. of MnCl ₂ +Aq at room temp.	
% MnCl ₂	Sp. gr.
8.007	1.0960
15.650	1.1963
30.330	1.3372
40.132	1.4530

(Wagner, W. Ann. 1883, 18. 273.)		
Sp. gr. of MnCl ₂ +Aq at t°.		
t°	% MnCl ₂	Sp. gr.
14.5	5.0	1.0457
14.5	11.99	1.1076
14.0	14.98	1.1379
14.5	19.92	1.1891
14.0	23.10	1.2246
14.6	28.51	1.2888

(Long, W. Ann. 1880, 11. 38.)	
Sp. gr. of MnCl ₂ +Aq at 25°.	
Concentration of MnCl ₂ +Aq	Sp. gr.
1-normal	1.0513
1/2- "	1.0259
1/4- "	1.0125
1/8- "	1.0063

(Wagner, Z. phys. Ch. 1890, 5. 38.)

Solubility of MnCl₂+KCl in H₂O at t°.

	% MnCl ₂	% KCl	Solid phase
	40.23	...	MnCl ₂ , 4H ₂ O
	35.94	9.41	MnCl ₂ , 4H ₂ O+MnCl ₂ , KCl, 2H ₂ O+KCl
	...	23.06	KCl
4	44.46	...	MnCl ₂ , 4H ₂ O
	43.28	8.66	MnCl ₂ , 4H ₂ O+MnCl ₂ , KCl, 2H ₂ O
	38.65	13.79	MnCl ₂ , 4H ₂ O+MnCl ₂ , 2KCl, 2H ₂ O+KCl
	...	26.91	KCl
8	50.14	6.01	MnCl ₂ , 4H ₂ O+MnCl ₂ , 2H ₂ O+MnCl ₂ , KCl, 2H ₂ O
6	51.86	...	MnCl ₂ , 2H ₂ O
	49.95	6.67	MnCl ₂ , 2H ₂ O+MnCl ₂ , KCl, 2H ₂ O
	44.05	12.49	MnCl ₂ , KCl, 2H ₂ O+MnCl ₂ , 2KCl, 2H ₂ O
	36.85	18.77	MnCl ₂ , 2KCl, 2H ₂ O+MnCl ₂ , 4KCl
	...	31.57	KCl

(Süss, Z. Kryst. Min. 1912, 51. 262.)

l. in liquid NH₃. (Franklin, Am. Ch. 3, 20. 828.)

Solutions of MnCl ₂ in 75% alcohol saturated at t° contain:			
t°	% MnCl ₂	t°	% MnCl ₂
10	23.1	43.75	37.5
25	36.1	87.5	32.2
		(B.-pt.)	

Solutions of MnCl_2 in absolute alcohol saturated at t° contain:

t°	% MnCl_2	t°	% MnCl_2
11.25	33.3	76.25	36.2
37.5	33.3	(B.-pt.)	

(Brandes, *l. c.*)

MnCl_2 crystallises from above solutions on standing.

When 15–20 vols. ether are added to 1 vol. absolute alcohol sat. with MnCl_2 , MnCl_2 is completely pptd. (Döbereiner.)

Insol. in oil of turpentine.

Sol. in urethane. (Castoro, *Z. anorg.* 1899, 20. 61.)

Sl. sol. in benzonitrile. (Naumann, *B.* 1914, 47. 1369.)

Difficultly sol. in methyl acetate. (Naumann, *B.* 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, *B.* 1910, 43. 314.)

+ H_2O . Solubility in $\text{HCl} + \text{Aq}$ decreases with increasing amt. of HCl . It is greater when hot than cold, but is not inconsiderable even when HCl is conc. 1 l. conc. $\text{HCl} + \text{Aq}$ sat. at 12° dissolves 190 g. MnCl_2 from $\text{MnCl}_2 + \text{H}_2\text{O}$. (Ditte, *C. R.* 1881, 92. 243.)

+ $\frac{1}{2}\text{H}_2\text{O}$. $\text{MnCl}_2 + 4\text{H}_2\text{O}$ effloresces to $\text{MnCl}_2 + \frac{1}{2}\text{H}_2\text{O}$ in a dry atmosphere and under low pressure and not to $\text{MnCl}_2 + 2\text{H}_2\text{O}$. (Sabatier, *Bull. Soc.* 1894, (3) 11. 547.)

+ $2\text{H}_2\text{O}$.

Solubility in H_2O at t° .

t°	Pts. MnCl_2 per 100 pts. H_2O	Sp. gr. of sat. solution
60	108.6	1.6108
70	110.6	1.6134
80	112.7

(Dawson and Williams, *Z. phys. Ch.* 1899, 31. 63.)

Sat. aqueous solution of $\text{MnCl}_2 + 2\text{H}_2\text{O}$. Contains 51.86% MnCl_2 at 62.6° . (Süss, *Z. Krist.* 1912, 51. 262.)

+ $4\text{H}_2\text{O}$. Deliquescent.

100 pts. H_2O at t° dissolve:

t°	Pts. $\text{MnCl}_2 + 4\text{H}_2\text{O}$	t°	Pts. $\text{MnCl}_2 + 4\text{H}_2\text{O}$
8	151	87.5	641
31.25	265	106.25	656
62.5	641

(Brandes, *l. c.*)

Sol. in 0.8 pt. H_2O at 18.75° . (Abl.)

Pptd. from solution in 9.17 mols. H_2O . (Kuznetsoff, *C. C.* 1899, I. 246.)

Sat. aq. solution contains at:

— 22° — 5° + 7° 17° 19°
34.7 37.8 40.4 41.2 42.3% MnCl_2

35° 55° 57° 80° 100° 140°
44.4 48.2 50.0 51.0 53.7 54.7% MnCl_2

(Étard, *A. ch.* 1894, (7) 2. 537.)

Solubility in H_2O at t° .

t°	Pts. MnCl_2 per 100 pts. H_2O	Sp. gr. of solution
25	77.18	1.499
30	80.71	1.504
40	88.59	1.534
50	98.15	1.574
*57.65	105.40	1.609

* Temp. of transition into $\text{MnCl}_2 + \frac{1}{2}\text{H}_2\text{O}$. (Dawson and Williams, *Z. phys. Ch.* 1899, 31. 63.)

Sat. aqueous solution of MnCl_2 contains 40.23% MnCl_2 at 6° ; 44.6% at 28.4° . (Süss, *Z. Krist.* 1912, 51. 262.)

100 pts. 75% alcohol dissolve at t°

t°	Pts. $\text{MnCl}_2 + 4\text{H}_2\text{O}$	t°	Pts. $\text{MnCl}_2 + 4\text{H}_2\text{O}$
10	53	43.75	144
25	132	87.5	100

(Brandes, *l. c.*)

Insol. in absolute ether, which ~~ab~~ not abstract crystal H_2O .

Insol. in boiling oil of turpentine. (B.)

Sol. in conc. $\text{HNO}_3 + \text{Aq}$.

+ $5\text{H}_2\text{O}$. (Müller-Erbach, *B.* 1891, 3181.)

+ $6\text{H}_2\text{O}$. Pptd. from solution in 11. H_2O at -21° . (Kuznetsoff, *C. C.* 1907, 246.)

Manganese trichloride, MnCl_3 .

Immediately decomp. by H_2O ; sol. ether and in abs. alcohol. (Holmes, *Chem. Soc.* 1907, 29. 1285.)

Manganese tetrachloride, MnCl_4 .

Has not been isolated.

Sol. in H_2O , alcohol, or ether. (J. B. 1865. 225.)

Composition is Mn_2Cl_4 . (Christen pr. (2) 34. 41.)

Manganese hydrogen tetrachloride (manganic acid), $\text{MnCl}_4 \cdot 2\text{HCl}$.

Sol. in ether; decomp. by H_2O . (J. B. 1865. 225.)

Manganese heptachloride, $\text{MnCl}_7(\text{?})$.

Decomp. by H_2O . (Dumas, *Berz.* 1845, 112.)

e formula MnO_2Cl (?). (Aschoff, J.)

ous mercuric chloride, MnCl_2 , HgCl_2 , I_2O .

escent in moist air. Easily sol. in . Bonsdorff.)

, 2HgCl_2 . (Varet, C. R. 1896, 123,

ous potassium chloride, MnCl_2 , KCl I_2O .

escent. Very sol. in H_2O , but is thereby. (Remsen and Saunders, J. 14. 129.)

, $2\text{KCl}+2\text{H}_2\text{O}$. (Süss, Z. Kryst. 262.)

c potassium chloride, MnCl_2 , $2\text{KCl}+$

p. by H_2O . Sol. in HCl apparently decomp. (Rice, Chem. Soc. 1898,

, 2KCl . Very easily decomp. nd Best, Z. anorg. 1899, 22. 186.)

, MnCl_2 , 5KCl . Easily decomp. nd Best, Z. anorg. 1899, 22. 185.)

ous rubidium chloride, MnCl_2 , Cl .

froy.)

). Easily sol. in H_2O . Insol. in conc. $\text{HCl}+\text{ppt. anhydrous salt}$ ieous solution. (Godeffroy, Arch. 3) 12. 40.)

ns only $2\text{H}_2\text{O}$. (Saunders, Am. Ch.).)

ous thallic chloride, MnCl_2 , 2TiCl_3+).

e cryst. from H_2O . (Gewecke, A. . 224.)

ous stannic chloride.

lorostannate, manganous.

us chloride hydrazine, MnCl_2 , I_4 .

Franzen, Z. anorg. 1908, 60. 285.)

us chloride hydroxylamine, MnCl_2 , 2OH .

table; insol. in alcohol. (Feldt, B. 405.)

us fluoride, MnF_2 .

ol. in H_2O containing HF . (Ber-

in H_2O ; decomp. by boiling with ol. in liquid NH_3 ; easily sol. in cold nc. HNO_3 and HCl ; slowly sol in decomp. by fused K_2CO_3 , KOH , nd KClO_3 ; insol. in alcohol and

ether. Slowly sol. in acetic acid. (Moissan, C. R. 1900, 130. 1160.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 828.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Manganese trifluoride, MnF_3 .

Completely sol. in a little H_2O , but decomp. by dilution or boiling. (Berzelius.)

+ $6\text{H}_2\text{O}$. Efflorescent. (Christensen, J. pr. (2) 35. 57.)

Sol. in H_2SO_4 , HCl , HNO_3 ; decomp. by H_2O ; insol. in most organic solvents. (Moissan, C. R. 1900, 130. 626.)

Manganomanganic fluoride, $\text{Mn}_2\text{F}_8+10\text{H}_2\text{O}$.

Sol. in a little H_2O , but decomp. by dilution. (Nicklès, C. R. 67. 448.)

Manganese tetrafluoride, MnF_4 .

Not isolated. Sol. in absolute alcohol or ether; decomp. by H_2O . (Nicklès, C. R. 65. 107.)

Probably does not exist. (Christensen, J. pr. (2) 35. 161.)

Manganese heptafluoride, MnF_7 (?).

Sol. in H_2O with decomp. (Wöhler.)

Manganese sesquifluoride with MF .

See also Fluomanganate, M.

Manganic nickel fluoride, 2NiF_2 , Mn_2F_8+ $8\text{H}_2\text{O}$.

(Christensen, J. pr. (2) 34. 41.)

Manganic potassium fluoride, Mn_2F_8 , $4\text{KF}+$ $2\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in conc. $\text{HCl}+\text{Aq}$, dil. HNO_3+Aq , conc. $\text{H}_2\text{SO}_4+\text{Aq}$, $\text{H}_3\text{PO}_4+\text{Aq}$, $\text{H}_2\text{C}_2\text{O}_4+\text{Aq}$, $\text{H}_2\text{C}_4\text{H}_4\text{O}_6+\text{Aq}$, and dil. $\text{HF}+\text{Aq}$. (Christensen, J. pr. (2) 35. 72.)

MnF_4 , 2KF . Difficultly sol. in H_2O . Decomp. by much H_2O . (Nicklès, C. R. 65. 107.)

True composition is Mn_2F_8 , 4KF , also with $2\text{H}_2\text{O}$. (Christensen, J. pr. (2) 34. 41.)

MnF_4 , 4KF . (Nicklès.)

See also Fluomanganate, potassium.

Manganic rubidium fluoride.

See Fluomanganate, rubidium.

Manganic silver fluoride, 2AgF , Mn_2F_8+ $14\text{H}_2\text{O}$.

Sol. in $\text{HF}+\text{Aq}$. (Christensen, J. pr. (2) 34. 41.)

Manganic sodium fluoride, Mn_2F_8 , 4NaF .

Decomp. by much H_2O . Not as sol. in $\text{HF}+\text{Aq}$ as the K salt. (Christensen, J. pr. (2) 35. 161.)

Manganomanganic thalious fluoride, 5TlF , 2MnF_2 , MnF_2 .

Decomp. by H_2O .

Sl. sol. in dil., easily sol. in conc. HF .

Sol. in conc. HCl , dil. HNO_3 , and cold or hot conc. H_2SO_4 .

Sol. in warm H_2O , containing H_2SO_4 .

Sol. in dil. tartaric and oxalic acids. (Ephraim, B. 1909, 42. 4458.)

Manganous stannic fluoride.

See Fluostannate, manganous.

Manganic zinc fluoride, 2ZnF_2 , $\text{Mn}_2\text{F}_7 + 8\text{H}_2\text{O}$

See Fluomanganate, zinc.

Manganous zirconium fluoride.

See Fluozirconate, manganous.

Manganous fluoride ammonia, 3MnF_2 , 2NH_3 .

(Moissan, C. R. 1900, 130. 1161.)

Manganous hydroxide, MnO_2H_2 .

2.15×10^{-4} g.-mol. are sol. in 1 l. H_2O at 18° . (Sackur, Z. Elektrochem, 1909, 15. 846.)

Solubility in $\text{H}_2\text{O} = 0.6 \times 10^{-4}$ g. mol. (Herz, Z. anorg. 1899, 22. 284.)

1 l. H_2O dissolves 2×10^{-4} mol. MnO_2H_2 . (Tamm, Z. phys. Ch. 1910, 74. 500.)

Very sl. sol. in H_2O or alkalies. (Fresenius.) Easily sol. in acids. Insol. in NaOH , or $\text{KOH} + \text{Aq}$. Sol. in NH_4 salts + Aq . Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in $\text{NaOH} + \text{Aq}$ in presence of glycerine. (Donath, Dingl. 229. 542.)

Not pptd. by $\text{NH}_4\text{OH} + \text{Aq}$ in presence of $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$; by $\text{KOH} + \text{Aq}$ in presence of cane sugar; by $\text{KOH} + \text{Aq}$ in presence of Na citrate.

Solubility of MnO_2H_2 in organic Na salts + Aq . (0.5 normal.)

Na tartrate, 0.0068 mol. per l.

Na malate, 0.0042 " " "

Na citrate, 0.0126 " " "

(Tamm, Z. phys. Ch. 1910, 74. 496.)

Min. *Pyrochroite*.

Manganomanganic hydroxide, Mn_2O_4 , $x\text{H}_2\text{O}$.

Not attacked by boiling $\text{NH}_4\text{Cl} + \text{Aq}$. Behaves towards acids as Mn_2O_3 .

Manganic hydroxide, Mn_2O_3 , H_2O .

Insol. in hot or cold dil. $\text{H}_2\text{SO}_4 + \text{Aq}$.

Sol. in conc. H_2SO_4 at somewhat over 100° . (Carius.)

Sol. in tartaric, oxalic, and malic acids, with subsequent decomp. Insol. in formic, acetic, benzoic, or hippuric acids. (Hermann, Pogg. 74. 303.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$. Insol. in cane sugar + Aq . (Pescher.)

Min. *Manganite*. Sol. in conc. $\text{HCl} + \text{Aq}$. Sl. sol. in conc. H_2SO_4 .

Manganese dihydroxide, MnO_2 , H_2O .

See Manganous acid.

Manganous iodide, MnI_2 .

Anhydrous. Nearly insol. in (Walden, Z. anorg. 1902, 29. 374.)

Sol. in POCl_3 . (Walden, Z. anorg. 25. 212.)

Moderately sol. in liquid NH_3 . (Fr. Am. Ch. J. 1898, 20. 828.)

+ $4\text{H}_2\text{O}$. Very deliquescent, and H_2O . (Kuznetzoff, C. C. 1900, II. 52 + $9\text{H}_2\text{O}$. (Kuznetzoff.)

Manganous mercuric iodide, MnI_2 , $6\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. without in alcohol and acetone. (Dobroszerdo 1901, I. 363.)

3MnI_2 , 5HgI_2 , + $20\text{H}_2\text{O}$.

A sat. solution in H_2O at 17° has a ratio 1.4 MnI_2 , HgI_2 , + $10.22 \text{H}_2\text{O}$ and 2.98. (Duboin, C. R. 1906, 142. 133)

Very sol. without decomp. in methyl, isopropyl, isobutyl, and allyl ethyl acetate and ethyl cyanide. Less sol. in amyl, propyl and isobutyl acetates, acetone, acetic acid, formic acid, ppt. of HgI_2 , ethyl benzoate, ethyl butyl alcohol, amyl alcohol and nitrobenzene. Sl. decomp. by glycerine. Insol. in nitrate, ethylene bromide, toluene, CHCl_3 , CCl_4 , ethyl iodide, monobromobenzene, monochlorobenzene. (Duboin, A. ch. (8), 16. 278.)

Manganese nitride, Mn_3N_2 .

Sol. in HNO_3 only on heating. It dissolves only in presence of Pt. Aq dissolves slowly. H_2SO_4 acts only with conc. Insol. in acetic acid. (P. M. 1894, 15. 398.)

Mn_3N_2 . Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$ and NH_3 + Aq ; insol. in HCl ; sol. in HNO_3 , + Aq decomp. (Prelinger, M. 1894, 15. 398)

Mn_3N_2 . Easily attacked by acids and alkalies. (Wedekind, B. 1908, 41. 377)

Manganous oxide, MnO .

Insol. in H_2O . Easily sol. in acids. Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$.

Manganic oxide (Manganese sesquioxide, Mn_2O_3 .

Decomp. by boiling with HNO_3 , + MnO , which dissolves, and MnO_2 , insol. (Berthier); also by boiling with $\text{H}_2\text{SO}_4 + \text{Aq}$. (Turner.) Sol. in hot H_2SO_4 or $\text{HCl} + \text{Aq}$. Sol. in cold H_2O without decomp. If perfectly pure, in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, but if it contains MnO , it dissolves. (Rose.) Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$.

Insol. in acetone. (Naumann, B. 184329.)

lity in (calcium sucrate+sugar)+

lution containing 418.6 g. sugar and CaO dissolves 0.50 g. Mn_2O_3 ; containing 174.4 g. sugar and CaO dissolves 0.32 g. Mn_2O_3 . (Bodenf. B. 1865. 600.)

Braunite.

sol. Solution in H_2O containing 0.21 litre is precipitated by $KNO_3 + Aq$; $K_2SO_4 + Aq$ (1 : 1100); $(NH_4)_2SO_4$ (1 : 1500); $NaCl + Aq$ (1 : 1580); $MgSO_4$ (1 : 40,983); $BaCl_2 + Aq$ (1 : 58,823); $-Aq$ (1 : 147,929); $(NH_4)_2Al_2(SO_4)_4 + Aq$ (1 : 362,318); $K_2Cr_2(SO_4)_4 + Aq$ (1 : 416,350); $HCl + Aq$ (1 : 61,350); $HC_2H_3O_2$ (1 : 62,500). (Spring and de Bull. Soc. (2) 48. 170.)

manganic oxide, Mn_2O_3 .

in H_2O . Boiling dil. or conc. HNO_3 + gives out MnO (Berthier); also boiling $O_2 + Aq$. (Turner.) Sol. in hot HCl (Otto.) $NH_4Cl + Aq$ dissolves out (Rose.) Sol. without decomp. in hot conc. $H_3PO_4 + Aq$, and cold conc. H_2SO_4 , citric, and tartaric acids + Aq .

Hausmannite.

sesquioxide, MnO_2 .

Pyrolusite. Insol. in H_2O . Very sol. in conc. H_2SO_4 with evolution of O_2 in cold $HCl + Aq$; decomp. by hot q . Sol. in aqua regia. Sol. in $SO_2 + I_2O_5 + Aq$. (Karsten.)

insol. in HNO_3 , or dil. $H_2SO_4 + Aq$, except in presence of organic reducing substances. Sol. by citric acid, and more easily by $acid$. (Bolton.)

insol. in hot conc., but insol. in dil. HNO_3 (Deville.) When pure it is insol. in $H_2SO_4 + Aq$, but if a small quantity of $acid$ is added much MnO_2 dissolves.

decomp. by boiling $NH_4Cl + Aq$. Sol. in a mixture of nitrososulphuric acid and conc. $HCl + Aq$. (Bornträger, Rep. h. 1887. 741.)

sol. in acetone. (Naumann, B. 1904, 37. Schmidt, C. C. 1899, II. 1014.)

sesquioxides, Mn_2O_3 , Mn_3O_4 , etc.

Manganite, manganous.

Sesquioxide, MnO_2 .

luculent. Sol. in H_2O , with subsequent decomp. Decomp. by ether. Sol. in H_2SO_4 . (Franke, J. pr. (2) 36. 31.)

Sesquioxide, MnO_2 (?).

sol. in H_2O with decomp. Decomp. by ether. (Franke, J. pr. (2) 36. 166.)

Manganese heptoxide, Mn_2O_7 .

Very unstable; takes up H_2O from air. Sol. in H_2O with evolution of heat and rapid decomposition. Sol. in conc. H_2SO_4 without decomp. (Aschoff.)

Manganese oxychloride, $3Mn_2O_3, MnCl_2$.

Insol. in H_2O . (Saint-Gilles, C. R. 55. 329.) $MnCl_2, MnO$ (?). (Gorgeu, A. ch. (6) 4. 515.)

MnO_2Cl . See **Manganyl chloride.**

Manganic oxyfluoride, $MnOF_2$.

Sol. in absolute ether.

$MnOF_2, 2HF$ = fluoxymanganic acid. (Nicklès, C. R. 659. 107.)

Manganic oxyfluoride potassium fluoride.

See **Fluoxymanganate, potassium.**

Manganic sesquioxifluoride potassium fluoride.

See **Sesquifluoxymanganate, potassium.**

Manganous oxyiodide, $MnI_2, MnO + 6H_2O$.

Sol. in H_2O with decomp. (Kuznetzoff, C. C. 1913, I. 1659.)

Manganese oxysulphide, MnO, MnS .

Sol. in acids. (Arfvedson, Pogg. 1. 50.)

Manganese phosphide, Mn_3P_2 .

Insol. in dil. acids; sol. in hot conc. HNO_3 . (Wedekind, B. 1907, 40. 1268.)

Sol. in aqua regia; insol. in HNO_3 . (Granger, C. R. 1897, 124. 191.)

$Mn_3P_2, HCl + Aq$ dissolves out Mn_3P_2 and leaves Mn_7P_2 , which is sol. in $HNO_3 + Aq$. (Wöhler and Merkel, A. 86. 371.)

Not attacked by boiling H_2O or by HCl . Easily sol. in warm HNO_3 or aqua regia. (Wedekind and Veit, B. 1907, 40. 1268.)

xMn_3P_2, yMn_4P_3 . Easily sol. in aqua regia; partly sol. in H_2SO_4 or $HCl + Aq$. (Struve, J. pr. 79. 321.)

Mn_4P_3 . Insol. in $HCl + Aq$. Sol. in $HNO_3 + Aq$. (Schrötter, W. A. B. 1849, 1. 305.)

Manganous phosphoselenide, MnS, P_2Se .

Insol. in H_2O . Sol. in $HCl + Aq$ or $HNO_3 + Aq$. Insol. in cold, sl. decomp. by hot alkalies + Aq . (Hahn, J. pr. 93. 436.)

$2MnSe, P_2Se_3$. Insol. in cold, slowly sol. in hot $HCl + Aq$. Not decomp. by alkalies.

$2MnSe, P_2Se_3$. Easily decomp. by acids. (Hahn.)

Manganese selenide, MnSe.

Decomp. by H_2O and min. acids. (Wedekind, B. 1911, 44. 2667.)

Cryst. Sl. decomp. by H_2O at 100° ; easily sol. in dil. acids. (Fonzes-Diacon, C. R. 1900, 130. 1025.)

Manganese silicide.

Sol. in HF; only very sl. sol. in other acids. (Warren, C. N. 1898, 78. 319.)

Mg_2Si_2 . Sol. in $HCl + Aq$ with evolution of SiH_4 . (Wöhler, A. 106. 54.)

Mn_2Si . Insol. in H_2O . (Vigouroux, C. R. 1895, 121. 772.)

Easily sol. in HF. (Wedekind, B. 1911, 44. 2668.)

Easily sol. in dil. acids, HF and HNO_3 . Insol. in $KOH + Aq$. (Vigouroux, A. ch. 1897, (7) 12. 179.)

Easily sol. in HF when heated; in HCl when red hot. Sol. in dil. min. acids with decomp. (Vigouroux, C. R. 1895, 121. 772.)

Insol. in HNO_3 ; sol. in dil. or conc. HCl . Slowly decomp. by alkali hydroxides. (Lebeau, C. R. 1903, 136. 91.)

Mn_2Si . Easily sol. in molten alkali. (Vigouroux, C. R. 1895, 121. 772.)

$MnSi$. Slowly attacked by hot conc. HCl . Not acted upon by dil. or conc. HNO_3 or H_2SO_4 . (Lebeau, C. R. 1903, 136. 91.)

$MnSi_2$. Not attacked by HNO_3 or H_2SO_4 . Easily sol. in cold HF; decomp. by conc. alkalis + Aq . (Lebeau, C. R. 1903, 136. 233.)

Manganous sulphide, MnS .

Anhydrous. Insol. in H_2O . Sol. in weak acids, even in acetic acid.

1 l. H_2O dissolves 71.60×10^{-4} moles MnS at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Min. *Alabandite*. Sol. in $HCl + Aq$.

+ $\frac{1}{2}H_2O$. *Green.* Decomp. by boiling with H_2O . Sol. in weak acids, as acetic or sulphurous acid. Very sl. sol. in $(NH_4)_2S + Aq$. (Wackenroder.)

Sol. in NH_4 salts + Aq . 100 ccm. of sat. $NH_4Cl + Aq$ at 12° dissolve 0.43 g. MnS . (Clermont and Guyot, C. R. 85. 37.)

+ $\frac{3}{2}H_2O$. *Flesh-colored.* Less sol. in NH_4 salts, or acetic acid + Aq than the preceding salt. 100 ccm. of sat. $NH_4Cl + Aq$ at 12° dissolve 0.088 g. (Clermont and Guyot.)

Neither green nor flesh-coloured MnS contains H_2O . (Antony and Donnini, Gazz. ch. it. 23. 560.)

MnS is not pptd. in presence of alkali citrates, tartrates, or grape sugar; cane or milk sugar do not prevent precipitation. (Spiller.) Not pptd. in presence of $Na_4P_2O_7$. (Rose.)

Manganese sulphide, Mn_2S_4 .

Decomp. by H_2O . Sol. in cold d (Gautier and Hallopeau, C. R. 18 809.)

Manganese disulphide, MnS_2 .

(Senarmont, J. pr. 51. 385.)

Min. *Hauerite*. Decomp. by hot l with separation of S.

Manganous phosphorus sulphide, $Mn_2S_2P_2$.

Sol. in $HCl + Aq$ with decomp. (A. 46. 147.)

Manganous potassium sulphide, $3MnS \cdot K_2S$.

Nearly insol. in water, alcohol, Easily sol. in acids. (Völcker, A. 59

Manganous sodium sulphide, $3MnS \cdot Na_2S$.

Insol. in H_2O , alcohol, or ether dil. acids, and $SO_2 + Aq$. (Völcker.

$2MnS \cdot Na_2S$. Decomp. by H_2O . er, Pogg. 151. 446.)

Manganese telluride, $MnTe$.

Decomp. by H_2O and min. acids kind, B. 1911, 44. 2667.)

Manganic acid, H_2MnO_4 .

Known only in solution, which poses rapidly. (Franke, J. pr. (2)

Barium manganate, $BaMnO_4$.

Insol. in H_2O ; decomp. by acid (scherlich.)

Didymium manganate, $Di_2(MnO_4)_3$.

Insol. in H_2O . Sol. in H (Frerichs and Smith, A. 191. 331.)

Does not exist. (Cleve, B. 11. 9)

Lanthanum manganate, $La_2(MnO_4)_3$.

Ppt. (Frerichs and Smith, A. 191

Does not exist. (Cleve, B. 11. 9

Manganese manganate, $Mn_2O_7 \cdot 3MnO_2$.

$3MnO_2$.

See Manganese dioxide.

Lead manganate, $PbMnO_4 \cdot 2H_2O$.

Ppt. (Jolles, C. C. 1888. 58.)

Potassium manganate, K_2MnO_4 .

Sol. in water containing alkali decomp., but decomp. by pure H_2C recrystallised from dil. $KOH + Aq$.

Solubility in KOH + Aq at t°.

at	t°	Mol. K_2MnO_4 in 1 l. of sat. solution
KOH	0	0.907
	10	1.013
	20	1.140
	30	1.252
	45	1.424
KOH	0	0.554
	17	0.681
	25	0.733
	30	0.772
	40	0.852
	45	0.889
	51	0.938
	60	1.003
	70	1.074
	80	1.143
KOH	0	0.155
	15	0.224
	23	0.261
	30	0.303
	40	0.362
	45	0.388
	60	0.469
	70	0.528
	80	0.587
KOH	0	0.063
	10	0.070
	20	0.078
	30	0.096
	40	0.119
	50	0.142
	60	0.167
	70	0.196
	80	0.222
KOH	0	0.0145
	10	0.0152
	20	0.0160
	30	0.0215
	40	0.0305
	50	0.0462
	63	0.0620
	70	0.0700
	80	0.0830

r, Z. Elektrochem. 1912, 18. 724.)

ethyl acetate. (Naumann, B. 1910,

manganate permanganate,
 lnO_4 , $KMnO_4$.without decomp. in 20% KOH + Aq.
A. ch. (3) 61. 355.)manganate, $Na_2MnO_4 + 10H_2O$. H_2O , with partial decomp. (Gen-
r. 82. 58.)a manganate, $SrMnO_4$.in H_2O . (Frombers.)

Permanganic acid.

See Permanganic acid.

Manganicyanhydric acid, $H_2Mn(CN)_6$.

Not known in the free state.

Barium manganicyanide, $Ba_2[Mn(CN)_6]$.Sol. in H_2O . (Fittig and Eaton.)Barium potassium manganicyanide barium
cyanide, $2KBaMn(CN)_6$, $3Ba(CN)_2 +$
 $8H_2O$.Decomp. by H_2O . (Lehmann, Dissert.
1898.)Calcium manganicyanide, $Ca_2[Mn(CN)_6]$.Sol. in H_2O . (Fittig and Eaton.)Potassium manganicyanide, $K_2Mn(CN)_6$.Sol. in H_2O . (Christensen, J. pr. (2) 31.
163.)Sodium manganicyanide, $Na_2Mn(CN)_6 +$
 $2H_2O$.Sol. in H_2O . (Fittig and Eaton.)

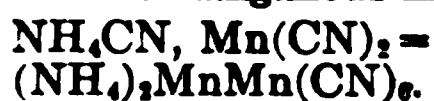
Manganimanganic acid.

Barium manganimanganate, $Ba_2Mn_2O_7 +$
 H_2O .Insol. in H_2O . Identical with Rosenstiehl's
"basic barium manganate," (J. Pharm. 1864,
46. 344). (Auger and Billy, C. R. 1904, 138.
501.)Lithium manganimanganate, $Li_2Mn_2O_7 +$
 H_2O .Insol. in H_2O . (Auger and Billy.)Manganiperiodic acid, H_2O , Mn_2O_7 , I_2O_7 .Wholly insol. in H_2O , in hot dil. or conc.
 HNO_3 , and in hot dil. H_2SO_4 . (Price, Am.
Ch. J. 1903, 30. 182.)Potassium manganiperiodate, K_2O , Mn_2O_7 ,
 I_2O_7 .Apparently entirely insol. and unchanged
when boiled with H_2O , dil. or conc. HNO_3 ,
or dil. H_2SO_4 . (Price.)Sodium manganiperiodate, Na_2O , Mn_2O_7 ,
 I_2O_7 .Apparently insol. and unchanged when
treated with boiling H_2O , boiling dil. or conc.
 HNO_3 , and boiling dil. H_2SO_4 . (Price.)Manganocyanhydric acid, $H_2Mn(CN)_6$.Most easily decomp. Sl. sol. in alcohol.
Insol. in ether. (Descamps, A. ch. (5) 24.
185.)

Ammonium cuprous manganocyanide,
 $(\text{NH}_4)_2\text{Cu}_2\text{Mn}(\text{CN})_6$.

Sol. in H_2O ; decomp. by acids and alkalies; very unstable. (Straus, Z. anorg. 1895, 9. 14.)

Ammonium manganous manganocyanide,



Sol. in $\text{NH}_4\text{CN} + \text{Aq.}$ (Fittig and Eaton, A. 145. 157.)

Barium manganocyanide, $\text{Ba}_2\text{Mn}(\text{CN})_6$.

Sol. in cold H_2O . (Fittig and Eaton.)

Calcium manganocyanide, $\text{Ca}_2\text{Mn}(\text{CN})_6$.

Very deliquescent. Sol. in H_2O ; insol. in alcohol. (Fittig and Eaton.)

Cuprous potassium manganocyanide,



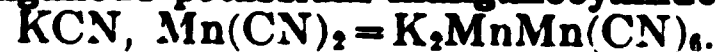
Sol. in H_2O with sl. decomp. Easily decomp. by acids and alkalies. (Straus, Z. anorg. 1895, 9. 12.)

Cuprous sodium manganocyanide,



Sol. in H_2O with only sl. decomp. Partially decomp. by acids. (Straus.)

Manganous potassium manganocyanide,



Ppt. Sol. in $\text{KCN} + \text{Aq.}$

Potassium manganocyanide, $\text{K}_4\text{Mn}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Very efflorescent. Sol. in H_2O ; decomp. by boiling.

Potassium manganocyanide chloride,



Easily sol. in H_2O . (Descamps.)

Sodium manganocyanide, $\text{Na}_4\text{Mn}(\text{CN})_6 + 8\text{H}_2\text{O}$.

Very efflorescent. Easily sol. in H_2O . (Fittig and Eaton.)

Strontium manganocyanide, $\text{Sr}_2\text{Mn}(\text{CN})_6$.

As the Ba comp. (Descamps.)

Permanganomolybdic acid.

See Permanganomolybdic acid.

Permanganotungstic acid.

See Permanganotungstic acid.

Manganosulphuric acid.

See Sulphate, manganic.

Manganous acid, $\text{H}_2\text{MnO}_3 = \text{MnO}_2, \text{H}_2\text{O}$.

Insol. in H_2O . (Franke, J. pr. (2) 36. 451.)
 $2\text{MnO}_2, \text{H}_2\text{O}$ (?). Min. Wad.

Barium manganite, $\text{BaO}, 5\text{MnO}_2$.

Sl. sol. in $\text{HCl} + \text{Aq}$, less sol. in HN (Rissler, Bull. Soc. (2) 30. 111.)

$\text{BaO}, 7\text{MnO}_2$. (Rousseau, C. R. 1

$\text{BaO}, 2\text{MnO}_2$. Insol. in H_2O .

BaO, MnO_2 . Insol. in H_2O . (F C. R. 102. 425.)

$\text{Ba}(\text{H}_2\text{Mn}_2\text{O}_{10})_2$. (Morawski and pr. (2) 18. 92.)

Calcium manganite, $\text{CaO}, 5\text{MnO}_2$.

Easily sol. in $\text{HCl} + \text{Aq}$, less in H (Rissler.)

$3\text{CaO}, \text{MnO}_2$ (?). Decomp. by H in $\text{HCl} + \text{Aq}$ with evolution of Cl . sol. in cold HNO_3 , but sol. on heating, A. ch. 1897, (7) 12. 275.)

$2\text{CaO}, \text{MnO}_2$. Sol. in dil. m (Rousseau, C. R. 116. 1060.)

$\text{CaO}, 2\text{MnO}_2$. (Rousseau, C. R.

$\text{CaO}, 3\text{MnO}_2$.

CaO, MnO_2 . Sol. in fuming HCl not in dil. $\text{HNO}_3 + \text{Aq}$. (Rousseau, 1060.)

Chromium manganite, $\text{Cr}_2\text{O}_3, \text{Cr}_2(\text{MnO}_3)_3$.

Slowly decomp. by acids. (C anorg. 1905, 44. 458.)

Cobaltous manganite, CoO, MnO_2 .

Ppt. (Salinger, Z. anorg. 1903, 1 + $4\text{H}_2\text{O}$. Ppt. (Salinger.)

Cobalt copper manganite, $\text{CoO}, \text{Cu} + 4\text{H}_2\text{O}$.

Min. Asbolite. Sol. in $\text{HCl} +$ evolution of Cl .

Cupric manganite, $\text{CuO}, 4\text{MnO}_2$.

(Gorgeu, Bull. Soc. 1903, (3) 29.

$\text{CuO}, 8\text{MnO}_2 + 3\text{H}_2\text{O}$. (Baubig 1897, 124. 955.)

Cupric manganous manganite, $4\text{Cu} + 7\text{MnO}_2 + 8\text{H}_2\text{O}$.

Ppt. (Salinger, Dissert. 1902.)

$\text{Mn}_2\text{O}_3, 3\text{CuO}$. Sol. in $\text{HCl} + \text{Aq}$. der, Am. Ch. J. 9. 269.)

Lead manganite, $\text{PbO}, 5\text{MnO}_2$.

Not attacked by conc. acids; so regia. (Rissler.)

Magnesium manganite, $2\text{MgO}, \text{Mn}$

(Lemoine, Ann. Min. (7) 3. 5.)

+ $x\text{H}_2\text{O}$. (Vollard.)

Manganous manganite, $\text{Mn}_2\text{O}_3 = 2\text{MnO}_2$.

(Reissig, A. 103. 27.)

$\text{Mn}_2\text{O}_{11} = \text{MnO}, 5\text{MnO}_2$. (Vel Soc. 38. 581.)

2MnO. Decomp. by dil. H₂SO₄ +
ke, J. pr. (2) 36. 166.)

MnO + H₂O. Min. *Varvicite*.

3 zinc manganite, MnO, ZnO,

Bull. Soc. 1903, (3) 29. 1168.)
nO, 2MnO₂. (Gorgeu.)

manganite, K₂O, 2MnO₂.

H₂O.

nO₂.

nO₂ + 3H₂O.

[nO₂ + 3H₂O = KH₂Mn₄O₁₀. (Mor-
stingl, J. pr. (2) 18. 91.)

t exist. (Wright and Menke,
37. 22.)

4nO₂.

1nO₂ + 6H₂O. Sol. in conc. HCl +
seau, C. R. 114. 72.)

manganite, AgH₂Mn₄O₁₀.

ki and Stingl, J. pr. (2) 18. 92.)

. Ppt. (Gorgeu, C. R. 110. 958.)

entous) manganite, Ag₄O,
(?).

cold dil. HNO₃ + Aq, and separ-
on warming. Insol. in NH₄OH +
Pogg. 101. 229.)

entoargentic) manganite, Ag₄O,
Mn₂O₃ (?).

nganite, Na₂O, 5MnO₂.

H₂O. (Rousseau, C. R. 103. 261.)

2MnO₂. Insol. in H₂O. (Rous-

(Rousseau, C. R. 112. 525.)

4nO₂ + 5H₂O. (Rousseau.)

MnO₂ + 8H₂O. (Rousseau.)

nanganite, MnO₂, SrO.

H₂O.

SrO. Insol. in H₂O. (Rousseau,
167.)

3rO. Sol. in HCl, or HNO₃ + Aq.
Ill. Soc. (2) 30. 110.)

nite, ZnO, 5MnO₂.

H₂O. (Rissler.)

[nO₂. (Gorgeu, Bull. Soc. 1903,
3.)

nO₂ + 7½H₂O. (Salinger, Dissert.

2MnO₂ + 25H₂O. Insol. in H₂O.

chloride, MnO₂Cl.

by H₂O. (Aschoff, J. pr. 81. 29.)

Melanocobaltic chloride,

Co₂(NH₃)₆Cl₄NH₂Cl, or

Co₂(NH₃)₆Cl₄NH₃.

Very sl. sol. in cold H₂O or very dil. HCl +
Aq. Decomp. by long standing or warming.
Cold conc. HCl or dil. H₂SO₄ + Aq does not
attack, but decomp. on warming. HNO₃ +
Aq decomp. on warming. Sol. in cold H₂SO₄
or NH₄OH + Aq; from both solutions it can
be precipitated by HCl + Aq. (Vortmann, B.
10. 1455.)

— chloroplatinate, Co₂(NH₃)₆NH₂Cl₄,
PtCl₄.

Ppt. (Vortmann, B. 15. 1902.)

Co₂(NH₃)₆NH₂Cl₄(OH)₂, PtCl₄. Ppt.
(Vortmann.)

— mercuric chloride,

Co₂(NH₃)₆(NH₂)Cl₄(OH)₂, 3HgCl₂ +
H₂O.

Ppt. Difficultly sol. in cold H₂O, quite
easily in warm H₂O acidified with HCl.
(Vortmann.)

— chloride chromate,

Co₂(NH₃)₆NH₂Cl₄Cr₂O₇ + H₂O.

Sol. in hot H₂O. (Vortmann.)

Mercuriammonium comps.

See Mercury ammonium comps.

Mercuriammonium bromide, Hg(NH₂)Br.

See Dimercuriammonium ammonium bro-
mide.

Mercuriammonium chloride, Hg(NH₂)Cl.

See Dimercuriammonium ammonium chlo-
ride.

**Mercuriammonium oxydimercuriammonium
chloride, 4Hg(NH₂)Cl, NH₂(HgOHg)Cl.**

(Millon.)

Correct composition is *Dimercuriammon-
ium ammonium chloride*, NHg₂Cl, NH₄Cl,
which see. (Balestra, Gazz. ch. it. 21, 2.
294.)

Hg(NH₂)Cl, 2NH₂(HgOHg)Cl. (Millon.)

Correct composition is *Dimercuriammon-
ium mercuric chloride*, 2NHg₂Cl, HgCl₂ +
H₂O, or *Dimercuriammonium hydrogen chlor-
ide*, NHg₂Cl, HCl. (Balestra.)

**Mercuriammonium nitrate, 2NH₃, 2HgO,
N₂O₅ = NH₂HgNO₃ + ½H₂O.**

Easily decomp. by HCl, or alkali sul-
phides + Aq. Sl. sol. in HNO + Aq. Insol.
in H₂SO₄, NH₄OH, or KOH + Aq. (Mits-
cherlich.)

Is *dimercuriammonium ammonium ni-
trate*, NHg₂NO₃, NH₄NO₃ + H₂O. (Pesci,
Gazz. ch. it. 20. 485.)

Mercuriammonium oxydimercuriammonium nitrate, 3HgO , 2NH_3 , $\text{N}_2\text{O}_5 = \text{NH}_4\text{HgNO}_3$, $(\text{NHg}_2\text{OH})\text{NO}_3 + \text{H}_2\text{O}$.

Decomp. by boiling with H_2O , which dissolves out NH_4NO_3 . Sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$ containing NH_4OH . (Mitscherlich.)

Is *dimercuriammonium ammonium nitrate*, $3\text{NHg}_2\text{NO}_3$, $\text{NH}_4\text{NO}_3 + 2\text{H}_2\text{O}$. (Pesci, Gazz. ch. it. 20. 485.)

Mercuriammonium oxydimercuriammonium sulphate, $(\text{NH}_4\text{Hg})_2\text{SO}_4$, $3(\text{NHg}_2\text{OH})_2\text{SO}_4$.

Boiling H_2O dissolves out H_2SO_4 . Gradually decomp. by boiling $\text{KOH} + \text{Aq}$. Completely sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. Sol. in conc. or dil. HCl , or very dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Insol. in conc. or dil. $\text{HNO}_3 + \text{Aq}$ or conc. H_2SO_4 . (Schneider.)

Correct formula is $7(\text{NHg}_2)_2\text{SO}_4$, $(\text{NH}_4)_2\text{SO}_4 + 12\text{H}_2\text{O}$, *dimercuriammonium ammonium sulphate*. (Pesci, Gazz. ch. it. 20. 485.)

Mercuridiammonium chloride (fusible white precipitate), $\text{Hg}(\text{NH}_2)_2\text{Cl}_2$.

Is *dimercuriammonium ammonium chloride*, Hg_2NCl , $3\text{NH}_4\text{Cl}$, which see. (Rammelsberg J. pr. 38. 558.)

Mercuridiammonium mercuric chloride, $\text{Hg}(\text{NH}_2)_2\text{Cl}_2$, HgCl_2 .

Insol. in H_2O , but gradually decomp. by boiling therewith. (Rose, Pogg. 20. 158.) Partly sol. in H_2O . (Kane.)

Mercuridiammonium iodide, $\text{Hg}(\text{NH}_2)_2\text{I}_2$.

H_2O extracts all the NH_3 . Partly sol. in little alcohol. Partly sol. in ether without decomp. (Nessler.)

Correct composition is *dimercuriammonium ammonium iodide*, NHg_2I , $3\text{NH}_4\text{I}$. (Pesci, Gazz. ch. it. 20. 485.)

Mercuridiammonium cupric iodide, 4NH_3 , CuI_2 , HgI_2 .

Decomp. by H_2O . Sol. in alcohol + $\text{HC}_2\text{H}_3\text{O}_2$. (Jørgensen, J. pr. (2) 2. 347.) $2\text{Hg}(\text{NH}_2)_2\text{I}_2$. CuI_2 . (Decomp. by H_2O . (Jørgensen.)

Mercuridiammonium iodide, $\text{Hg}(\text{NH}_2)_2\text{I}_2$.

Decomp. by H_2O . Partly sol. in a little alcohol. Partly sol. in ether. (Nessler.)

Correct composition is *dimercuriammonium ammonium iodide*, NHg_2I , $3\text{NH}_4\text{I}$. (Pesci.)

Mercuridiammonium mercuric iodide, $\text{Hg}(\text{NH}_2)_2$, HgI_2 , or NH_3 , HgI_2 .

Decomp. by H_2O or dil. acids. (Caillot and Corriol, J. Pharm. 9. 381.)

Correct composition is *dimercuriammonium ammonium mercuric iodide*, $3\text{NHg}_2\text{I}$, $8\text{NH}_4\text{I}$, 4HgI_2 . (Pesci, Gazz. ch. it. 20. 485.)

Mercuridiammonium sulphate, $\text{Hg}(\text{NH}_2)_2$, SO_4 .

Decomp. with H_2O . Does not exist. (Pesci, Gazz. ch. 485.)

+ H_2O . Decomp. by H_2O . Easily HCl , very dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, or HNO_3 . Insol. in conc. $\text{HNO}_3 + \text{Aq}$. Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$ or $\text{NH}_4\text{Cl} + \text{Aq}$. Decomp. by Aq . (Schneider, J. pr. 75. 136.)

Correct composition is $(\text{NHg}_2)_2\text{SO}_4 + 12\text{H}_2\text{O}$, *dimercuriammonium sulphate*. (Pesci.)

Dimercuriammonium acetate, $\text{NHg}_2\text{C}_2\text{H}_3\text{O}_2$.

Insol. in H_2O or alcohol. Sol. in $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Balestra, Gazz. 22. 2. 563.)

Dimercuriammonium ammonium, $\text{NHg}_2\text{C}_2\text{H}_3\text{O}_2$, $3\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{H}_2\text{O}$.

Deliquescent; sol. in a little H_2O decomp., but decomp. into $\text{NHg}_2\text{C}_2\text{H}_3\text{O}_2$ and $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$ by excess of H_2O . (Balestra, Gazz. 22. 2. 563.)

— **arsenate**, $\text{NHg}_2\text{H}_2\text{AsO}_4$.

(Hirzel, Zeit. Pharm. 1853. 3.)

— **bromate**, $\text{NHg}_2\text{BrO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Ppt. (Rammelsberg, Pogg. 55. 82) Is *oxydimercuriammonium bromate*, $(\text{NH}_2\text{Hg}_2\text{O})\text{BrO}_3$.

— **bromide**, NHg_2Br .

Insol. in H_2O or HNO_3 . Sol. in F (Pesci, Gazz. ch. it. 19. 509.)

Sol. in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ with excess of NH_3 . (Balestra, Gazz. ch. it. 22. 2. 563.)

Sol. in ammoniacal solutions of anhydrous salts and in aq. acids. (Franklin, Chem. Soc. 1905, 27. 839.)

— **ammonium bromide**, NHg_2Br .

Decomp. by H_2O . (Pesci, Gazz. ch. it. 19. 509.)

$4\text{NHg}_2\text{Br}$, $5\text{NH}_4\text{Br}$. Decomp. by H_2O . Insol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Sol. in dil. $\text{HCl} + \text{Aq}$. Insol. in $\text{HNO}_3 + \text{Aq}$.

NHg_2Br , $3\text{NH}_4\text{Br}$. Decomp. by H_2O . Easily sol. in $\text{HCl} + \text{Aq}$. Insol. in $\text{HNO}_3 + \text{Aq}$. (Pesci.)

Sol. in NH_4Br , NH_4Cl , or NH_4I sol. in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$.

— **mercuric bromide**, $2\text{NHg}_2\text{Br}$, I .

Ppt. Sol. in HBr and in HCl (Franklin, Chem. Soc. 1902, 31. 649.)

— **carbonate**, $(\text{NHg}_2)_2\text{CO}_3 + 2\text{H}_2\text{O}$.

Ppt. Not decomp. by KOH or easily by K_2S , or $\text{KI} + \text{Aq}$. (Rammelsberg, J. pr. (2) 38. 567.)

curiammonium chloride, NHg_2Cl .

attacked by boiling H_2O . Sl. attacked by dil. $\text{HCl} + \text{Aq}$, but is gradually destroyed thereby. Decomp. by hot $\text{KOH} + \text{Aq}$.

) in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ with evolution of S .

O_2 . Nearly insol. in H_2O ; easily sol. in O_2 , and $\text{HCl} + \text{Aq}$. Not decomp. by $-\text{Aq}$. Decomp. by KCl , NaCl , or $\text{KI} + \text{Rammelsberg}$, Pogg. 48. 181.)

hydrogen chloride, NHg_2Cl , 2HCl .

Correct composition of mercuric chlorammonium chloride. (Balestra, Gazz. ch. it. 21, 1901, 17. 96.)

Decomp. by H_2O .

HCl . Decomp. by H_2O . (Balestra, l. c.)

HCl , 4HCl . Sol. in H_2O . (Ray, Proc. Soc. 1901, 17. 96.)

ammonium chloride, NHg_2Cl , NH_4Cl . (Balestra, l. c.)

Correct composition of what has been called mercuric chloramide, $\text{Hg}(\text{NH}_2)\text{Cl}$. (Rammelsberg, J. pr. 38. 558.)

l. in cold, decomp. by hot H_2O . (Mikolajewski, ch. (3) 18. 413.) Sol. in 600 pts. H_2O . (Wittstein.) Sol. in 719.98 pts. H_2O . (Abl.) Insol. in alcohol.

in acids, even in $\text{HCl} + \text{Aq}$, also in HNO_3 , $(\text{NH}_4)_2\text{SO}_4$, and $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Pelouze and Fremy.)

in warm NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq}$.

sol. in alkali chlorides + Aq , which partly decomp. (Miahle, A. ch. (3) 5. 180.)

Decomp. by $\text{KOH} + \text{Aq}$. Sol. in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$, with evolution of NH_3 . (Balestra, l. c.)

a freshly prepared is sol. in conc. $\text{HCl} + \text{Aq}$. (Saha and Choudhuri, Z. 1910, 67. 359.)

in excess of $(\text{NH}_4)_2\text{HPO}_4 + \text{Aq}$. Insol. in excess of $\text{Na}_2\text{HPO}_4 + \text{Aq}$. (Carnegie and Z. N. 1897, 76. 175.)

l. in excess of $\text{NH}_4\text{OH} + \text{Aq}$. (Carnegie and Burt.)

HCl , $3\text{NH}_4\text{Cl}$ (Fusible white precipitate)

Correct composition of what has been called mercuridiammonium chloride, $(\text{NH}_4)_2\text{HgCl}_2$. (Rammelsberg, J. pr. (2) 38. 566.)

Decomp. by hot H_2O . Sol. in acids, even in $\text{HCl} + \text{Aq}$. Not decomp. by cold, but by boiling $\text{KOH} + \text{Aq}$. (Weyl.)

in warm, less in cold $\text{NH}_4\text{OH} + \text{Aq}$. (Mitscherlich.)

in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$, with evolution of S . (Balestra.)

in 10% HNO_3 , H_2SO_4 , and acetic acid. (Hofmann and Marburg, A. 1899, 305. 212.)

Dimercuriammonium mercuric chloride, $2\text{NHg}_2\text{Cl}$, HgCl_2 .

Insol. in, and not decomp. by boiling H_2O , alkalies, conc. HNO_3 , or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in boiling $\text{HCl} + \text{Aq}$. (Mitscherlich, J. pr. 19. 453.)

Sl. decomp. by H_2O , readily by $\text{KOH} + \text{Aq}$. (Gaudechon, A. ch. 1911, (8) 22. 212.)

Ppt. Sol. in HBr . (Ray, Proc. Chem. Soc. 1902, 18. 86.)

— chloride ammonia, NHg_2Cl , $\frac{1}{2}\text{NH}_3$.

Decomp. by water and by $\text{NH}_4\text{OH} + \text{Aq}$. (Gaudechon, A. ch. 1911, (8) 22. 212.)

— chromate.

See Oxydimercuriammonium chromate.

— hydroxide, NHg_2OH .

Takes up H_2O to form $\text{NHg}_2\text{OH} + \text{H}_2\text{O}$ or $(\text{NHg}_2\text{OH})_2\text{O}$, oxydimercuriammonium hydroxide, which also see.

Sol. in warm HCl or $\text{HNO}_3 + \text{Aq}$.

— iodate, NHg_2IO_3 , $2\text{NH}_4\text{IO}_3$.

Insol. in HNO_3 . (Rammelsberg, J. pr. (2) 38. 568.)

— iodide, NHg_2I .

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. Decomp. by boiling with $\text{KOH} + \text{Aq}$ or $\text{KCl} + \text{Aq}$. (Weyl, Pogg. 121. 601.) Decomp. by hot KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Balestra.)

Decomp. by dilute HCl . Sol. in ammonium salts + Aq . (Franklin, Z. anorg. 1905, 46. 21.)

+ H_2O . See Oxydimercuriammonium iodide.

— ammonium iodide, NHg_2I , $3\text{NH}_4\text{I}$.

Correct composition of mercuridiammonium iodide, $\text{Hg}(\text{NH}_2)_2\text{I}_2$. (Pesci, Gazz. ch. it. 20. 485.)

$3\text{NHg}_2\text{I}$, $8\text{NH}_4\text{I}$, 4HgI_2 . Correct formula for mercuridiammonium mercuric iodide, $\text{Hg}(\text{NH}_2)_2\text{I}_2$, HgI_2 . (Pesci.)

— nitrate, NHg_2NO_3 .

Insol. in H_2O . (Rammelsberg, J. pr. (2) 38. 566.)

Sol. in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$, with evolution of NH_3 . (Balestra, Gazz. ch. it. 22. 2. 560.)

+ H_2O . (Hofmann and Marburg, A. 1899, 305. 212.)

Sl. sol. in HNO_3 . (Ray, Z. anorg. 1902, 33. 209.)

— ammonium nitrate, NHg_2NO_3 , NH_4NO_3 , + H_2O .

Correct formula for mercuriammonium nitrate, $\text{NH}_2\text{HgNO}_3 + \frac{1}{2}\text{H}_2\text{O}$. (Pesci, Gazz. ch. it. 20. 485.)

NHg_2NO_3 , $2\text{NH}_4\text{NO}_3 + 2\text{H}_2\text{O}$. Correct for-

mula for oxydimercuriammonium ammonium nitrate, $(\text{NHg}_2\text{OH}_2)\text{NO}_3$, $2\text{NH}_4\text{NO}_3 + \text{H}_2\text{O}$. (Pesci.)

NHg_2NO_3 , $3\text{NH}_4\text{NO}_3$. Decomp. by cold H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Pesci.)

$3\text{NHg}_2\text{NO}_3$, $\text{NH}_4\text{NO}_3 + 2\text{H}_2\text{O}$. Correct formula for mercuriammonium oxydimercuriammonium nitrate, NH_2HgNO_3 , $(\text{NHg}_2\text{OH}_2)\text{NO}_3 + \text{H}_2\text{O}$. (Pesci.)

Dimercuriammonium nitrite, NHg_2NO_2 .

Readily sol. in warm HCl or HBr . (Ray, Chem. Soc. 1902, 81. 648.)

$+ \frac{1}{2}\text{H}_2\text{O}$. Ppt. Sol. in HCl . (Ray, Proc. Chem. Soc. 1902, 18. 85.)

$+ \text{H}_2\text{O}$. (Hofmann and Marburg, A. 1899, 305. 214.)

— **oxide**, $(\text{NHg}_2)_2\text{O}$.

Slowly decomp. by H_2O . Sol. in HCl , or $\text{HNO}_3 + \text{Aq.}$ Decomp. by hot KOH , or $\text{KCl} + \text{Aq.}$ (Weyl, Pogg. 121. 601.)

Sol. in $\text{KCN} + \text{Aq}$ by heating 4–5 hours at 130° . Not completely sol. in HCl owing to formation of Hg_2Cl_2 . (Gaudechon, C. R. 1907, 144. 1419.)

— **phosphate**, $(\text{NHg}_2)_2\text{PO}_4$, $2\text{NHg}_2\text{OH} + 10\text{H}_2\text{O}$.

(Rammelsberg, J. pr. (2) 38. 567.)

Sec Oxydimercuriammonium phosphate.

— **ammonium salicylate**,

$2\text{NHg}_2\text{C}_6\text{H}_4\text{OHCO}_2$, $5\text{NH}_4\text{C}_6\text{H}_4\text{OHCO}_2$.

Decomp. by H_2O . Sol. in $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$, HCl , or $\text{KI} + \text{Aq.}$ (Balestra.)

— **selenate**, $(\text{NHg}_2)_2\text{SeO}_4 + 2\text{H}_2\text{O}$.

Ppt. Insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Cameron and Davy, C. N. 44. 63.)

— **sulphate**, $(\text{NHg}_2)_2\text{SO}_4 + 2\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in $\text{HCl} + \text{Aq.}$ (Rammelsberg, J. pr. (2) 38. 565.) Sol. (Kane), insol. (Hirzel) in $\text{HNO}_3 + \text{Aq.}$

Sol. in KI , or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ with evolution of NH_3 . (Balestra.)

$+ \text{H}_2\text{O}$. Insol. in H_2O ; sol. in HCl . (Ray, Chem. Soc. 1905, 87. 9.)

— **ammonium sulphate**, $(\text{NHg}_2)_2\text{SO}_4$, $3(\text{NH}_4)_2\text{SO}_4 + 4\text{H}_2\text{O}$.

Correct formula for mercuriammonium sulphate, 2NH_2 , HgO , $\text{SO}_3 + \text{H}_2\text{O}$. (Pesci, Gazz. ch. it. 20. 485.)

$5(\text{NHg}_2)_2\text{SO}_4$, $14(\text{NH}_4)_2\text{SO}_4 + 16\text{H}_2\text{O}$. (Pesci.)

$7(\text{NHg}_2)_2\text{SO}_4$, $(\text{NH}_4)_2\text{SO}_4 + 12\text{H}_2\text{O}$. Correct formula for mercuriammonium oxydimercuriammonium sulphate, $(\text{NHg}_2\text{H}_2)_2\text{SO}_4$, $3(\text{NHg}_2\text{OH}_2)_2\text{SO}_4$. (Pesci.)

Dimercuriammonium tartrate, $(\text{NHg}_2)_2\text{C}_4\text{H}_4\text{O}_6 + 2\frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in HCl , KI , $\text{N. NH}_4\text{C}_2\text{H}_3\text{O}_2$, or $(\text{NH}_4)_2\text{C}_4\text{H}_4\text{O}_6 + \text{Aq.}$ (Balestra, Gazz. ch. it. 22, 2. 563.)

— **ammonium tartrate**, $2(\text{NHg}_2)_2(\text{NH}_4)_2\text{C}_4\text{H}_4\text{O}_6 + \text{H}_2\text{O}$.

As above. (B.)

Trimercuriammonium sulphate, $(\text{NHg}_2)(\text{NHgH}_2)\text{SO}_4 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Millon.)

Does not exist. (Pesci, Gazz. ch. 485.)

Dimercuriarsonium mercuric chloride, $\text{AsHg}_2\text{Cl}_2 = \text{AsHg}_2\text{Cl}$, HgCl_2 .

Decomp. by H_2O . Decomp. by $\text{HNO}_3 + \text{Aq.}$ (Rose, Pogg. 51. 423.)

Mercurimidosulphonic acid, $(\text{HO}_2\text{S})_2\text{N}_2\text{Hg}$.

Very unstable. (Berglund, B. 9. 2)

Barium mercurimidosulphonate, $\text{Ba}_2(\text{SO}_3)_2\text{N}_2\text{Hg} + 5\text{H}_2\text{O}$.

(Berglund, B. 9. 256.)

Cadmium —, $\text{Cd}_2\text{HgN}_2(\text{SO}_3)_4 + 12\text{H}_2\text{O}$

Unstable; sl. sol. in H_2O . (Berglund, Soc. (2) 25. 452.)

Cobalt —, $\text{Co}_2\text{HgN}_2(\text{SO}_3)_4 + 15\text{H}_2\text{O}$

Sol. in H_2O . (B.)

Copper —, $\text{Cu}_2\text{HgN}_2(\text{SO}_3)_4 + 15\text{H}_2\text{O}$

Very sol. in H_2O . (B.)

Magnesium —, $\text{Mg}_2\text{HgN}_2(\text{SO}_3)_4 +$

Very sol. in H_2O . (B.)

Manganous —, $\text{Mn}_2\text{HgN}_2(\text{SO}_3)_4 +$

Unstable. (B.)

Mercuric —, $(\text{Hg}_2\text{O})_2\text{HgN}_2(\text{SO}_3)_4$.

Nearly insol. in H_2O . (B.)

Nickel —, $\text{Ni}_2\text{HgN}_2(\text{SO}_3)_4 + 15\text{H}_2\text{O}$

(B.)

Potassium —, $(\text{KO}_2\text{S})_2\text{N}_2\text{Hg} + 4\text{H}_2\text{O}$

Precipitate. (Raschig, A. 241. 161)

Potassium silver —, $(\text{AgSO}_3)_2(\text{KSO}_3) + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Berglund.)

Sodium —, $(\text{NaSO}_3)_2\text{HgN}_2 + 5\text{H}_2\text{O}$

More sol. in H_2O than K salt. (B.)

Strontium —, $\text{Sr}_2(\text{SO}_3)_2\text{HgN}_2 + 15\text{H}_2\text{O}$

More sol. than Ba salt. (B.)

urimidosulphonate,
 $3\text{O}_2, \text{HgN}_2 + 15\text{H}_2\text{O}$.

sol. in H_2O . (B.)

riphosphonium mercuric bro-
 e, $2\text{PHg}_2\text{Br}$, HgBr_2 .

ult, C. R. 1907, 145. 1176.)

iphosphonium mercuric chloride,
 $\frac{1}{2}, \text{PHg}_2\text{Cl}$.

ult, C. R. 1907, 145. 1176.)

H_2O . Decomp. by hot, slowly by
 into Hg , HCl , and H_3PO_3 . Decomp.
 or alkalies. (Rose, Pogg. 40. 75.)

iphosphonium mercuric iodide,
 PHg_2I .

decomp. by cold or warm H_2O ,
 by $\text{MOH} + \text{Aq}$. Not attacked by
 $\text{H}_2\text{SO}_4 + \text{Aq}$. Rapidly attacked by
 and aqua regia. (Lemoult, C. R.
 479.)

iphosphonium mercuric nitrate,
 $\frac{1}{2}, 6\text{HgO}$, $3\text{N}_2\text{O}_5 = 2[\text{PHg}_2\text{NO}_3,$
 $\text{O}_2], 3\text{HgO}$.

Pogg. 40. 75.)

iphosphonium mercuric sulphate,
 $\frac{1}{2}, 6\text{HgO}$, $4\text{SO}_3 + 4\text{H}_2\text{O} = (\text{PHg}_2)_2\text{SO}_4,$
 $\text{O}_4, 2\text{HgO} + 4\text{H}_2\text{O}$.

aqua regia. (Rose, Pogg. 40. 75.)

acid.

mercurate (?).

illet, A. ch. 1. 61.)

mercurate, K_2O , 2HgO .

ly decomp. by H_2O ; less rapidly by
 alcohol. (St. Meunier, C. R. 60.

mercurate, Na_2O , HgO .

off, Bull. Soc. (2) 34. 328.)

ammonium chloride,

H_2N , Cl .

Pogg. 20. 158.)

of Hg , HgNH_2Cl , and NH_4Cl .
 J. pr. (2) 39. 201.)

te, $(\text{NHg}_2\text{H}_2)\text{NO}_3$, "*Hahnemann's*
mercury."

not HCl , and $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. De-
 $\text{NH}_4\text{OH} + \text{Aq}$, or NH_4 salts + Aq .
 mixture of mercurous salts and Hg .

ammonium chloride,

H_2N , Cl .

decomp. (Rose, Pogg. 20. 158.)

of Hg , NH_2HgCl , and NH_4Cl .
 J. pr. (2) 39. 201.)

Mercurodiammonium fluoride,
 $\text{Hg}_2(\text{NH}_2)_2\text{F}_2$ (?).

Decomp. by H_2O . (Finkener, Pogg. 110.
 147.)

Mercuriosulphonic acid.

Mercuriosulphonates, $\text{Hg}(\text{SO}_3\text{M})_2$.

Correct composition for the double sul-
 phites, HgSO_3 , M_2SO_3 . (Divers and Shimid-
 zu, Chem. Soc. 49. 583; Barth, Z. phys. Ch.
 9. 195.)

Mercuroxy-comps.

See Oxymercur- comps.

Mercury, Hg .

Not attacked by H_2O . Not attacked by
 boiling conc. HCl or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Easily
 sol. in dil. or conc. $\text{HNO}_3 + \text{Aq}$; also in HBr
 or $\text{HI} + \text{Aq}$.

Not attacked by pure HNO_3 unless heated,
 but readily attacked by cold dil. $\text{HNO}_3 + \text{Aq}$
 containing NO . (Millon.)

Anhydrous H_2SO_4 attacks Hg gradually
 at ord. temp. (Berthelot, C. R. 1897, 125.
 749.)

H_2SO_4 attacks only when hot and conc.
 (Ditte, A. ch. 1890, (6) 19. 68.)

Conc. H_2SO_4 does not attack dry or moist
 Hg either with or without air. (Pitman, J.
 Am. Chem. Soc. 1898, 20. 100.)

H_2SO_4 attacks Hg at 20° if it contains
 99.7%; does not attack if it contains only
 95.6%. (Baskerville, J. Am. Chem. Soc.
 1898, 20. 515.)

Insol. in $\text{H}_2\text{SO}_4 + \text{Aq}$ alone or in presence
 of HCl or dil. H_2SO_4 . (Berthelot, A. ch.
 1898, (7) 14. 198.)

Not attacked by $\text{HF} + \text{Aq}$ at any temp.
 (Gay-Lussac.)

HI dissolves Hg rapidly at ord. temp.
 (Norris and Cottrell, Am. Ch. J. 1896, 18.
 99.)

More rapidly attacked by $\text{HBr} + \text{Aq}$ than
 by $\text{HCl} + \text{Aq}$. Rapidly acted upon by HI in
 absence of O . (Bailey, Chem. Soc. 1888, 53.
 760.)

Not attacked by pure $\text{HCl} + \text{Aq}$, but in
 presence of O , $\text{Hg}_2\text{OCl}_2 + \text{H}_2\text{O}$ is formed.
 Action is apparently less in sunlight than in
 the dark. (Bailey, Chem. Soc. 1888, 53. 759.)

Small amts. of ferric salts hinder action
 of HNO_3 on Hg , but it is hastened by pres-
 ence of $\text{Mn}(\text{NO}_3)_2$ or NaNO_3 . (Ray, Chem.
 Soc. 1911, 99. 1015.)

HNO_3 under 33% does not attack Hg if
 metal and acid are kept in motion and HNO_3
 is absent. (Veley, B. 1895, 28. 928.)

Rapidly sol. in HClO . (Balard, Dissert.
 1834.)

Alkali chlorides + Aq in presence of air
 decomp. Hg ; action is not increased by heat.
 (Miahle.)

Insol. in alkali chlorides + Aq in neutral or

alkaline solution. (Bhaduri, Z. anorg. 1897, 13. 407.)

Very sol. in conc. solution of I in KI + Aq. (Varet, Bull. Soc. 1897, (3) 17. 451.)

Slowly sol. in KCl or KI + Aq in presence of air. (Palmaer, Z. phys. Ch. 1907, 59. 136.)

Slowly sol. in Na_2S + Aq in presence of air. (Palmaer, Z. phys. Ch. 1907, 59. 137.)

Hg is appreciably sol. in the sulpho salts of Mo, W, V, As, Sb and Sn. (Storch, B. 1883, 16. 2015.)

Persulphates in alkali or neutral solution attack Hg. $(\text{NH}_4)_2\text{S}_2\text{O}_8$ in NH_4OH solution has strongest action. This dissolves Hg by repeated shaking at high temp. (Tarugi, Gazz. ch. it. 1903, 33, (1) 127.)

Insol. in KCN + Aq. (Elmer, J. pr. 1888, (2) 37. 442.)

Slowly sol. in KCN + Aq in presence of air. (Palmaer, Z. phys. Ch. 1907, 59. 136.)

Sol. in considerable quantity in 6% KCN + Aq. (Goyder, C. N. 1894, 69. 268.)

Most sol. in $\text{K}_4\text{Fe}(\text{CN})_6$ + Aq when KOH is present. (Smith, J. Am. Chem. Soc. 1905, 27. 544.)

Hg dissolves in Br_2 in the presence of KBr al. faster than in I_2 , but in CuBr_2 much more slowly. (Van Name and Edgar, Am. J. Sci. 1910, (4) 29. 255.)

Not attacked by PCl_5 even at 350° . (Moissan, A. ch. 1885 (6) 6. 457.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0075 g. Hg in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Mercurous acetylde, $\text{Hg}_2\text{C}_2 + \text{H}_2\text{O}$.

(Burkard and Travers, Chem. Soc. 1902, 81. 1271.)

Mercuric acetylde, basic, 2HgO , $3\text{HgC}_2 + 2\text{H}_2\text{O}$.

(Burkard and Travers, Chem. Soc. 1902, 81. 1272.)

Mercuric acetylde, HgC_2 .

Very sol. in HCl with evolution of H_2C_2 . (Keiser, Am. Ch. J. 1893, 15. 535.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O , alcohol and ether. Sol. in NH_4 acetate + Aq, and in KCN + Aq. Sl. attacked by cold HCl, easily by hot. Easily sol. in HNO_3 . Dil. H_2SO_4 attacks slowly; conc. causes explosion. (Plimpton and Travers, Chem. Soc. 1894, 65. 267.)

Mercuric acetylde chloride, HgC_2 , $\text{HgCl}_2 + \frac{1}{2}\text{H}_2\text{O}$.

Not acted upon by dil. HCl. Decomp. by fuming HNO_3 or aqua regia. Insol. in alcohol and ether. (Keiser, Am. Ch. J. 1893, 15. 538.)

Mercuric acetylde mercuromercuric chloride, HgC_2 , HgCl , $\text{HgCl}_2 + \text{H}_2\text{O}$.

Insol. in all ordinary solvents. (Biltz and Mumm, B. 1904, 37. 4420.)

Mercurous amidofluoride, HgNH_2F .

Insol. in H_2O , HNO_3 and H_2SO_4 dil. HCl.

Probably *dimercuriammonium* amidofluoride, NHg_2F , NH_4F . (Böhm, Z. 1904, 43. 327.)

Mercury ammonium comps.

See—

Mercurioammonium comps., NH_2Hg

Dimercuroammonium comps., NH_2

Mercurous chloramide, $\text{Hg}(\text{NH}_2)\text{Cl}$

Dimercuriammonium comps., NHg

Mercuric chlor-, brom-, etc., $\text{Hg}(\text{NH}_2)\text{R}$.

Mercuridiammonium comps., $\text{Hg}(\text{NH}_2)_2$

Mercuriammonium comps., HgNH_2

Dimercuridiammonium comps., Hg_2

Trimercuriammonium comps., N_3Hg

Oxydimercuriammonium comps., $(\text{NH}_2\text{Hg}_2\text{O})\text{R}$.

Mercurous arsinchloride, AsHgCl .

Decomp. by H_2O . (Capitaine, J. 25. 559.)

Mercurous arsinchloride chloride, $\text{A} = 2\text{AsHgCl}$, Hg_2Cl_2 (?).

Decomp. by H_2O . (Capitaine.)

Mercurous azoimide, HgN_2 .

Wholly insol. in H_2O . (Curtius 3324.)

1 l. H_2O dissolves 0.25 g. (Web Krupko, B. 1913, 46. 2050.)

Mercuric azoimide, HgN_2 .

Sol. in H_2O especially when hot thelot and Vieille, Bull. Soc. 1894, 747.)

Moderately sol. in H_2O . (Web Krupko, B. 1913, 46. 2050.)

Mercuric bromamide, $\text{Hg}(\text{NH}_2)\text{Br}$.

Insol. in H_2O and alcohol. Sl. NH_4OH + Aq. (Mitscherlich, J. pr.)

Correct composition is *dimercurium ammonium bromide*, Hg_2NBr , which see. (Pesci, Gazz. ch. it. 19. 5)

Mercurous bromide, Hg_2Br_2 .

Solubility in $\text{H}_2\text{O} = 9.1 \times 10^{-4}$ g per l. (Bodländer, Z. phys. Ch. 1898)

Solubility in $\text{H}_2\text{O} = 7 \times 10^{-4}$ m liter at 25° . (Sherrill, Z. phys. Ch. 735.)

ty in H₂O at 25° = 1.4×10⁻⁷ equivalent. (Thompson, J. Am. Chem. Soc. 762.)

n H₂O and dil. acids. Decomp. by Sol. in hot conc. H₂SO₄ with of SO₂. Sl. sol. in hot HNO₃+Aq. gr. (Stromann, B. 20. 2818.)

p. into Hg and HgBr₂ by boiling H₂Br, or NH₄Cl+Aq; also by ammonium carbonate or succinate, but not by ammonium sulphate or nitrate. (Witt-

Hg(NO₃)₂+Aq. (Wackenroder, A.

ly decomp. by alkali chlorides+Aq; t of contact of air this decomp. is d HgBr₂ is formed, while in the air the resulting product. Much more lecomp. in hot than cold solutions. A. ch. (3) 5. 177.)

tion of HgBr in 0.1-N KBr contains mg. Hg ions in 1300 l. in alcohol. in benzonitrile. (Naumann, B. 1370.)

in methyl acetate (Naumann, 42. 3790); ethyl acetate. (Naumann, 43. 314.)

in acetone. (Naumann, B. 1904, 37. Eidmann, C. C. 1899, II. 1014.)

c bromide, HgBr₂.

50 pts. H₂O at ordinary temp., and 25 pts. H₂O. (Wittstein.) Sol. in 240 pts. H₂O at Abl.)

94 pts. H₂O at 9°, and in 4-5 pts. at Lassaigne, J. chim. méd. 12. 177.)

lity in g-equivalents per litre = (Bodländer, Z. phys. Ch. 1898, 27.

H₂O dissolves about 4 g. at ord. temp. Z. phys. Ch. 1902, 41. 731.)

H₂O dissolves 0.017 mol. at 25°. (Jansert. 1902.)

lity in H₂O at 25° = 0.017 mol. Sherrill, Z. phys. Ch. 1903, 43.

lity at ord. temp. = 0.8%; at bpt. = (Larine, J. Pharm. 1904, (6) 20.

ility in cold H₂O = 4%, but solution l by heating contains more Hg on of decomp. into HBr and oxybromicario, C. C. 1907, II. 1224.)

H₂O dissolves about 5-6 g. at ord. (Gaudechon, A. ch. 1911, (8) 22.

queous solution at 25° contains 0.017 Herz and Paul, Z. anorg. 1913, 83.

np. by warm HNO₃, or H₂SO₄+Aq. warm H₂SO₄. (Ditte, A. ch. (5)

is sol. in 1 mol. warm HI+Aq. in cold conc. HCl and in 1/2 mol. hot Cl. (Löwig.)

Solubility of HgBr ₂ in KBr+Aq at 25°.			
Mols. per liter			
KBr		HgBr ₂	
0.		0.017	
0.05		0.055	
0.10		0.088	
0.5		0.0359	
0.866		0.611	
2		1.407	
3		2.096	
4		2.339	

(Sherrill, Z. phys. Ch. 1903, 43. 705.)

Solubility in various salts+Aq at 25°.			
Salt	In 10 ccm. of the solution		
	Millimols Hg Br ₂	Millimols salt	
NaBr	0.17	0	
	0.78	1.18	
	2.85	5.96	
	5.40	11.42	
	12.76	24.48	
	15.50	29.97	
	23.06	52.46	
KBr	0.17	0	
	0.98	2.09	
	4.72	7.70	
	13.60	23.80	
	19.30	34.70	
CaBr ₂	0.17	0	
	1.17	0.72	
	6.76	6.45	
	13.58	18.92	
	27.66	24.79	
	36.66	37.54	
SrBr ₂	0.17	0	
	1.04	0.62	
	4.71	3.28	
	9.02	6.68	
	17.70	14.01	
	22.38	18.72	
BaBr ₂	0.17	0	
	3.70	2.74	
	5.40	3.96	
	7.59	5.79	
	14.78	10.96	

(Herz and Paul, Z. anorg. 1913, 82. 434.)

Solubility in 0.1-N Hg(NO ₃) ₂ +Aq is about 20 g. per liter. (Morse, Z. phys. Ch. 1902, 41. 731.)			
Sol. in KBr or NaBr+Aq. • (Jander, Dissert. 1902.)			
Solubility in 10 cc. Br ₂ +Aq at 25°.			
Millimol Br ₂	0.753	1.797	2.231
" Hg	0.1844	0.1947	0.2120

(Herz and Paul, Z. anorg. 1914, 85. 215.)

Sat. solution in liquid SO_2 contains about 1.5% HgBr_2 at 159.4° . (Niggli, Z. anorg. 1912, 75. 182.)

Moderately sol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913, 84. 27.)

1 ccm. of sat. solution in abs. alcohol at ord. temp. contains 0.0958 g. HgBr_2 ; at bpt. contains 0.1262 g. (Hamper, Ch. Z. 1887, 11. 905.)

Solubility of HgBr_2 in alcohols + Aq at t° .

Alcohol	t°	G. HgBr_2 per 100 g. alcohol
Methyl alcohol	0	41.15
	10	49.5
	19	66.3
	22	60.9
	39	71.3
	65	90.8
	97	139.1
Ethyl alcohol	0	25.2
	10	26.3
	19	29.7
	39	31.9
	65	44.5
	89	66.9
Propyl alcohol	0	14.6
	10	15.6
	19	15.5
	39	20.8
	65	31.3
	86.5	42.7
Isobutyl alcohol	0	4.61
	10	5.63
	23	6.65
	39	9.58
	65	15.80

(Timofeiew, Dissert. 1894.)

Much more sol. than HgI_2 in alcohol. 100 g. of solution sat. at 0° contain 13.33–13.05 g. HgBr_2 ; 16.53 g. at 25° ; 22.63 g. at 50° . (Reinders, Z. phys. Ch. 1900, 32. 522.)

Solubility of HgBr_2 in methyl alcohol + Aq at 25° .

P = g. alcohol in 100 g. alcohol + Aq.

HgBr_2 = millimols. HgBr_2 in 10 cc. of the solution.

P	HgBr_2	Sp. gr.	P	HgBr_2	Sp. gr.
0	0.167	1.0022	47.06	0.700	0.9401
10.60	0.201	0.9857	64.00	1.90	0.9386
30.77	0.358	0.9588	78.05	4.07	0.9744
37.21	0.422	0.9508	100	13.96	1.2275

(Herz and Anders, Z. anorg. 1907, 52. 165.)

Solubility of HgBr_2 in ethyl alcohol + Aq at 25° .
P = g. alcohol in 100 g. alcohol + Aq.
 HgBr_2 = millimols. HgBr_2 in 10 cc. of solution.

P	HgBr_2	Sp. gr.
0	0.167	1.0022
20.18	0.187	0.9717
40.69	0.440	0.9435
70.01	1.829	0.9214
100	6.337	0.9873

(Herz and Anders.)

Solubility in mixtures of methyl and propyl alcohol at 25° .

P = % propyl alcohol in the solvent.

G = g. HgBr_2 in 10 ccm. of the solution.

S = Sp. gr. of the sat. solution.

P	G	S $25^\circ/4^\circ$
0	5.02	1.227
11.11	4.728	1.1954
23.8	4.153	1.1524
65.2	2.530	1.0257
91.8	1.635	0.9437
93.75	1.586	0.9368
96.6	1.466	0.9275
100	1.873	0.9213

(Herz and Kuhn, Z. anorg. 1908, 60. 153.)

Solubility in mixtures of ethyl and propyl alcohol at 25° .

P = % propyl alcohol in the solvent.

G = g. HgBr_2 in 10 ccm. of the solution.

S = Sp. gr. of the sat. solution.

P	G	S $25^\circ/4^\circ$
0	2.28	0.9873
8.1	2.225	0.9802
17.85	2.106	0.9740
56.6	1.763	0.9487
88.6	1.476	0.9269
91.2	1.464	0.9239
95.2	1.406	0.9227
100	1.378	0.9213

(Herz and Kuhn, Z. anorg. 1908, 60. 161.)

Solubility of HgBr_2 in mixtures of methyl and ethyl alcohol at 25° .

P = % methyl alcohol in the mixtures.

HgBr_2 = g. HgBr_2 in 10 ccm. of the solution.

S $25^\circ/4^\circ$ = sp. gr. of the sat. solution.

P	HgBr_2	S $25^\circ/4^\circ$
0	2.28	0.9873
4.37	2.31	0.9932
10.4	2.54	1.009
41.02	3.33	1.060
80.69	4.57	1.185
84.77	4.68	1.198
91.25	4.86	1.211
100	5.02	1.227

(Herz and Kuhn, Z. anorg. 1908, 60. 162.)

Cold sat. solution in anhydrous ether contains 0.00567 g. in 1 cc.; hot solution contains 32 g. (Hampe, Ch. Z. 1887, 11. 905.)

Solubility in organic solvents at 18°–20°. 100 g. chloroform dissolve 0.126 g. HgBr_2 . 100 g. tetrachlormethane dissolve 0.003 g. Br_2 .

100 g. bromoform dissolve 0.679 g. HgBr_2 .

100 g. ethyl bromide dissolve 2.310 g. Br_2 .

100 g. ethylene dibromide dissolve 2.340 g. Br_2 .

(Sulc, Z. anorg. 1900, 25. 401.)

Solubility in CS_2 at t° .

t°	100 pts. sat. solution contain pts. HgBr_2
–10	0.049
5	0.068
0	0.087
+ 5	0.105
10	0.122
15	0.140
20	0.187
25	0.232
30	0.274

(Arctowski, Z. anorg. 1894, 6. 267.)

100 g. boiling methyl acetate (bpt. 56.2–6.7°) dissolves 24 g. HgBr_2 . (Schroeder and Leiner, J. pr. 1909, (2) 79. 49.)

1 g. HgBr_2 is sol. in 4.56 g. methyl acetate at 18°. Sp. gr. 18°/4° of sat. solution = 1.09. Naumann, B. 1909, 42. 3795.)

100 g. anhydrous ethyl acetate or sat. with H_2O at 18° dissolve 130.5–135 g. HgBr_2 . (Hamers, Dissert. 1906.)

Solubility of HgBr_2 in ethyl acetate + Aq at 25°.

P = g. ethyl acetate in 100 g. ethyl acetate + Aq.

HgBr_2 = millimols HgBr_2 in 10 cc. of the solution.

P	HgBr_2	Sp. gr.
0	0.167	1.0022
4.39	0.159	1.0018
96.76	7.42	1.1159
100	3.93	1.0113

(Herz and Anders, Z. anorg. 1907, 52. 172.)

1 pt. HgBr_2 sol. in 7.66 pts. ethyl acetate at 18°. (Naumann, B. 1910, 43. 315.)

Easily sol. in acetone. (Oppenheim, B. 2. 72.)

Sol. in acetone. (Eidmann, C. C. 1899, 1. 1014.)

Acetone dissolves much more HgBr_2 than HgI_2 . 100 g. sat. solution at 25° contains 34.58 g. HgBr_2 . (Reinders, Z. phys. ch. 1900, 32. 514.)

Solubility in diethyl oxalate is much greater than that of HgI_2 and is equal to 12% at 100°. (Reinders, Z. phys. Ch. 1900, 32. 507.)

Solubility in benzene = 0.0194 mol. per l. at 25°. (Sherrill, Z. phys. Ch. 1903, 43. 735.)

Sol. in allyl mustard oil. (Mathews, J. phys. Chem. 1905, 9. 647.)

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Solubility in aniline.

S = temp. of solidification.

Mols. HgBr_2 per 100	S	Mols. HgBr_2 per 100	S	Mols. HgBr_2 per 100	S
4.9	9°	25.4	115.5°	49.6	123
10.	43.5	33.9	117	54.9	124
12.5	57	39.5	108	58.8	134
14.9	68	41.9	113	64.0	133
19.7	89	43.9	118		
23.4	106	46.2	121		

(Staronka, Anz. Ak. Wiss. Krakau, 1910, 372.)

Solubility of HgBr_2 in quinoline.

S = temp. of solidification.

Mols. HgBr_2 per 100	S	4.4	8.9	14.3	17.6
		88°	111°	127°	134°

(Staronka, Anz. Wiss. Krakau, 1910. 372.)

Mol. weight determined in ethyl sulphide. (Werner, Z. anorg. 1897, 15. 30.)

+4 H_2O . (Thomsen.)

Mercuric perbromide, HgBr_4 .

(Herz and Paul, Z. anorg. 1914, 85. 216.)

Mercuric hydrogen bromide (Bromomercuric acid), HgBr_2 , $\text{HBr} = \text{HHgBr}_2$.

Decomp. by H_2O . (Neumann, M. 10. 236.)

Mercuric nickel bromide, basic, HgBr_2 , NiBr_2 , $6\text{NiO} + 20\text{H}_2\text{O}$.

(Mailhe, A. ch. 1902, (7) 27. 369.)

Mercuric platinum bromide.

See Bromoplatinate, mercuric.

Mercuric potassium bromide, HgBr_2 , KBr .

Sol. in H_2O , but decomp. by a large amount, with separation of one half of the HgBr_2 . (v. Bonsdorff, Pogg. 19. 339.)

2HgBr_2 , $\text{KBr} + 2\text{H}_2\text{O}$. Permanent. Sol. in H_2O and alcohol. (v. Bonsdorff.)

Mercuric sodium bromide, HgBr_2 , NaBr .

Deliquescent. (v. Bonsdorff.)

(Varet, C. R. 1890, 111, 527.)

HgBr₂, 2NaBr.

Very sol. in H₂O. (Vicario, J. Pharm. 1907, (6) 26. 145.)

2HgBr₂, NaBr+3H₂O. Sol. in H₂O and alcohol. (Berthemot.)

Mercuric strontium bromide, HgBr₂, SrBr₂.

Sol. in all proportions of H₂O. (Löwig, Mag. Pharm. 33. 7.)

2HgBr₂, SrBr₂. Decomp. by H₂O into HgBr₂ and HgBr₂, SrBr₂. (Löwig.)

Mercuric zinc bromide.

Deliquescent in moist air. (v. Bonsdorff.)

Mercuric zinc bromide cyanide ammonia.

See Cyanide zinc bromide ammonia, mercuric.

Mercuric bromide ammonia, HgBr₂, 2NH₃.

Decomp. by boiling H₂O. Sol. in min. acids and acetic acid. (Naumann, B. 1910, 43. 316.)

Mercuric bromide cadmium oxide,

HgBr₂, CdO+H₂O.

(Mailhe, A. ch. 1902, (7) 27. 371.)

Mercuric bromide cupric oxide,

HgBr₂, CuO+3H₂O.

(Mailhe, Bull. Soc. 1901, (3) 25. 791.)

Mercuric bromide hydrazine, HgBr₂, N₂H₄.

(Hofmann and Marburg, A. 1899, 305. 215.)

Mercuric bromide potassium chloride,

HgBr₂, 2KCl.

Decomp. by H₂O. (Harth, Z. anorg. 1897, 14. 345.)

Mercuric bromide zinc oxide, HgBr₂, ZnO+8H₂O.

(Mailhe, C. R. 1901, 132, 1274.)

Mercuric bromoiodide, HgBrI.

Sol. in alcohol and ether. Can be recrystallised from ether without decomp. (Oppenheim, B. 2. 571.)

Mercurous chloramide, Hg₂(NH₂)Cl.

Insol. in boiling H₂O or NH₄OH+Aq. (Kane, A. ch. (2) 72. 215.)

Mixture of Hg and HgNH₂Cl. (Barfoed, J. pr. (2), 39. 201.)

Mercuric chloramide, Hg(NH₂)Cl.

Composition is dimercuriammonium chloride, Hg₂NCl, NH₄Cl see.

Mercuric chloramide oxymercuriam chloride, 4Hg(NH₂)Cl, (NHg₂ (Millon.)

Correct composition is dimercuriam ammonium chloride, NHg₂Cl, NH₄Cl see. (Balestra, Gazz. ch. it. 21. (2) 294.)

Hg(NH₂)Cl, (NHg₂OH₂)Cl. (M

True composition is dimercuriam mercuric chloride, 2Hg₂NCl, HgCl or dimercuriammonium hydrogen (NHg₂Cl, HCl. (Balestra.)

Mercuric chloramide chloride, Hg₂(NH₂)Cl, HgCl.

Properties as mercuric chloramid comp. by cold HCl+Aq. (Millon.)

True composition is dimercuriam hydrogen chloride, NHg₂Cl, 2HCl lestra, Gazz. ch. it. 21. (2) 294.)

Mercuric chloramide chromate,

2Hg(NH₂)Cl, HgCrO₄.

Decomp. by hot H₂O. Easily sol. i or HCl+Aq. (Jäger and Krüss, B. 2

Mercurous chloride, Hg₂Cl₂.

Almost absolutely insol. in cold, but ally sl. decomp. by boiling H₂O.

Calculated from electrical conduc Hg₂Cl₂+Aq, 1 l. H₂O dissolves 3.1 mg at 18°. (Kohlrausch and Rose, Z. p 12. 241.)

1 l. H₂O dissolves 2 mg. Hg₂Cl₂ (Kohlrausch, Z. phys. Ch. 1904, 50. 150.)

1 l. H₂O dissolves 1.4 mg. at 0.5°; 2 18°; 2.8 mg. at 24.6°; 7 mg. at 43°. rausch, Z. phys. Ch. 1908, 64. 150.)

When finely divided, is 10% more when coarsely crystalline. (Sauer, Ch. 1904, 47. 184.)

Solubility in H₂O=0.8×10⁻⁴ g. mc Z. Elektrochem. 1904, 10. 301.)

Sl. sol. with decomp. in boiling 1 from air, 20 ccm. H₂O affording HgCl₂ after boiling 1 hour with (Miahle, A. ch. (3) 5. 176.) Hg₂(Aq containing 1 pt. Hg₂(NO₃), to pts. H₂O give ppt. of Hg₂Cl₂ with H Sol. with decomp. in conc. HCl+HNO₃+Aq, aqua regia, or Cl₂+Aq senius.) Insol. in cold dil. acids, but sol. on heating.

The solubility of Hg₂Cl₂ in HCl increases slowly with time, and finally i point where it increases very rapidly takes place sooner the more dil. t Presence of Hg₂(NO₃),+Aq helps t bility. Why not oxidation to (Varenne, C. R. 92. 1161.)

ality of Hg_2Cl_2 in $\text{HCl} + \text{Aq}$ at 25° .
phase = $\text{Hg}_2\text{Cl}_2 + 0.1 \text{ g. Hg.}$

G. per l.	G. per l.		Sp. gr. of solutions
	H_2O	Hg_2Cl_2	
60	0.034
46	0.048
33	0.207	1.042	...
4	0.399	1.069	...
2	0.548	1.091	...
3	0.654	1.114	...
7	0.675	1.119	...
3	0.670	1.132	...
6	0.673	1.153	...

de and Archibald, Z. phys. Ch. 1902, 40. 385.)

conc. H_2SO_4 does not dissolve or
Boiling H_2SO_4 dissolves with evolution of SO_2 . (Vogel.)

ty of Hg_2Cl_2 in chlorides + Aq at 25° .
phase = $\text{Hg}_2\text{Cl}_2 + 0.1 \text{ g. Hg.}$

G. per liter	G. per liter		Sp. gr. of solutions
	NaCl	Hg_2Cl_2	
1	5.85	0.0041	...
	58.50	0.041	1.040
	119.	0.129	1.078
	148.25	0.194	1.093
	222.3	0.380	1.142
	292.5	0.643	1.188
1	104.15	0.044	1.088
	156.22	0.088	1.134
	208.30	0.107	1.174
	312.54	0.231	1.263
1	39.96	0.022	...
	55.5	0.033	...
	111	0.081	1.064
	138.75	0.118	1.105
	195.36	0.231	1.151
	257.52	0.322	1.205
	324.67	0.430	1.243
	432.9	0.518	1.315
	499.5	0.510	1.358

de and Archibald, Z. phys. Ch. 1902, 40. 385.)

n cold $\text{HCN} + \text{Aq}$ with separation of

n alkali chlorides + Aq. $\text{NH}_4\text{Cl} + \text{Aq}$
e out HgCl_2 at ord. temp., much more
 50° . Dil. $\text{NH}_4\text{Cl} + \text{Aq}$ decomposes
wly than conc. Access of air hastens
L. (Miahle.)

i heated several hours to $40-50^\circ$, 100
 $\text{H}_2\text{Cl} + 833 \text{ pts. H}_2\text{O}$ form 0.75 pt.
from 25 pts. Hg_2Cl_2 ; 100 pts. $\text{NaCl} +$
 H_2O form 0.33 pt. HgCl_2 from 25
 Cl_2 ; 100 pts. $\text{KCl} + 833 \text{ pts. H}_2\text{O}$ form

0.25 pt. HgCl_2 from 25 pts. HgCl_2 ; 100 pts.
 $\text{BaCl}_2 + 833 \text{ pts. H}_2\text{O}$ form 0.33 pt. HgCl_2
from 25 pts. Hg_2Cl_2 . (Miahle, J. Pharm. 26.
108.)

Other chlorides act as NH_4Cl , only less
vigorously. (Pettenkofer.)

By boiling 1 pt. Hg_2Cl_2 10 times with a
solution of 1 pt. NaCl each time, the Hg_2Cl_2 is
finally completely decomp. (Henne.)

Boiling $\text{BaCl}_2 + \text{Aq}$ or $\text{CaCl}_2 + \text{Aq}$ dissolve
traces. $\text{K}_2\text{SO}_4 + \text{Aq}$, $\text{KNO}_3 + \text{Aq}$, or
 $\text{KHC}_4\text{H}_4\text{O}_6 + \text{Aq}$ do not dissolve. (Petten-
kofer.)

Sol. in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. Insol. in NH_4
nitrate, or succinate + Aq. (Wittstein.)

Sol. in hot $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$, and still more
in hot $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$; on cooling it crystal-
lizes out completely. 25 g. Hg_2Cl_2 dissolve in
1.5 l. H_2O containing 50 g. $\text{Hg}(\text{NO}_3)_2$. (De-
bray, C. R. 70. 995.)

Sol. in $\text{PtCl}_2 + \text{Aq}$.

Decomp. by $\text{NH}_4\text{OH} + \text{Aq}$.

Decomp. by KOH , or $\text{NaOH} + \text{Aq}$.

Sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Faktor, C. C.
1906, I. 1524.)

Very sl. sol. in NH_4 succinate. (Witt-
stein.)

Insol. in SbCl_3 . (Klemensiewicz, C. C.
1906, II. 1850.)

Very sol. in liquid NH_3 . (Franklin, Am.
Ch. J. 1898, 20. 829.)

Insol. in alcohol or ether. More sol. in
 H_2O containing pepsin and an acid than in
 H_2O , and is not converted thereby into
 HgCl_2 . (Torsellini, Ann. Chim. Ch. farm.
(4) 4. 105.)

Small amts. are sol. with decomp. in al-
cohol, ether and CHCl_3 . 1 g. CHCl_3 dis-
solves 0.0046 g. Hg_2Cl_2 . (MacLagan, Arch.
Pharm. 1884, 222, 788.)

Formic acid (95%) dissolves at 16.5° ,
0.02%; at 18° , 0.0003%. (Aschan, Ch. Z.
1913, 37. 1117.)

Insol. in methyl acetate (Naumann, B.
1909, 42. 3790); ethyl acetate. (Hamers,
Dissert. 1906; Naumann, B. 1904, 37. 3602.)

Somewhat sol. in hydroxylamine hydro-
chloride. (Adams, Am. Ch. J. 1902, 28. 1216.)

Insol. in benzonitrile. (Naumann, B. 1914,
47. 1370.)

Insol. in acetone and in methylal. (Eid-
mann, C. C. 1899, II. 1014.)

Solubility in organic solvents at $18-20^\circ$.

100 g. chloroform dissolve traces of HgCl_2 .

100 g. bromoform dissolve 0.055 g. HgCl_2 .

100 g. ethyl bromide dissolve traces of
 HgCl_2 .

100 g. ethylene dibromide dissolve traces of
 HgCl_2 . (Sulc. Z. anorg. 1900, 25. 401.)

Mercuric chloride, HgCl_2 .

Permanent.

Sol. in 18.5 pts. H_2O at 13.8° , and 2-3 pts. at 100° .
(J. Davy, 1822.) Sol. in 3 pts. boiling H_2O . (Wenzel.)
Sol. in 18.23 pts. H_2O at 10° , and 3 pts. at 100° . (M. R.
and P.) Sol. in 18.46 pts. at 18.75° . (Abt.) Sol. in 16
pts. cold, and 3 pts. warm H_2O . (Dumas.)

100 pts. H ₂ O dissolve pts. HgCl ₂ at t°:					
t°	Pts. HgCl ₂	t°	Pts. HgCl ₂	t°	Pts. HgCl ₂
0	5.73	40	9.62	80	24.30
10	6.57	50	11.34	90	37.05
20	7.39	60	13.86	100	53.96
30	8.43	70	17.29

(Poggiale, A. ch. (3) 8. 468.)

Solubility of HgCl₂ in H₂O.

t°	% HgCl ₂	t°	% HgCl ₂
+ 1	3.9	80	23.6
4.5	4.8	87	28.2
4.5	4.8	100	39.3
7.5	5.1	121	59.7
13.8	5.2	127	69.7
25.1	7.1	140	77.0
29.5	7.6	150	78.4
38.0	9.9	159	80.2
49	11.3	160	81.7
61	15.1	165	81.8

(Étard, A. ch. 1894, (7) 2. 557.)

71.17 g. HgCl₂ are sol. in 1 l. H₂O at 25°.
(Morse, Z. phys. Ch. 1902, 41. 726.)

Solubility at 25°=0.267 mol. in 1 l. H₂O.
(Jander, Z. Elektrochem. 1903, 8. 688.)

Solubility in H₂O at 25°=0.263 mol. liter.
(Sherrill, Z. phys. Ch. 1903, 43. 735.)

Sat. HgCl₂+Aq at 25° contains 6.9%
HgCl₂. (Foote, Am. Ch. J. 1906, 35. 238.)

HgCl₂+Aq contains 3.95 g. HgCl₂ in 100 g.
of solution at 0°; 7.67 g. at 30°. (Schreine-
makers, Ch. Weekbl. 1910, 7. 202.)

HgCl₂+Aq sat. at (?) contains 6.8%
HgCl₂. (Abe, J. Tok. Chem. Soc. 1912, 33.
1087.)

HgCl₂+Aq sat. at 35° contains 8.51%
HgCl₂. (Schreinemakers and Thonus, Ver.
K. Akad. Wet. Amsterdam, 1912, 21. 333.)

1 l. aqueous solution sat. at 25° contains
0.265 mol. HgCl₂. (Herz and Paul, Z. anorg.
1913, 82. 431.)

H₂O dissolves 7.39% at 20°. (Aschan,
Ch. Z. 1913, 37. 1117.)

* HgCl₂+Aq sat. at 8° has 1.041 sp. gr. (Anthon,
1837.)

Sp. gr. of HgCl₂+Aq at 20°.

% HgCl ₂	Sp. gr.	% HgCl ₂	Sp. gr.
1	1.0072	4	1.0323
2	1.0148	5	1.0411
3	1.0236

(Schröder, calculated by Gerlach, Z. anal.
27. 306.)

Sp. gr. of HgCl ₂ +Aq at 15°		
% HgCl ₂	Sp. gr.	% HgCl ₂
8	1.071	11
9	1.0815	12
10	1.095	13

(Mendelejeff, calculated by Gerlach
27. 306.)

Sp. gr. of HgCl₂+Aq.

% HgCl ₂	Sp. gr.		
	at 0°	at 10°	at 20°
4.72	1.04070	1.04033	1.03856
3.57	1.03050	1.03022	1.02885
2.42	1.02035	1.02018	1.01856
1.22	1.01008	1.00990	1.00835

(Schröder, B. 19. 161 R.)

Sp. gr. of HgCl₂+Aq at room t
aining:

0.226 3.55% 1
1.0233 1.0328

(Wagner, W. Ann. 1883, 18.

Sp. gr. of HgCl₂+Aq at 2

Concentration of HgCl ₂ +Aq.	
1/4-normal	
1/8- "	

(Wagner, Z. phys. Ch. 1890,

Sp. gr. at 16°/4° of HgCl₂+Aq
4.5256% HgCl₂=1.03806.

Sp. gr. at 16°/4° of HgCl₂+Aq
4.2224% HgCl₂=1.03491. (Sch
phys. Ch. 1893, 11. 768.)

HgCl₂+Aq containing 6.04%
sp. gr. 20°/20°=1.0523.

HgCl₂+Aq containing 6.08%
sp. gr. 20°/20°=1.0528.

(Le Blanc and Rohland, Z. phy
19. 282.)

Sat. HgCl₂+Aq boils at 101.1°. (Griff

B.-pt. of HgCl₂+Aq

% HgCl ₂	B.-pt.	% HgCl ₂
4.8	100.10°	11.04
9.0	100.16	15.2

(Skinner, Chem. Soc. 61. 3

Solubility in HCl+Aq is greater t
(Dumas.)

Sol. in 0.5 pt. HCl+Aq of 1.158 sp. gr. s
ing a solution of 2.412 sp. gr. (Davy. 28

Solubility of HgCl_2 in $\text{HCl} + \text{Aq.}$

Pts. HgCl_2 dissolved by 100 pts. liquid	Pts. HCl in 100 pts. H_2O	Pts. HgCl_2 dissolved by 100 pts. liquid
6.8	21.6	127.4
46.8	31.0	141.9
73.7	50.0	148.0
87.8	68.0	154.0

Atte, A. ch. (5) 22. 551.)

$\text{HCl} + \text{Aq}$ at 0° . $\frac{\text{HgCl}_2}{2} = \frac{1}{2}$ mols.

(in mgs.) in 10 ccm. solution; HCl
 HCl ditto; $\text{H}_2\text{O} = \text{grms. H}_2\text{O}$

HCl	Sp. gr.	H_2O
4.3	1.117	9.704
9.9	1.238	9.340
17.8	1.427	9.816
26.9	1.665	8.135
32.25	1.811	7.714
34.25	1.874	7.679
41.5	2.023	7.131
48.1	2.066	6.893
70.875	2.198	6.431

Agel, A. ch. (6) 17. 362.)

sp. by H_2SO_4 or $\text{HNO}_3 + \text{Aq.}$

a. H_2SO_4 , and in more than 500 pts. hot
1.41 sp. gr. without decomp. (J. Davy)

H_2SO_4 , HNO_3 , HIO_3 , or H_2CrO_4
decomp. (Millon, A. ch. (3) 10. 373.)
sol. in HNO_3 , but not decomp.
(Furze.)

Solubility of HgCl_2 in $\text{NH}_4\text{Cl} + \text{Aq}$ at 30° .

of liquid %	Solid phase
70.5	NH_4Cl
56.76	"
50.29	"
41.88	"
32.50	"
25.16	$\text{NH}_4\text{Cl} + \text{HgCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O}$
25.09	$\text{HgCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O}$
24.15	"
22.61	"
21.03	$\text{HgCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O} +$
21.13	$\text{HgCl}_2, \text{NH}_4\text{Cl}, \text{H}_2\text{O}$
24.61	$\text{HgCl}_2, \text{NH}_4\text{Cl}, \text{H}_2\text{O}$
25.12	"
26.47	"
27.17	"
27.60	"

Solubility of HgCl_2 in $\text{NH}_4\text{Cl} + \text{Aq}$
at 30° .—Continued.

Composition of liquid phase			Solid phase
% HgCl_2	% NH_4Cl	% H_2O	
55.55	15.94	28.51	$\text{HgCl}_2, \text{NH}_4\text{Cl}, \text{H}_2\text{O}$
55.85	15.35	28.80	"
56.71	14.22	29.07	"
57.04	14.10	28.86	$\text{HgCl}_2, \text{NH}_4\text{Cl}, \text{H}_2\text{O} +$
56.98	14.14	28.88	$3\text{HgCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O}$
56.83	13.90	29.27	$3\text{HgCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O}$
56.26	13.04	30.70	"
56.43	11.88	31.69	"
56.70	11.05	32.23	"
57.05	9.92	33.02	"
58.55	9.23	32.22	"
58.65	9.20	32.15	$3\text{HgCl}_2, 2\text{NH}_4\text{Cl}, \text{H}_2\text{O} +$
51.83	8.76	39.41	$9\text{HgCl}_2, 2\text{NH}_4\text{Cl}$
46.00	7.52	46.48	$9\text{HgCl}_2, 2\text{NH}_4\text{Cl}$
39.02	6.28	54.70	"
35.60	5.26	59.14	"
35.10	5.18	59.72	"
32.90	5.06	62.04	"
29.65	3.62	66.73	$9\text{HgCl}_2, 2\text{NH}_4\text{Cl} + \text{HgCl}_2$
40.12	5.13	54.75	HgCl_2
21.00	2.29	76.71	"
7.67	0	92.33	"

(Meerburg, Z. anorg. 1908, 59. 139.)

1 pt. sat. $\text{NaCl} + \text{Aq}$ dissolves 1.29 pts.
 HgCl_2 at 14° . (Voit, A. 104. 354.)

Sat. $\text{NaCl} + \text{Aq}$ (20 grains $\text{H}_2\text{O} + 7$ grains NaCl) dis-
solves 32 grains HgCl_2 at 15.5° , and 3 grains more on
warming. Sp. gr. of solution = 2.14 (Davy. 1823.)

Sat. $\text{KCl} + \text{Aq}$ (21 grains $\text{H}_2\text{O} + 7$ grains KCl) dis-
solves 8 grains HgCl_2 on being gently heated (Davy.)

Sat. $\text{BaCl}_2 + \text{Aq}$ (21 grains $\text{H}_2\text{O} + 8.7$ grains $\text{BaCl}_2 +$
 $2\text{H}_2\text{O}$) dissolves 16 grains HgCl_2 at 15.5° , and 4 grains
more on heating. Sp. gr. of solution = 1.9 (Davy.)

$\text{MgCl}_2 + \text{Aq}$ (31 grains $\text{HCl} + \text{Aq}$ of 1.58 sp. gr.
neutralised with MgO) dissolves 40 grains HgCl_2 , and
25 grains more on gently heating. Sp. gr. of solution =
2.83. (Davy.)

Sol. in sat. KCl , $\text{NaCl} + \text{Aq}$, and in MnCl_2 , ZnCl_2 ,
 CoCl_2 , FeCl_2 , NiCl_2 , and $\text{CuCl}_2 + \text{Aq}$. (v. Bonsdorff,
Pogg. 17. 123.)

The solubility in H_2O is greatly increased
by the addition of cupric chloride. 8.5%
 HgCl_2 is sol. in pure H_2O and 52.8% HgCl_2
is sol. in 18.06% $\text{CuCl}_2 + \text{Aq}$. (Schreine-
makers, C. C. 1913, I. 1858.)

Solubility of HgCl ₂ + KCl at 25%.					
Composition of solution		Composition of undissolved residue			Solid phase
% KCl	% HgCl ₂	% KCl	% HgCl ₂	% H ₂ O	
26.46	0	100	0	KCl
26.24	15.04	...	3.63	KCl + 2KCl, HgCl ₂ , H ₂ O
26.23	15.02	...	26.15	"
26.33	15.02	...	52.01	"
26.33	14.92	...	61.04	"
23.74	18.91	34.61	61.66	3.73	2KCl, HgCl ₂ , H ₂ O
22.36	21.39	34.77	62.02	3.21	"
21.39	23.88	34.05	61.84	3.35	"
20.32	27.62	...	65.24	2KCl, HgCl ₂ , H ₂ O + KCl, HgCl ₂ , H ₂ O
20.26	27.38	...	73.98	"
17.85	25.34	21.89	75.10	3.01	KCl, HgCl ₂ , H ₂ O
9.26	18.95	21.02	73.36	5.62	"
7.80	19.56	20.76	73.06	6.18	"
6.84	22.81	20.75	74.54	4.71	"
6.66	24.32	20.54	73.99	5.47	"
6.52	25.13	...	76.46	KCl, HgCl ₂ , H ₂ O + KCl, 2HgCl ₂ , 2H ₂ O
6.64	25.16	...	80.60	"
6.27	25.11	12.09	83.20	4.71	KCl, 2HgCl ₂ , 2H ₂ O
5.77	24.73	11.87	83.18	4.95	"
4.68	24.75	...	84.46	KCl, 2HgCl ₂ , 2H ₂ O + HgCl ₂
4.66	25.17	...	93.58	"
4.69	24.82	...	98.50	"
0	6.90	0	100	0	HgCl ₂

(Foote and Levy, Am. Ch. J. 1906, 35. 239.)

Solubility of HgCl ₂ + KCl at 20°.		
G. per 100 g. H ₂ O		Solid phase
KCl	HgCl ₂	
0	7.39	HgCl ₂
1.12	11.63	"
2.39	15.72	"
4.05	22.17	"
4.84	25.16	HgCl ₂ + 2HgCl ₂ , KCl
5.60	25.13	2HgCl ₂ , KCl
6.71	25.66	"
7.39	26.41	2HgCl ₂ , KCl + HgCl ₂ , KCl
7.46	24.70	HgCl ₂ , KCl
8.95	19.93	"
15	22.87	"
17.57	26.12	"
20.35	29	"
26.31	34.83	"
30.32	39.10	"
34.12	42.82	HgCl ₂ , KCl + HgCl ₂ , 2KCl
34.18	39.34	HgCl ₂ , 2KCl
34.34	35.16	"
34.54	30.63	"
37.72	24.30	"
41.13	19.33	HgCl ₂ , 2KCl + KCl
39.66	15.76	KCl
37.87	10.28	"
35.32	2.1	"

(Tichomiroff, J. russ. Phys. Chem. Soc. 1907 39. 731.)

Solubility of HgCl ₂ + RbCl in H ₂ O. Solubility data are given showing double salt formed at 25°. (Foote and Levy, l.c.)			
Solubility in NaCl + Aq. 100 pts. NaCl + Aq. containing given % NaCl dissolve g. HgCl ₂ .			
% NaCl	g. HgCl ₂ at 15°	g. HgCl ₂ at 65°	g. HgCl ₂ at 100°
26	128	152	206
25	120	142	196
10	58	68	110
5	30	36	64
1	14	18	46
0.5	10	13	44

(Homeyer and Ritsert, Pharm. Ztg. 33. 731.)

Solubility of HgCl ₂ + NaCl at 25°.					
Composition of solution		Composition of undissolved residue			Solid phase
% NaCl	% HgCl ₂	% NaCl	% HgCl ₂	% H ₂ O	
26.5	0	100	0	0	NaCl
18.66	51.35	...	16.39	...	NaCl + NaCl, HgCl ₂ , 2H ₂ O
18.71	51.32	...	21.98	"
18.64	51.42	...	65.45	"
18.87	51.26	...	71.25	"
14.97	57.74	16.38	74.18	9.44	NaCl, HgCl ₂ , 2H ₂ O
14.03	59.69	16.36	74.21	9.43	"
13.25	62.16	16.16	74.70	9.14	"
13.17	62.59	15.96	74.76	9.28	"
12.97	62.50	...	78.20	NaCl, HgCl ₂ , 2H ₂ O + HgCl ₂
13.14	62.48	...	88.64	"
13.15	62.55	...	90.83	"

(Foote and Levy, Am. Ch. J. 1906, 35. 239.)

105.0 g. HgCl₂ are sol. in 1 l. of 0.1-Hg(NO₃)₂ + Aq at 25°. (Morse, Z. phys. C 1902, 4 . 726.)

Solubility in MCl + Aq at 25°.		
Salt	In 10 ccm. of the solution	
	Millimols HgCl ₂	Millimols MCl
LiCl	2.65	0
	3.51	4.14
	6.66	8.35
	10.21	12.71
	16.78	17.38
	22.14	22.65
	28.96	30.91
	30.62	35.27
NaCl	2.65	0
	3.72	2.12
	5.06	4.16
	7.48	6.71
	11.92	11.53
	20.22	19.41
	27.54	27.83
	34.34	31.62

Solubility in MCl + Aq at 25°.—Continued.

Salt	In 10 ccm. of the solution	
	Millimols HgCl ₂	Millimols salt
KCl	2.65	0
	3.55	1.74
	3.81	2.21
	8.36	6.83
MgCl₂	2.65	0
	3.74	1.68
	7.19	4.15
	11.31	5.70
	18.64	9.97
	25.69	13.20
	32.06	17.28
CaCl₂	2.65	0
	3.64	1.90
	7.66	4.02
	11.08	6.56
	18.11	9.64
	26.45	14.29
	33.04	17.23
 SrCl₂	2.65	0
	3.15	1.64
	5.63	3.11
	8.29	5.19
	13.42	7.24
	17.76	10.46
	22.93	13.86
BaCl₂	2.65	0
	6.97	3.85
	11.67	5.72
	16.20	7.76
	26.45	13.36
	53.48	30.30

(Herz and Paul, Z. anorg. 1913, 82. 433.)

Solubility in H₂O is increased by presence of I₂. (Herz and Paul, Z. anorg. 1914, 85. 214.)

Solubility in H₂O is increased by presence of hydroxylamine hydrochloride. (Adams, Am. Ch. J. 1902, 28. 213.)

Moderately sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in liquid CO₂. (Büchner, Z. phys. Ch. 1906, 54. 674.)

Abundantly sol. in H₂PtCl₄ + Aq. (Nilson, B. 1876, 9. 1146.)

Sol. in 2.5 pts. cold alcohol (Richter); 3 pts. (Karl); 2.5 pts. alcohol of 0.833 sp. gr. at ordinary temp., and 1.167 pts. on boiling (Berzelius); 2 pts. alcohol of 0.816 sp. gr. at 15.5 (sp. gr. of solution = 1.08) (J. Davy, Phil. Trans. 1822. 358).

At 10°, sol. in 2.57 pts. alcohol of 39° (Cartier), in 2.9 pts. alcohol of 38°; in 3.6 pts. alcohol of 35°; in 4.2 pts. alcohol of 30°; in 9.3 pts. alcohol of 22°; in 14.6 pts. alcohol of 14°. (N. E. Henry.)

Sol. in 25 mols. methyl, 13.1 mols. ethyl, and 20.3 mols. propyl alcohol at 8.5°; in

16.2 mols. methyl, 12.4 mols. ethyl, and 18 mols. propyl alcohol at 20°; in 6.8 mols. methyl, 10.6 mols. ethyl, and 14.6 mols. propyl alcohol at 38.2°. (Timofejew, C. R., 112. 1224.)

100 pts. absolute methyl alcohol dissolve 66.9 pts. HgCl₂ at 25°; 100 pts. absolute ethyl alcohol dissolve 49.5 pts. HgCl₂ at 25°. (de Bruyn, Z. phys. Ch. 10. 783.)

At 15°, 1 pt. by weight is sol. in:—

13.53 pts. H₂O.

1.5 " methyl alcohol of sp. gr. 0.7990

2.5 " ethyl " " " 0.8100

6.3 " propyl " " " 0.8160

(Rohland, Z. anorg. 1899, 18. 328.)

100 g. HgCl₂ + CH₃OH contain 1.2 g. HgCl₂ at the critical temp. (Centnerszwer, Z. phys. Ch. 1910, 72. 437.)

Solubility of HgCl₂ in methyl alcohol + Aq at 25°.

P = g. alcohol in 100 g. alcohol + Aq.

HgCl₂ = millimols HgCl₂ in 10 cc. of the solution.

P	HgCl ₂	Sp. gr.
0	2.67	1.0565
10.60	2.92	1.0441
30.77	4.18	1.0420
37.21	4.96	1.0507
47.06	7.27	1.0809
64.00	14.19	1.2015
78.05	21.11	1.3314
100	17.95	1.2160

(Herz and Anders, Z. anorg. 1907, 52. 165.)

100 cc. 90% ethyl alcohol dissolve 27.5° g. HgCl₂ at 15.5°. Sp. gr. 15° of sat. solution = 1.065. (Greenish and Smith, Pharm. J. 1903, 71. 881.)

100 g. 99.2% ethyl alcohol dissolve 33.4 g. HgCl₂ at 25°. (Osaka.)

Solubility of HgCl₂ in ethyl alcohol + Aq at 25°.

P = g. alcohol in 100 g. alcohol + Aq.

HgCl₂ = millimols HgCl₂ in 10 cc. of the solution.

P	HgCl ₂	Sp. gr.
0	2.67	1.0565
20.18	2.49	1.0214
40.69	3.94	1.0180
70.01	8.70	1.0616
100	13.61	1.1067

(Herz and Anders, Z. anorg. 1907, 52. 170.)

Solubility of HgCl₂ in ethyl alcohol + Aq at 25°.

% C ₂ H ₅ OH	% HgCl ₂	% C ₂ H ₅ OH	% HgCl ₂
0	6.80	45.84	15.36
5.08	6.65	49.86	18.18
14.49	6.41	53.61	21.40
21	6.55	57.26	24.51
26.25	7.31	60.55	27.67
31.53	8.51	63.95	29.86
36.85	10.32	67.39	32.40
41.36	12.69		

(Abe, J. Tok. Chem. Soc. 1912, 33. 1087.)

Solubility in alcohol is increased by presence of hydroxylamine hydrochloride. (Adams, Am. Ch. J. 1902, 28. 213.)

Solubility of HgCl₂ in a mixture of methyl and ethyl alcohol at 25°.

P = % methyl alcohol in the mixture.
HgCl₂ = g. HgCl₂ in 10 ccm. of the solution.
S = sp. gr. of the sat. solution.

P	HgCl ₂	S 25°/4°
0	3.686	1.107
4.37	3.943	1.130
10.4	4.261	1.157
41.02	5.837	1.294
80.69	6.167	1.321
84.77	5.782	1.288
91.25	5.385	1.254
100	4.862	1.216

(Herz and Kuhn, Z. anorg. 1908, 58. 161.)

Solubility in mixtures of methyl and propyl alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. HgCl₂ in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	4.862	1.2160
11.11	5.034	1.2278
23.8	5.714	1.2848
65.2	4.228	1.1568
91.8	2.509	1.0090
93.75	2.323	1.0029
96.6	2.152	0.9851
100	2.003	0.9720

(Herz and Kuhn, Z. anorg. 1908, 60. 157.)

Solubility in mixtures of propyl and alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. HgCl₂ in 10 ccm. of the solution
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	3.686	1.1070
8.1	3.667	1.0968
17.85	3.406	1.0857
56.6	2.711	1.0272
88.6	2.166	0.9654
91.2	2.160	0.9624
95.2	2.087	0.9772
100	2.003	0.9720

(Herz and Kuhn, Z. anorg. 1908, 60. 157.)
Sp. gr. of HgCl₂ + alcohol.

% HgCl ₂	Sp. gr. 25°/39°
0	0.7948
5.44	0.8346
6.52	0.8431

(Le Blanc and Rohland, Z. phys. Ch. 19. 283.)

Sp. gr. of alcoholic solution of HgCl₂

% HgCl ₂	Sp. gr.		
	at 0°	at 10°	at 20°
0.00	0.83135	0.82286	0.81435
1.22	0.8397	0.8312	0.8228
2.38	0.8484	0.8399	0.8314
4.42	0.8635	0.8549	0.8463
8.56	0.8966	0.8877	0.8789
12.43	0.9306	0.9213	0.9119
15.91	0.9629	0.9523	0.9425
19.32	0.9951	0.9852	0.9753
22.46	1.0285	1.0184	1.0083

(Schröder, B. 19. 161 R.)

Sp. gr. at 16°/4° of HgCl₂ + ethyl containing 23.5489% HgCl₂ = 0.9968 containing 11.8801% = 0.88572. (Schön phys. Ch. 1893, 11. 768.)

Sp. gr. at 16°/4° of HgCl₂ + amyl containing 10.9948% HgCl₂ = 0.8904 (Schönrock, Z. phys. Ch. 1893, 11. 70)

Sol. in 4 pts. ether (Karls); in 1 pt. ether (Henry); in 2.86 pts. ether of 0.744 (sp. gr. of solution = 1.08); the solvent is not increased by elevating the temperature. b.-pt. of ether is not raised. (J. Dav Ether extracts HgCl₂ from Hg (Orfila); very slightly if HgCl₂ + A (Lassaigne.)

Very sl. sol. in pure ether. (Poli 717.)
6.35 pts. are sol. in 100 pts. ether;
6.44 " " " " 100 " "
6.38 " " " " 100 " "

(Laszcynski, B. 1894, 27. 220)

Sol. in 7½–8 pts. ether. (Madsen, Ch. Z. **Supert.** 1897, 21. 169.)

Solubility in 100 cc. ether at 17° = 4.1–4.12 (Strömholm, J. pr. 1902, (2), 66. 450.)

The solubility of HgCl₂ in H₂O is only sl. affected by the presence of ether. An aqueous solution sat. with ether and HgCl₂ contains about 10% less HgCl₂ than a pure sat. aqueous solution. Partition coefficient for

$\frac{\text{HgCl}_2}{\text{H}_2\text{O}} = 4.9$ at 0°; 3.02 at 14.6°; 2.80 at 16.8°. (Strömholm, Z. phys. Ch. 1903, 44. 70.)

Solubility of HgCl₂ in ether + Aq at 25°.

% Ether	% H ₂ O	% HgCl ₂
87.86	5.22	6.92
1.2	93.6	5.2
5.2	90.5	4.3
5.4	91.8	2.8
5.4	93.1	1.5

(Abe, J. Tok. Chem. Soc. 1912, 33. 1087.)

Solubility of HgCl₂ in ether + ethyl alcohol at 25°.

% Alcohol	% HgCl ₂	% Alcohol	% HgCl ₂
67.57	32.43	27.16	36.29
58.59	32.50	22.48	34.08
51.02	37.39	15.20	28.55
44.79	37.96	8.97	20.67
38.69	38.24	0	5.49
32.84	37.75		

(Abe.)

4 pts. ether dissolve 1 pt. HgCl₂, but 4 pts. ether + 1.33 pts. camphor dissolve 1.33 pts. HgCl₂; 4 pts. ether + 4 pts. camphor dissolve 2 pts. HgCl₂; 4 pts. ether + 8 pts. camphor dissolve 4 pts. HgCl₂; 4 pts. ether + 16 pts. camphor dissolve 8 pts. HgCl₂. (Karls, Pogg. 10. 608.)

3 pts. alcohol dissolve 1 pt. HgCl₂, but 3 pts. alcohol + 1 pt. camphor dissolve 2 pts. HgCl₂; 3 pts. alcohol + 3 pts. camphor dissolve 3 pts. HgCl₂; 3 pts. alcohol + 6 pts. camphor dissolve 6 pts. HgCl₂. (Karls, l. c.)

Solution can be obtained containing 25 pts. camphor, 16 pts. HgCl₂, and only 4 pts. alcohol. Sp. gr. of solution = 1.326. (Simon, Pogg. 37. 553.)

100 pts. acetone dissolve 60 pts. HgCl₂ at 25°. (Krug and M'Elroy, J. Anal. Appl. Ch. 184.)

98.35 pts. HgCl₂ are sol. in 100 pts. acetone at 0°.

110.95 pts. HgCl₂ are sol. in 100 pts. acetone at 10°.

126.80 pts. HgCl₂ are sol. in 100 pts. acetone at 18°. (Laszcynski, B. 1894, 27. 2287.)

1 g. HgCl₂ is sol. in 0.70 g. acetone at 18°. Sg. gr. of sat. solution 18°/4° = 1.956. (Naumann, B. 1904, 37. 4334.)

Sat. solution in acetone contains 57.74 g. HgCl₂ in 100 g. solution at 25°. (Foote and Haigh, J. Am. Chem. Soc. 1911, 33. 461.)

Sp. gr. at 26.7°/4° of HgCl₂ + acetone containing 36.25% HgCl₂ = 1.1585. (Schönrock, Z. phys. Ch. 1893, 11. 769.)

Sp. gr. of HgCl₂ + acetone.

% HgCl ₂	Sp. gr. 20°/20°
0	0.8003
10.94	0.8847
21.05	0.9799

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 283.)

100 g. methyl acetate dissolve 46 g. at bpt. (56.5°). (Schroeder and Steiner, J. pr. 1909, (2) 79. 49.)

1 g. HgCl₂ is sol. in 2.35 g. methyl acetate at 18°. Sp. gr. 18°/4° of the sat. solution = 1.251. (Naumann, B. 1909, 42. 3793.)

Solubility in ethyl acetate.

Pts. sol. in 100 pts. ethyl acetate.

Pts. HgCl ₂	t°
28.92	0
29.03	13
30.71	35
31.87	48
32.77	60
35.98	83

(Laszcynski, B. 1894, 27. 2286.)

Solubility in ethyl acetate = 1 : 3.466 at 18°. (Alexander, Dissert. 1899.)

Solubility of HgCl₂ in ethyl acetate.

Temp.	0°	13°	30°	40.5°	50.2°
Mol. HgCl ₂ in 100 mols. C ₄ H ₈ O ₂	15.4	15.9	16.0	16.1	16.3

(Linebarger, Am. Ch. J. 1894, 16. 214.)

1 g. HgCl₂ is sol. in 3.5 g. ethyl acetate at 18°. Sp. gr. of sat. solution 18°/4° = 1.110. (Naumann, B. 1904, 37. 3602.)

Solubility of HgCl₂ in ethyl acetate and acetone at t°.

t°	Molecules HgCl ₂ sol. in 100 molecules of ethyl acetate	Molecules HgCl ₂ sol. in 100 molecules of acetone	Solid present in acetone
-15	9.10	14.5	HgCl ₂ , CH ₃ COCH ₃
0	9.25	14.3	
+10	18.7	
10	23.5	HgCl ₂
17	23.2	
25	9.15	22.8	

(Aten, Z. phys. Ch. 1906, 54. 121.)

Solubility of HgCl₂ in ethyl acetate+Aq at 25°.

P=g. ethyl acetate in 100 g. ethyl acetate +Aq.

HgCl₂=millimols. HgCl₂ in 10 cc. of the solution.

P	HgCl ₂	Sp. gr.
0	2.67	1.0565
4.39	2.72	1.0581
96.76	15.34	1.2371
100	9.75	1.1126

(Hers and Anders, Z. anorg. 1907, 52. 172.)

1 pt. is sol. in 2.05 pts. ethyl acetate at 18° or 100 g. ethyl acetate dissolve 48.7 g. HgCl₂. (Naumann, B. 1910, 43. 315.)

Easily sol. in glycerine; sol. in 14 pts. glycerine. (Fairley, Monit. Scient. (3) 9. 685.)

100 g. glycerine dissolve 80 g. HgCl₂ at 25°. (Moles and Maquina, Ann. Soc. Eshan. fis quin. 1914, 12. 383.)

Solubility in organic solvents

Solvent	t°	Sat. solution contains % HgCl ₂
Methyl alcohol	-34	7.6
	-20	11.5
	-15	12.8
	- 2	18.7
	+ 4	23.2
	12	27.6
	36	53.1
	51	61.0
	62	63.6
	64	63.7
	74	64.3
	100	68.7
	127	75.2

Solvent	t°	Sat. solution contains % HgCl ₂
Ethyl alcohol	-60	3.8
	-55	7.8
	-43	8.8
	-40	9.8
	-30	14.3
	-23	18.6
	-21	19.1
	-20	21.9
	-17	22.1
	-11	24.7
	- 9	27.0
	- 5	29.7
	0	29.0
	+ 3	30.0
	7	30.9
	10	31.3
	14	31.3
	19	32.0
	31	34.2
	43	36.4
	51	38.9
	62	42.1
	63	42.5
	68	44.7
	75	45.2
	80	48.0
	92	51.0
	93	51.4
	100	53.6
	115	60.6
	127	65.3
	138	67.8
N-propyl alcohol	-32	14.7
	-22	15.4
	-14	15.6
	0	16.4
	0	16.5
	+16	18.2
	41	23.8
	53	27.9
	62	29.4
	67	32.7
	78	36.4
	100	43.8
	127	52.7
Allyl alcohol	-21	20.6
	- 1	29.6
	+ 8	35.2
	22	48.7
Acetone	-23	51.4
	-18	52.9
	-15	56.6
	-10	56.7
	- 8	58.4
	- 4	59.1
	- 1	60.1
	+ 6	61.9
	12	61.4
	15	61.8
	27	62.0
	36	61.9
	54	62.1

Solubility in organic solvents.—Continued.

Solvent	t°	Sat. solution contains % HgCl ₂
alcohol	—21	12.4
	— 6	13.0
	+ 9	14.3
	21	15.9
	59	25.8
	82	33.1
alcohol	—11	5.5
	— 6	6.2
	0	6.7
	+11	7.5
	63	19.3
	98	32.1
	127	42.0
	145	47.2
	155	50.4
alcohol	—13	8.6
	+20	8.9
	51	14.0
	90	21.1
	106	35.1
	—47	5.6
	—40	5.8
	—35	6.1
	—30	5.9
	—19	5.6
	0	5.8
	+13	5.8
	83	8.4
	100	8.7
	115	9.0
acetate	—20	29.6
	— 3	29.2
	+24	30.0
	+46	31.0
acetate	—50	39.6
	—20	40.5
	—14	40.2
	— 6	40.0
	0	39.5
	+ 7	39.9
	19	40.2
	45	41.6
	66	44.0
	100	47.8
	131	50.1
	150	57.0
	180	59.3
acetate	—20	42.0
	+24	40.3
	55	41.5
acetate	+22	18.3
	48	18.5
pyrate	+20	12.6
	55	13.5
	71	15.1

Solubility in organic solvents.—Continued.

Solvent	t°	Sat. solution contains % HgCl ₂
Acetic acid	+21	2.7
	22	3.0
	33	5.0
	43	6.0
	50	6.7
	61	8.0
	87	11.0
	95	12.0
	98	12.5
	115	16.0
	116	17.0
	127	20.0
	145	28.3
	182	44.8
	207	55.2
Formic acid	21	2.0
	50	3.2
	90	7.3

Very sl. sol. in propionic and isobutyric acids.

(Étard, A. ch. 1894, (7) 2. 557 et seq.)

Solubility of HgCl₂ in organic solvents at t°.

Solvent	t°	% HgCl ₂
CHCl ₃	—20.5	0.01
	+44.2	0.12
C ₆ H ₆	+6.5	0.26
	18.0	0.53
	34.1	0.64
	54.1	1.02
	69.0	1.39
C ₆ H ₅ Cl	0	1.33
	12.5	1.55
	20.8	1.68
	25.3	1.73
	30.2	1.92
	33.0	2.05
	45.9	2.42
CH ₃ COOC ₂ H ₅	0	22.8
	6.5	22.7
	26.1	22.8
	38.5	23.5
	45.3	26.4

Dukelski, Z. anorg. 1907, 68. 129.

Solubility of HgCl_2 in mixed organic solvents at t° .

Solvent	t°	% HgCl_2
$\text{C}_6\text{H}_6 + \text{C}_2\text{H}_5\text{OH}$	-2.5	15.20
	0.0	15.40
	6.0	16.38
	20.5	18.40
	20.65	18.50
	24.5	19.33
	34.5	21.34
	54.4	24.84
	54.5	24.42
$\text{C}_6\text{H}_6 + 2\text{C}_2\text{H}_5\text{OH}$	-5.2	19.45
	0	20.13
	+9.1	21.65
	20.9	23.57
	24.4	24.19
	36.5	26.53
	53.7	31.27
	74.0	38.74
$\text{CHCl}_3 + \text{C}_2\text{H}_5\text{OH}$	-20.5	3.82
	-12.0	4.43
	0.0	4.89
	+8.0	5.37
	23.0	7.12
	38.5	8.51
	44.2	9.51
	45.6	9.98
$\text{CHCl}_3 + 2\text{C}_2\text{H}_5\text{OH}$	-20.5	6.60
	0.0	7.69
	+8.0	8.96
	23.0	10.66
	38.5	12.50
	44.2	14.40
$\text{CHCl}_3 + \text{CH}_3\text{OH}$	12.0	1.73
	0.0	3.51
	+8.0	5.63
	23.0	10.15
	24.9	10.71
	30.6	11.40
	38.5	12.02
$\text{CHCl}_3 + 2\text{CH}_3\text{OH}$	-12.0	3.33
	0.0	6.73
	+8.0	8.21
	23.0	16.56
	24.9	18.45
	30.6	19.70
	38.5	20.83
$\text{CCl}_4 + 2\text{CH}_3\text{OH}$	0.0	5.20
	7.7	6.69
	24.9	14.06
	30.6	19.40
	35.5	20.50
	36.1	21.80
	48.5	21.90

Solubility of HgCl_2 in mixed organic sol at t° —Continued.

Solvent	t°	%
$\text{C}_2\text{H}_5\text{Cl}_2 + \text{CH}_3\text{OH}$	0.0	13
	12.5	21
	20.8	29
	25.3	34
	30.2	36
	33.0	37
	37.4	37
	45.9	39
$\text{CH}_3\text{COOC}_2\text{H}_5 + \text{C}_6\text{H}_6$	0.0	9
	6.5	9
	25.7	9
	27.6	9
	35.5	10
	45.3	13
$\text{CH}_3\text{COOC}_2\text{H}_5 + \text{CHCl}_3$	0.0	3
	26.1	4
	36.1	4
	46.0	5
	48.5	5
$2\text{CH}_3\text{COOC}_2\text{H}_5 + \text{CCl}_4$	0.0	9
	10.3	9
	25.7	9
	27.6	9
	38.5	9
	45.3	11

(Dukelaki, Z. anorg. 1907, 53, 335.)

Solubility in organic solvents at $18^\circ/3$ 100 g. chloroform dissolve 0.106 g. HgCl_2 .

100 g. tetrachloromethane dissolve 0.0

 HgCl_2 .100 g. bromoform dissolve 0.486 g. HgCl_2 .

100 g. ethyl bromide dissolve 2.04

 HgCl_2 .

100 g. ethylene dibromide dissolve 1.5

 HgCl_2 . (Sule. Z. anorg. 1900, 25, 401.)Solubility of HgCl_2 in various organic solvents at 25° .G. = g. HgCl_2 dissolved in 1 mol. of sol

Solvent	G HgCl_2
Ethylene chloride	1.216
Tetrachlorethane	0.146
Chloroform	0.120
Dichlorethylene	0.110
Pentachlorethylene	
Trichlorethylene	0.034
Perchlorethylene	0.013
Carbon tetrachloride	Trace

(Hofman, et al., B. 1910, 43, 183.)

Very sl. sol in nitromethane at ord. t
 Very sol. on warming. (Bruner, B. 1901
 329b.)

Solubility in CS ₂ at t°.	
t°	100 pts. sat. solution contain pts. HgCl ₂
−10	0.010
− 5	0.014
0	0.018
+ 5	0.022
10	0.026
15	0.032
20	0.042
25	0.053
30	0.063

(Arctowski, Z. anorg. 1894, 6. 267.)

130 g. is dissolved in 100 g. sat. solution S₂ at 8°. (Arctowski, Z. anorg. 1894, 6.)
Formic acid (95%) dissolves 2.1% at 19°. (Kahlan, Ch. Z. 1913, 37. 1117.)
Sol. in molten urethane. (Castoro, Z. g. 1899, 20. 61.)
Sol. in ethyl sulphocyanate. (Kahlenberg, Z. phys. Ch. 1903, 46. 66.)

Solubility of HgCl₂ in benzene.
0 pts. C₆H₆ dissolve at:—
15° 41° 55° 84°
0.54 0.62 0.85 1.80 pts. HgCl₂.
(Laszcynski, B. 1894, 27. 2287.)

Solubility in C₆H₆=0.0197 mol./l. at 25°. (Grill, Z. phys. Ch. 1903, 43. 735.)
Sol. in C₆H₆, toluene, xylene, and other aromatic hydrocarbons. Insol. or only sl. sol. in petroleum ether, hexane, decane and CS₂. (Lewitsch, B. 1904, 37. 1563.)
Sol. in p-toluidine. (Werner.)
Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

Solubility of HgCl₂ in pyridine.
t = point of fusion.
Solid Phase = HgCl₂, 2C₅H₅N.

t°	% HgCl ₂	t°	% HgCl ₂	t°	% HgCl ₂
2.8	2.76	40.90	29.29	78.0	49.72
1.9	7.86	50.10	34.94	78.7	50.37
0.02	13.14	60.03	40.36	80.2	51.52
2.58	17.34	70.15	46.44	82.5	52.40
3.78	19.78	70.8	45.77	89.0	56.45
3.60	21.59	74.6	48.00	90.8	57.01
7.23	22.65	75.2	48.38	94.1	60.09
1.05	24.46	76.4	49.15		

Solubility of HgCl ₂ in pyridine.—Continued.					
t = point of fusion.					
Solid Phase = HgCl ₂ , C ₅ H ₅ N.					
t°	% HgCl ₂	t°	% HgCl ₂	t°	% HgCl
74.7	48.38	90.61	53.50	104.1	60.09
83.5	50.53	75.0°	56.45	104.2	60.72
86.5	52.37	99.5	56.07	104.7	58.97
87.3	52.02	99.5	57.01	107.	63.06
		100.5	57.84		

Solid Phase = 3HgCl ₂ , C ₅ H ₅ N.					
t°	% HgCl ₂	t°	% HgCl ₂	t°	% HgCl ₂
94.7	60.72	113.6	63.06	124.2	65.00
95.2	60.77	114.0	63.18	129.4	65.63
106.4	61.93	115.7	63.37	145.5	69.66
109.8	62.58	118.2	64.09		

(McBride, Z. phys. Ch. 1910, 14. 196.)

Solubility in pyridine.					
S = temp. of solidification.					
Mols. per 100	S	Mols. per 100	S	Mols. per 100	S
5.8	19	27.0	87	38.5	130
5.9	18.5	28.6	(98)	41.0	137
10.2	39.5	30.3	91.5	43.2	142
14.1	52	31.2	92	44.0	143.5
21.4	74.5	33.1	108	47.5	159
25.0	83	35.1	115.5	52.8	173

(Staronka, Anz. Ak. Wiss. Krakau, 1910. 372.)

Sp. gr. at 16°/4° of HgCl₂+pyridine containing 17.53% HgCl₂=1.1523; containing 6.57% HgCl₂=1.0388. (Schönrock, Z. phys. Ch. 1893, 11. 768.)
Mol. weight determined in benzonitrile, methyl- and ethyl-sulphide. (Werner, Z. anorg. 1897, 15. 31. 26 and 30.)
Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Easily sol. in oil of turpentine and other essential oils; sl. sol. in cold benzene, but much more on heating, crystallising on cooling. (Franchimont, B. 16. 387.)
Easily sol. in boiling creosote.
Insol. in olive oil.
Insol. in oils and fats but sol. when first dissolved in alcohol, free ether or anhydrous ketones. (Glock, Ch. Z. Repert. 36. 315.)
Extracted from HgCl₂+Aq by volatile oils.
Mercuric hydrogen chloride (Chloromercuric acid), HgCl₂, HCl=HHgCl₂.
Decomp. by H₂O. (Boullay, A. ch. 34. 243.)
Easily decomposed. (Neumann, M. 10. 236.)

HgCl_2 , $2\text{HCl}+7\text{H}_2\text{O}$. Decomp. by H_2O . (Ditte, A. ch. (5) 22. 551.)

3HgCl_2 , $4\text{HCl}+14\text{H}_2\text{O}$. As above.

2HgCl_2 , $\text{HCl}+6\text{H}_2\text{O}$. As above.

4HgCl_2 , $2\text{HCl}+9\text{H}_2\text{O}$. As above.

3HgCl_2 , $\text{HCl}+5\text{H}_2\text{O}$. As above.

Mercuric hydrazine chloride, HgCl_2 , $2(\text{N}_2\text{H}_4, \text{HCl})$.

Very sol. in H_2O . More sol. in hot alcohol than in cold; decomp. by HNO_3 . (Curtius, J. pr. 1894, (2) 50. 332.)

Mercuric nickel chloride, basic, HgCl_2 , 6NiO , $\text{NiCl}_2+20\text{H}_2\text{O}$, and HgCl_2 , 7NiO , NiCl_2 .

(Mailhe, A. ch. 1902, (7) 27. 369.)

Mercuric nickel chloride.

Deliquescent. (v. Bonsdorff.)

Mercuric nitrosyl chloride, HgCl_2 , NOCl .

Sol. in H_2O without effervescence. (Sudborough, Chem. Soc. 59. 659.)

Mercuric phosphoric chloride, 3HgCl_2 , 2PCl_3 .

Decomp. and dissolved by H_2O . (Baudrimont, A. ch. (4) 2. 45.)

Mercuric potassium chloride, 2HgCl_2 , $\text{KCl}+2\text{H}_2\text{O}$.

Very easily sol. in warm H_2O . A clear solution at 18° is filled with crystals at 15° . Sl. sol. in alcohol. (v. Bonsdorff, Pogg. 17. 122.)

HgCl_2 , $\text{KCl}+\text{H}_2\text{O}$. Easily sol. in H_2O ; sl. sol. in alcohol. (v. Bonsdorff, Pogg. 19. 336.)

HgCl_2 , $2\text{KCl}+\text{H}_2\text{O}$. As above.

Solubility determinations show that the double salts formed by mercuric and potassium chlorides at 25° are:

2KCl , $\text{HgCl}_2+\text{H}_2\text{O}$.

KCl , $\text{HgCl}_2+\text{H}_2\text{O}$. Can be recryst. without decomp.

KCl , $2\text{HgCl}_2+2\text{H}_2\text{O}$. Gives HgCl_2 on recryst. from H_2O . (Foote and Levy, Am. Ch. J. 1906, 35. 237.)

Mercurous rhodium chloride.

See Chlororhodite, mercurous.

Mercuric rubidium chloride, HgCl_2 , RbCl .

Sol. in H_2O .

HgCl_2 , 2RbCl . Sol. in H_2O and $\text{HCl}+\text{Aq}$. (Godeffroy, Arch. Pharm. (3) 12. 47.)

$+2\text{H}_2\text{O}$. Sol. in H_2O . (Godeffroy.)

2HgCl_2 , RbCl . Sol. in H_2O . (Godeffroy.)

Solubility determinations show that at 25° there exist five double mercuric rubidium chlorides with the following formulas:

RbCl , 5HgCl_2 . Gives HgCl_2 on recryst. from H_2O .

3RbCl , $4\text{HgCl}_2+\text{H}_2\text{O}$. Gives RbCl , 5HgCl_2 on recryst. from H_2O .

RbCl , $\text{HgCl}_2+\text{H}_2\text{O}$. Gives 3RbCl , 4HgCl_2 on recryst. from H_2O .

3RbCl , $2\text{HgCl}_2+2\text{H}_2\text{O}$. Gives 4HgCl_2 on recryst. from H_2O .

2RbCl , $\text{HgCl}_2+\text{H}_2\text{O}$. Gives 3RbCl , 4HgCl_2 on recryst. from H_2O .

(Foote and Levy, Am. Ch. J. 1906,

Mercurous silver chloride, HgCl , Ag

(Jones, J. Soc. Chem. Ind. 1893,

2HgCl , AgCl . Min. *Bordosite*.

J. Soc. Chem. Ind. 1893, 12. 983.)

3HgCl , AgCl . (Jones, Chem. Soc. 97. 338.)

Mercuric sodium chloride, HgCl_2 , Na

Sp. gr. at $16^\circ/4^\circ$ of aqueous solution containing 14.937% salt = 1.13310; α 11.0736% = 1.09528. (Schönrock,

Ch. 1893, 11. 782.)

$+\text{H}_2\text{O}$. (Linebarger, Am. Ch. J. 344.)

$+1\frac{1}{2}\text{H}_2\text{O}$. Sol. in 0.33 pt. H_2O (Schindler, Repert. 36. 240.)

Extremely easily sol. in alcohol.

Sol. in 275 pts. ether. Ether dissolves undecomposed salt out of H_2O (Lassaigne, A. ch. 64. 104.)

HgCl_2 , 2NaCl . Deliquescent. in H_2O . (Voit, A. 104. 354.)

2HgCl_2 , NaCl . Decomp. by H_2O solution. Sol. in acetone and acetone (Linebarger, Am. Ch. J. 1893, 15. 34)

Solubility determinations show only double salt formed by mercuric sodium chlorides between 10.3° and NaCl , $\text{HgCl}_2+2\text{H}_2\text{O}$. Can be recryst. from H_2O . (Foote and Levy, Am. Ch. J. 237.)

Mercuric strontium chloride, basic, $\text{HgO}+6\text{H}_2\text{O}$.

Decomp. by H_2O . (André, C. R. 1

Mercuric strontium chloride, 2HgCl_2 , $2\text{H}_2\text{O}$.

Easily sol. in H_2O . (v. Bonsdorff.)

3HgCl_2 , $\text{SrCl}_2+5-6\text{H}_2\text{O}$. Very sol. in H_2O . (Swan, Am. Ch. J. 1898, 20. 1

Mercurous sulphur chloride.

See Mercurous sulphochloride.

Mercuric thallous chloride, HgCl_2 , Tl

Easily sol. in H_2O . (Jørgensen, 6. 83.)

Mercurous stannous chloride, Hg_2Cl_2

Decomp. by H_2O . (Capitaine, J. 25. 549.)

Mercuric yttrium chloride, 3HgCl_2 , $9\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O . (A. 131. 179.)

inc chloride, HgCl_2 , ZnCl_2 .
in H_2O . (Harth, Z. anorg. 1897,
 ZnCl_2 . (Varet, C. R. 1896, 123.)

zinc chloride ammonia, HgCl_2 ,
 $2, 10\text{NH}_3 + 2\text{H}_2\text{O}$.
boiling H_2O , but decomp. thereby.
R. 112. 995.)
 ZnCl_2 , $6\text{NH}_3 + \frac{1}{2}\text{H}_2\text{O}$. As above.

chloride ammonia, HgCl_2 , 12NH_3 .
in ammonia. (Franklin, Am. Ch.
. 300.)

chloride cadmium oxide, HgCl_2 ,
 H_2O .
A. ch. 1902, (7) 27. 371.)

chloride cobaltous oxide, HgCl_2 ,
 $+\frac{1}{2}\text{H}_2\text{O}$.
C. R. 1901, 132. 1274.)

chloride cupric oxide, HgCl_2 , 3CuO
).
Bull. Soc. 1901, (3) 25. 791.)

chloride hydrazine, HgCl_2 , N_2H_4 .
stable. Decomp. by H_2O . Pptd.
sol solution by H_2O ; very sol. in
with decomp.
sol. in HCl or HNO_3 . Decomp. by
somewhat sol. in acetic acid. (Hof-
1897, 30. 2020.)

chloride hydroxylamine, HgCl_2 ,
 OH .

ely sol. in methyl and ethyl al-
l. in ether; decomp. by H_2O and
q. Sol. in NH_4OH , $\text{HCl} + \text{Aq}$.
m. Ch. J. 1902, 28. 210.)

chloride lead oxide, HgCl_2 , $2\text{PbO} +$
.
A. ch. 1902, (7) 27. 372.)

chloride strontium chromate,
 I_2 , HCl , SrCrO_4 .
 H_2O without decomp. (Imbert,
1897, (3) 17. 471.)

chloroiodide, 2HgCl_2 , HgI_2 .
 I_2O . (Liebig.)
 HgI_2 . Sl. sol. in hot H_2O with
comp. More easily sol. in alcohol.
. 12. 1187.)

fluoride, Hg_2F_2 .
. by H_2O with separation of Hg_2O .

luoride, $\text{HgF}_2 + 2\text{H}_2\text{O}$.
. by cold H_2O , with separation of
in dil. $\text{HNO}_3 + \text{Aq}$, and $\text{HF} + \text{Aq}$.
Pogg. 110. 628.)

Mercurous hydrogen fluoride, Hg_2F_2 , $4\text{HF} +$
 $4\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O . Sol. in
dil. acids and dil. HF . (Böhm, Z. anorg.
1905, 43. 327.)

Mercurous silicon fluoride.

See Fluosilicate, mercurous.

Mercurous fluoride ammonia, Hg_2F_2 , 2NH_3 .
Stable on air. (Finkener, Pogg. 110. 142.)

Mercurous hydroxide, HgOH .

Nearly insol. in cold, sol. in hot H_2O .
Sol. in $\text{NaOH} + \text{Aq}$. (Bhaduri, Z. anorg.
1897, 13. 410.)

Mercurous iodamide, $\text{Hg}_2(\text{NH}_2)\text{I}$.

(Rammelsberg, Pogg. 48. 184.)
Is a mixture of Hg and $\text{Hg}(\text{NH}_2)\text{I}$. (Bar-
foed.)

Mercurous iodide, Hg_2I_2 .

Sol. in over 2375 pts. H_2O . (Saladin, J.
chim. méd. 7. 530.)

Solubility in $\text{H}_2\text{O} = 2.6 \times 10^{-8}$ g.-equiv. per
liter (calculated). (Bodländer, Z. phys. Ch.
1898, 27. 58.)

Solubility in $\text{H}_2\text{O} = 3 \times 10^{-10}$ mols. per litre
at 25° . (Sherrill, Z. phys. Ch. 1903, 43. 735.)

Sol. in $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$. (Stromann, B. 20.
2815.)

Sol. in $\text{KI} + \text{Aq}$. Easily sol. in $\text{Hg}_2(\text{NO}_3)_2 +$
 Aq . Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in hot
 $\text{NH}_4\text{Cl} + \text{Aq}$, but less than HgI_2 . Less sol.
in NH_4NO_3 than in $\text{NH}_4\text{Cl} + \text{Aq}$. (Brett.)

Partially sol. with separation of Hg and
formation of HgI_2 , in cold $\text{KI} + \text{Aq}$, hot NaI ,
 CaI_2 , SrI_2 , BaI_2 , MgI_2 , ZnI_2 , and $\text{NH}_4\text{I} + \text{Aq}$;
in warm NaCl , KCl and $\text{NH}_4\text{Cl} + \text{Aq}$, and
slowly in hot $\text{HCl} + \text{Aq}$. (Boullay, A. ch. (2)
34. 358.)

Decomp. by alkali chlorides $+ \text{Aq}$. (Miahle,
A. ch. (3) 5. 177.)

Very easily sol. in liquid NH_3 . (Franklin,
Am. Ch. J. 1898, 20. 829.)

Not wholly insol. in alcohol, ether, or chlor-
oform. (MacLagan, Rep. anal. Ch. 1884. 378.)

Decomp. by boiling alcohol; 1000 g. boiling
alcohol decomp. about 3.15 g. Hg_2I_2 . (Fran-
çois, C. R. 1896, 121. 890.)

Boiling alcohol decomp. Hg_2I_2 to Hg and
 HgI_2 , which dissolves until 0.220 g. HgI_2 are
contained in 100 g. alcohol. (François, C. R.
1896, 121. 889.)

Insol. in cold ether. (François, J. Pharm.
1897, (6), 6. 445.)

Insol. in methylene iodide. (Retgers, Z.
anorg. 3. 345.)

Difficultly sol. in methyl acetate. (Nau-
mann, B. 1909, 42. 3790.)

Phenol at 180° decomp. it into Hg and HgI_2 ,
until a state of equilibrium is reached with

2.75 g. HgI_2 to 100 g. phenol, above which point HgI is sl. sol. (0.05 g. in 100 g.) in phenol- HgI_2 mixture. Decomp. by cold aniline more rapidly than by hot. Equilibrium is reached when 26.35 g. HgI_2 are present to 100 g. aniline at bpt. of aniline. Aniline containing more than 26 g. HgI_2 to 100 g. dissolves HgI in considerable quantity. (François, C. C. 1896, I, 470.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sl. sol. in allyl mustard oil. (Mathews, J. phys. Chem. 1905, 9. 647.)

Mercuric iodide, HgI_2 .

Sol. in 150 (?) pts. H_2O . (Wurtz.)

1 l. H_2O at 17.5° dissolves 0.0403 g. HgI_2 . (Bourgoin, A. ch. (6) 3. 429.)

Sol. in about 6500 pts. H_2O . (Hager.)

According to calculation from electrical conductivity of $\text{HgI}_2 + \text{Aq}$, HgI_2 is much less sol., 1 l. H_2O dissolving only 0.5 mg. HgI_2 at 18° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

1 l. H_2O dissolves 0.054 g. HgI_2 at 22° . (Rohland, Z. anorg. 1898, 15. 412.)

1 l. H_2O at 25° dissolves about 0.06 g. (Morse, Z. phys. Ch. 1902, 41. 731.)

1 l. H_2O at 18° dissolves 4×10^{-6} mol. (Abegg, Z. Elektrochem. 1903, 9. 553.)

Solubility in H_2O at $25^\circ = 0.00013$ mol. liter. (Sherrill, Z. phys. Ch. 1903, 43. 735.)

1 l. H_2O dissolves 0.4 mg. HgI_2 at 18° . (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)

1 l. H_2O at 18° dissolves 0.2 to 0.4 mg. (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

The yellow modification is always deposited from solution even in the presence of an excess of the red form. (Gernez, C. R. 1903, 136. 1323.)

Sol. in many acids, especially in HCl , and $\text{HI} + \text{Aq}$. Insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Berthelot.) Scarcely sol. in dil. $\text{HNO}_3 + \text{Aq}$.

Not attacked by cold H_2SO_4 , decomp. by hot. (Ditte, A. ch. 1879, (5) 17. 124.)

Sat. solution in $\text{H}_2\text{SO}_4 + \text{Aq}$ contains at critical temp. (158.2°), 0.7% HgI_2 . (Niggli, Z. anorg. 1912, 75. 182.)

Sol. in hot $(\text{NH}_4)_2\text{CO}_3$, $(\text{NH}_4)_2\text{SO}_4$, cold NH_4Cl , NH_4NO_3 , or ammonium succinate + Aq . (Wittstein.)

Sol. in HgCl_2 , $\text{Hg}(\text{NO}_3)_2$, or $\text{Hg}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$. Easily sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. Easily sol. in soluble iodides + Aq . More sol. in hot than in cold NaI or $\text{KI} + \text{Aq}$. When conc., 1 mol. KI in hot solution dissolves 3 mols. HgI_2 , but a portion separates on cooling. BaI_2 , SrI_2 , MgI_2 , and CaI_2 act in the same way. Easily sol. in cold, more sol. in hot $\text{ZnI}_2 + \text{Aq}$, 2 mols. HgI_2 being dissolved to 1 mol. ZnI_2 . In $\text{NH}_4\text{I} + \text{Aq}$, 3 mols. HgI_2 are dissolved to 2 mols. NH_4I . Abundantly sol. in hot KCl , NaCl , $\text{NH}_4\text{Cl} + \text{Aq}$, but separates out on cooling, and the trace remaining may be pptd. by H_2O , 2 g. KCl in solution dissolves 1.166 g. HgI_2 . Sol. in $\text{HgCl}_2 + \text{Aq}$, and very easily sol. in

alcoholic solution of HgCl_2 . (Boullay, A. ch (2) 34. 346.)

Solubility in $\text{MI} + \text{Aq}$ at 25° .

Salt	In 10 cc. of the solution	
	Millimols HgI_2	Millimols salt
NaI	4.12	7.94
	6.22	13.85
	9.45	22.25
KI	1.27	3.03
	1.80	3.90
	5.10	10.34
	7.00	15.54
	12.24	25.19
CaI_2	0.50	0.53
	2.61	2.52
	4.40	4.68
	4.58	4.84
	17.06	17.99
SrI_2	2.12	2.54
	3.20	3.55
	5.82	5.39
	6.94	6.08
BaI_2	0.59	0.99
	7.42	7.48
	8.98	9.78
	14.62	15.08

(Herz and Paul, Z. anorg. 1913, 82. 434.)

Solubility of $\text{HgI}_2 + \text{KI}$ in H_2O .

Temp. = 20° .		
% KI	% HgI_2	Solid phase
50.9	19.3	KI
44.4	32.4	"
39	48	"
37.4	53.6	$\text{KI} + \text{KHgI}_2$
37.8	52.6	KHgI_2
35.1	52.2	"
35.5	51.2	$\text{KHgI}_2, \text{H}_2\text{O}$
26.7	50.3	$\text{KHgI}_2 + \text{HgI}_2$
26.6	49.4	HgI_2
23.7	40.2	"
14.9	22.5	"
Temp. = 30°		
60.6	...	KI
40	53	$\text{KI} + \text{KHgI}_2$
39.6	52.7	KHgI_2
40	52.2	"
40.2	51.2	"
39.3	50.3	"
33.7	49.8	"
33	52	"
31.4	51.7	$\text{KHgI}_2, \text{H}_2\text{O}$
29.1	52.2	"

(Dunningham, Chem. Soc. 1914, 105. 300)

ery sol. in KSCN + Aq. (Philipp, Pogg. 131. 93.)
l. in 1.09 pts. cryst. Na₂S₂O₃ + Aq. r and Ulm, M. 1882, 3. 197.)
ery sol. in hot CaCl₂ + Aq, less sol. in l₂, KCl and NaCl + Aq. (Lea, Z. anorg. 12. 341.)
ubility in normal Hg(NO₃)₂ + Aq = g. per litre. (Morse, Z. phys. Ch. 1902, 31.)
xtremely sol. in cold conc. NH₄Br + Aq. esmann, B. 1903, 36. 1602.)
l. in alkali sulphites + Aq. (Barth, Z. Ch. 1892, 9. 215.)
l. in Ca(OCl)₂ + Aq; sol. in KOH + Aq. sens, A. ch. (3) 26. 222.)
l. in liquid SO₂. (Walden and Cent- rwer, C. C. 1902, I. 344.)
ry easily sol. in liquid NH₃. (Franklin, Ch. J. 1898, 20. 829.)
l. in SOCl₂, S₂Cl₂, SO₂Cl₂, warm AsCl₃, , warm POCl₃. (Walden, Z. anorg. 25. 212.)
sily sol. in AsBr₃. (Walden, Z. anorg. 29. 374.)
sol. in liquid CO₂. (Büchner, Z. phys. 1906, 54. 674.)
ore sol. in alcohol than in H₂O. 1 l. containing 10% of 90% alcohol dis- s 0.08 g. HgI₂. 1 l. of alcohol of 80° B. ves 2.851 g. HgI₂, 1 l. absolute alcohol ves 11.86 g. HgI₂. (Bourgoin, A. ch. . 429.)
. in 130 pts. cold, and 15 pts. hot 90% ol. (Hager.)
) pts. absolute methyl alcohol dissolve pts. at 19.5°; 100 pts. absolute ethyl ol dissolve 2.09 pts. at 19.5°. (de n, Z. phys. Ch. 10. 783.)
0842 pt. is sol. in 1 pt. alcohol at 15°. tier and Charpy, C. R. 1890, 111.
) g. methyl alcohol dissolve 3.7 g. HgI₂ °; ethyl alcohol, 1.86 g.; propyl alcohol, g.; isobutyl alcohol, at 22.5°, 0.51 g. ofeiew, Dissert. 1894.)
15-20°, 100 g. methyl aleohol dissolve g. HgI₂; ethyl alcohol, 1.42 g.; propyl ol, 0.826 g. (Rohland, Z. anorg. 1898, 12.)

ility of HgI₂ in ethyl alcohol + Aq at 25°.
=g. alcohol in 100 g. alcohol + Aq.
l₂=millimols. HgI₂ in 100 cc. of the ion.

A	HgI ₂	Sp. gr.
0	3.86	0.80325
5.82	2.56	0.80950
2.44	1.92	0.81536
6.74	1.38	0.82996
8.75	0.935	0.84654
7.63	0.45	0.87214

rs and Knoch, Z. anorg. 1905, 45. 266.)

Solubility of HgI₂ in methyl alcohol + Aq at 25°.

P = g. alcohol in 100 g. alcohol + Aq.
HgI₂ = millimols. HgI₂ in 10 cc. of the solution.

P	HgI ₂	Sp. gr.
0	0.0013	...
47.06	0.0098	0.9187
64.00	0.0347	0.8834
78.05	0.0981	0.8519
100	0.571	0.8155

(Herz and Anders, Z. anorg. 1907, 52. 165.)

Solubility of HgI₂ in ethyl alcohol + Aq at 25°.

P = g. alcohol in 100 g. alcohol + Aq.
HgI₂ = millimols. HgI₂ in 10 cc. of the solu- tion.

P	HgI ₂	Sp. gr.
70.01	0.061	0.8636
100	0.386	0.8032

(Herz and Anders, Z. anorg. 1907, 52. 170.)

At 15°, 1 pt. by weight is sol. in:—
24813 pts. H₂O.
30.8 pts. methyl alcohol of sp. gr. at 0.7990.
70.3 " ethyl " " " " 0.8100.
121.0 " propyl " " " " 0.8160.

(Rohland, Z. anorg. 1899, 18. 328.)

Solubility of HgI₂ in mixtures of methyl and ethyl alcohol at 25°.

P = % methyl alcohol in the mixtures.
HgI₂ = g. HgI₂ in 10 ccm. of the solution.
S 25°/4° = Sp. gr. of the sat. solution.

P	HgI ₂	S 25°/4°
0	0.180	0.8038
4.37	0.193	0.8039
10.4	0.208	0.8046
41.02	0.232	0.8077
80.69	0.289	0.8131
84.77	0.296	0.8140
91.25	0.298	0.8146
100	0.316	0.8156

(Herz and Kuhn, Z. anorg. 1908, 58. 164.)

Solubility in mixtures of methyl and propyl alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. HgI₂ in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	0.316	0.8156
23.8	0.304	0.8155
91.8	0.169	0.8101
93.75	0.167	0.8110
96.6	0.153	0.8108
100	0.142	0.8116

(Herz and Kuhn, Z. anorg. 1908, 60. 158.)

Solubility in mixtures of propyl and ethyl alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. HgI₂ in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	0.180	0.8038
8.1	0.173	0.8036(?)
17.85	0.165	0.8043
56.6	0.155	0.8075
91.2	0.152	0.8099
95.2	0.144	0.8108
100	0.142	0.8116

(Herz and Kuhn, Z. anorg. 1908, 60. 161.)

Solubility in 100 pts. amyl alcohol equals:

0.66 pts. at 13°.
3.66 " " 71°.
5.30 " " 100°.
9.57 " " 133.5°.

(Laszcynski, B. 1894, 27. 2287.)

Sp. gr. at 16°/4° of HgI₂ + alcohol containing 1.8358% HgI₂ = 0.80718; containing 1.7119% = 0.80597. (Schönrock, Z. phys. Ch. 1893, 11. 770.)

Somewhat sol. in ether. Sol. in 77 pts. ether. (Saladin.) Sol. in 60 pts. ether. (Hager.)

Sol. in cold ether. (François, J. Pharm. 1897, (6) 6. 445.)

Very sl. sol. in anhydrous ether. (Hampe.)

0.62 pt. is sol. in 100 pts. ether at 0°.

0.97 pt. is sol. in 100 pts. ether at 36°.

(Laszcynski, B. 1894, 27. 2286.)

Solubility in ether = 0.3% at ord. temp. (Marsh, Chem. Soc. 1910, 97. 2299.)

Nearly insol. in ether. (Dunningham, Chem. Soc. 1914, 105. 368.)

Data are given on the system HgI₂ + KI + ether. (Dunningham.)

Solubility at 23° in chloroform = 0.071%; in ether = 0.551%; in acetone = 2.005%; in ethyl alcohol = 2%; in methyl alcohol = 3.975%; in benzene = 0.247%. (Beckmann and Stock, Z. phys. Ch. 1895, 17. 130.)

Solubility in organic solvents at t°.

Solvent	t°	100 g. of the solvent dissolve g. HgI ₂
Chloroform	61	0.163
Tetrachlormethane	75	0.094
Ethylene dichloride	85.5	1.200
Isobutyl chloride	69	0.328
Ethyl bromide	38	0.773
Methyl alcohol	66	6.512
Ethyl alcohol	78	4.325
Isopropyl alcohol	81	2.266
Isobutyl alcohol	ca. 100	2.433
Methyl formate	36-38	1.166
Ethyl formate	52-55	2.150
Methyl acetate	56-59	2.500
Ether	35	0.470
Acetone	56	3.249
Acetal	ca. 100	2.000
Chloral	96	...
Epichlorhydrin	ca. 100	6.113
Hexane	67	0.072
Benzene	80	0.825
Ethyl acetate	74-78	4.200

(Sulc, Z. anorg. 1900, 25. 402.)

Solubility in organic solvents at 18-20°

100 g. chloroform dissolve 0.040 g. HgI₂.
100 g. tetrachlormethane dissolve 0.008 g. HgI₂.

100 g. bromoform dissolve 0.486 g. HgI₂.
100 g. ethyl bromide dissolve 0.643 g. HgI₂.

100 g. ethyl iodide dissolve 2.041 g. HgI₂.
100 g. ethylene dibromide dissolve 0.748 g. HgI₂.

(Sulc, Z. anorg. 1900, 25. 401.)

1 pt. ethylene bromide dissolves 0.008 pts. HgI₂ at 15°. (Gautier and Charpy, C. R. 1890, 111. 647.)

100 pts. methylene iodide CH₂I₂ dissolve 2.5 pts. HgI₂ at 15°, 16.6 pts. at 100°, and 1 pt. at 180°. (Retgers, Z. anorg. 3. 252.)

1 l. sat. solution in CCl₄ at 15° contain 0.170 g. HgI₂. (Dawson, Chem. Soc. 1909, 95. 874.)

Sol. in 340 pts. glycerine. (Fairley, Mon. Scient. (3) 9. 685.)

100 pts. acetone dissolve 2.09 pts. HgI₂ at 25°. (Krug and M'Elroy, J. Anal. Ch. 1884, 84.)

Sol. in acetone and in methylal. (Eidman, C. C. 1899, II, 1014.)

Solubility in 100 pts. acetone equals:

2.83 pts. HgI₂ at — 1°.
3.36 " " " 18°.
4.73 " " " 40°.
6.07 " " " 58°.

(Laszcynski, B. 1894, 27. 2287.)

100 g. methyl acetate solution, sat. at 18° contain 1.10 g. HgI₂. (Beaold, Diss. 1906.)

g. boiling methyl acetate slowly dissolves 2.3 g. HgI_2 . (Schroeder and Steiner, 1909, (2) 79. 49.)

Solubility in ethyl acetate at t° .

sol. in 100 pts. ethyl acetate	t°
1.49	- 2
1.56	+17.5
1.64	21
2.53	40
3.19	55
4.31	76

(Laszczynski, B. 1894, 27. 2286.)

g. ethyl acetate anhydrous, or sat. H_2O at 18° , dissolve at 18° , 14.70 g. Solubility increases somewhat with temperature. (Hamers, Dissert. 1906.)

Solubility of HgI_2 in ethyl acetate + Aq at 25° .

g. ethyl acetate in 100 g. ethyl acetate

, = millimols. HgI_2 in 10 cc. of the solvent.

	HgI_2	Sp. gr.
.39	0.0028	0.9973
.76	0.412	0.9063
	0.241	0.9011

and Anders, Z. anorg. 1907, 52. 172.)

HgI_2 is sol. in 68.03 pts. ethyl acetate at 18° . (L. Mann, B. 1910, 43. 316.)

Solubility in diethyl oxalate is 12.5% at 25° and 2.5% at 100° . (Reinders, Z. phys. Chem. 1900, 32. 507.)

Solubility in CS_2 at t° .

t°	100 pts. sat. solution contain pts. HgI_2
-10	0.107
- 5	0.141
0	0.173
+ 5	0.207
10	0.239
15	0.271
20	0.320
25	0.382
30	0.445

Arctowski, Z. anorg. 1894, 6. 267.)

Solubility in CS_2 .

100 g. of the sat. solution contain at:

-86.5° -93° -116°
0.024 0.023 0.017 g. HgI_2 .

(Arctowski, Z. anorg. 1896, 11. 274.)

0.0028 pt. is sol. in 1 pt. CS_2 at 15° . (Gautier and Charpy, C. R. 1890, 111. 647.)

1 l. sat. solution in CS_2 at 15° contains 3.127 g. HgI_2 . (Dawson, Chem. Soc. 1909, 95. 874.)

Very sol. in liquid methylamine. (Gibbs, J. Am. Chem. Soc. 1906, 28. 1419.)

Abundantly sol. in methylamine. (Fitzgerald, J. phys. Chem. 1912, 16. 633.)

Somewhat sol. in allyl mustard oil. (Mathews, J. phys. Chem. 1905, 9. 647.)

Sol. in $\text{Sb}(\text{CH}_3)_3 + \text{Aq}$.

Very sl. sol. in Na citrate + Aq. (Spiller.)

1 pt. C_6H_6 dissolves 0.00217 pts. HgI_2 at 15° . (Gautier and Charpy, C. R. 1890, 111. 647.)

Solubility in 100 pts. benzene equals:

0.22 pts. at 15° .

0.88 " " 60° .

0.95 " " 65° .

1.24 " " 84° .

(Laszczynski, B. 1894, 27. 2284.)

1 l. C_6H_6 dissolves 0.00493 mol. HgI_2 at 25° . (Sherrill, Z. phys. Ch. 1903, 43. 735.)

100 g. boiling phenol dissolve 10 g. HgI_2 . (François, C. R. 1895, 121. 769.)

Sl. sol. in phenol with 20% H_2O . Not very sol. in acetic acid at 119° , in amyl acetate at 133° , in amyl bromide at 119° . Rather sol. in diethyl oxalate at 186° , in ethylene bromide at 131° , in amyl alcohol at 137° , in amyl iodide at 150° , in CHBr_3 at 151° , in iodo-benzol at 190° , in oil of turpentine at 160° . Very sol. in benzaldehyde at 179° , in methylene iodide at 182° . (Reinders, Z. phys. Ch. 1900, 32. 506.)

1000 pts. oil of bitter almonds dissolve 4 pts. HgI_2 at ord. temp.; 1000 pts. olive oil, 4 pts.; 1000 pts. poppy oil, 10 pts.; 1000 pts. nut oil, 15 pts.; 1000 pts. castor oil, 20 pts.; 1000 pts. lard oil, 4.5 pts.; 1000 pts. vaseline, 2.5 pts.; 1000 pts. benzene, 4 pts. Sol. in phenol. (Mehn, Pharm. J. 3. 327; B. 19. 8 R.)

Solubility in aniline.

S = Temp. of solidification.

Mols. per 100	S	Mols. per 100	S	Mols. per 100	S
5.9	12°	19.9	48.5°	33.0	128°
8.2	22.5	25.8	53.5	35.6	140
10.3	29	29.3	105	37.5	147
14.9	41.5	31.7	122	39.2	156
16.6	45	32.4	(55)		

(Staronka, Anz. Ak. Wiss. Krakau, 1910. 372.)

Solubility of HgI_2 in aniline at t° .		
t°	g. HgI_2 per 100 g aniline	Solid phase
-8.5	23.35	$\text{HgI}_2 \cdot 2\text{C}_6\text{H}_5\text{NH}_2$
+0.4	28.69	"
17.8	42.85	"
21.1	47.55	"
26.9	55.47	"
30.1	62.05	"
36.2	75.80	"
42.9	96.49	"
46.8*	" + HgI_2 (red)
48.8	128.1	HgI_2 (red)
63.6	163.8	"
70.82	184.1	"
76.2	201.6	"
95.9	246.7	"
108.*	HgI_2 (red) + HgI_2 (yellow)
115.7	281.8	HgI_2 (yellow)
137.2	285.2	"
181.1	279	"
199.1	863.2	"

* Transition point.

(Pearce and Fry, J. phys. Ch. 1914, 18, 667.)

Very sol. in boiling alcoholic solution of aniline. (Vohl, Dissert. 1871.)

Abundantly sol. in hot benzonitrile and other aromatic nitriles. (Werner, Z. anorg. 1897, 15, 7.)

Sol. in benzonitrile (0.98 g. in 100 g. at 18°). 20 times more sol. by addition of 5 g. KI to 100 cc. benzonitrile. (Naumann, B. 1914, 47, 1375.)

Sol. in pyridine. (Naumann, B. 1904, 37, 4609.)

Solubility of HgI_2 in pyridine.

S = temp. of solidification.

Mols. per 100	S	Mols. per 100	S	Mols. per 100	S
5	10°	34.6	107°	51.3	93.5°
9.8	42.5	38.0	103	51.6	96
15.14	66.5	43.0	97	52.7	108
19.3	83	46.7	88.5	53.2	109
26.3	102.5	48.5	89	55.4	122
29.6	107	50.6	89	67.9	135

(Staronka, Anz. Ak. Wiss. Krakau, 1910, 372.)

Sp. gr. at $16^\circ/4^\circ$ of HgI_2 + pyridine containing 10.43% HgI_2 = 1.1482; containing 7.99% = 1.1053. (Schönrock, Z. phys. Ch. 1893, 11, 770.)

Solubility of HgI_2 in quinoline S = temp. of solidification.

Mols. per 100	S	Mols. per 100	S	Mols. per 100
4.7	100°	29.8	151°	43.0
9.1	115.5	31.4	153	46.1
13.2	133.5	35.4	156	48.8
23.1	138	37.7	160	49.5
26.7	145	41.6	165	54.4

(Staronka, Anz. Ak. Wiss. Krakau, 1910, 372.)

Mol. weight determined in methyl- and ethyl-sulphide. (W. anorg. 1897, 15, 20.)

More or less sol. at high temp. in (bpt. $160-230^\circ$), bromnaphthalene, toluidine and amyl alcohol. (Re phys. Ch. 1900, 32, 503.)

Yellow modification.

100 g. of sat. solution in acetone contain 3.0 g. HgI_2 . (Reinders, Z. 1900, 32, 514.)

Red modification.

Solubility in alcohol equals:

0.717-0.724 g. in 100 g. solution

1.044-1.084 g. " "

2.10-2.20 g. " "

(Reinders, Z. phys. Ch. 1900, 32, 514.)

100 g. of sat. solution in acetone contain 1.95 g. HgI_2 . (Reinders, Ch. 1900, 32, 514.)

HgI_2 is moderately sol. in abs. its b.-pt. The solution has a decolor. On cooling, yellow crystals out. They soon change to the red.

Readily sol. in hot amyl alcohol crystals separate from the solution on cooling.

Readily sol. in allyl alcohol, yellow solution, from which yellow crystals separate on cooling.

Sl. sol. in acetone, giving a yellow solution. On cooling yellow plates separate and rapidly turn red.

Sol. in phenol at 150°C . Solution low color and yellow crystals separate on cooling.

Readily sol. in boiling benzene. solution is yellow. The yellow crystals separate out on cooling, and change to the red.

Sol. in toluene giving yellow solution which yellow crystals separate out. They rapidly change to red.

Readily sol. in naphthalene at 100° above its transition point. Solution and on cooling yellow crystals separate.

Readily sol. in hot pseudo-cumene a yellow solution. On cooling yellow crystals separate.

Readily sol. in ethyl iodide at

yellow solution, from which yellow crystals separate on cooling, which change to red rapidly.

Only sl. sol. in ethyl bromide, giving yellow solution from which yellow crystals separate on cooling, which change rather slowly to the red.

Sparingly sol. in isopropyl bromide.

Moderately sol. in isobutyl bromide, giving pink solution from which yellow crystals separate on cooling, which change slowly to red.

Sl. sol. in ethylidene chloride. On sudden cooling at 18° the iodide crystallizes out in yellow plates, which quickly change to red.

Sparingly sol. in propyl chloride, giving a pink solution, from which yellow crystals separate on cooling.

Readily sol. in ethyl cyanide, giving a yellow solution. On cooling yellow crystals separate and rapidly change to red.

Moderately sol. in benzene cyanide, giving a deep yellow solution. On sudden cooling the solution deposits yellow crystals, which rapidly turn red.

Rapidly sol. in benzoic acid at high temperatures.

Sparingly sol. in ethyl acetate, giving yellow solution.

Sol. in ethyl propionate.

Very sol. in ethyl butyrate, giving a yellow solution. On cooling the iodide crystallized from the solution.

Sl. sol. in ethyl isobutyrate.

Readily sol. in methyl salicylate, giving a yellow solution.

Sparingly sol. in phenyl salicylate, giving yellow solution. On cooling yellow crystals separate out, which gradually change to red. (Kastle, Am. Ch. J. 1899, 22. 474.)

Mercuriomercuric iodide, $\text{Hg}_4\text{I}_6 = \text{Hg}_2\text{I}_2 \cdot 2\text{HgI}_2$.

Insol. in H_2O or alcohol. Partially sol. in $\text{KI} + \text{Aq}$, in hot NaCl , and $\text{NH}_4\text{Cl} + \text{Aq}$, and in hot $\text{HCl} + \text{Aq}$, though very slowly. (Boullay, A. ch. (2) 34. 345.)

Mercury periodide, HgI_4 .

Sol. in $\text{KI} + \text{Aq}$. Decomp. by cold H_2O or alcohol. (Jørgensen, J. pr. (2) 2. 347.)

Mercuric hydrogen iodide (Iodomeric acid), HI , $\text{HgI}_2 = \text{HHgI}_2$.

Crystallizes from $\text{HI} + \text{Aq}$. (Boullay.)

Easily decomp. (Neumann, M. 10. 236.)

$3\text{HgI}_2, 2\text{HI} + \text{H}_2\text{O}$. (François, Dissert. 1901.)

Mercuric nickel iodide, $\text{HgI}_2, \text{NiI}_2 + 6\text{H}_2\text{O}$.

Sol. in alcohol, ether, and acetone; not decomp. by H_2O . (Dobroserdoff, C. C. 1901, II. 332.)

$2\text{HgI}_2, \text{NiI}_2 + 6\text{H}_2\text{O}$. Hygroscopic; decomp. by H_2O ; sol. in acetone and ether. (Dobroserdoff, C. C. 1901, II. 332.)

Mercuric potassium iodide, $\text{HgI}_2, \text{KI} + 1\frac{1}{2}\text{H}_2\text{O}$.

Deliquescent (v. Bonsdorff). Permanent; decomp. by H_2O into 2KI , HgI_2 , and HgI_2 (Boullay); sol. in alcohol, ether, and conc. $\text{HC}_2\text{H}_3\text{O}_2$, but decomp. by other acids (Berthelot, J. Pharm. 14. 186). Sp. gr. of sat. solution in $\text{H}_2\text{O} = 2.4$ to 3.1 .

$+ \text{H}_2\text{O}$. Sol. in H_2O with decomp. Can be cryst. from alcohol. Very sl. sol. in dry ether. Very sol. in wet ether. (Marsh, Chem. Soc. 1910, 97. 2297.)

$\text{HgI}_2, 2\text{KI}$. Sol. in H_2O . (Thomsen and Bloxam, Chem. Soc. 41. 379.)

Sat. solution of $\text{KI} + \text{HgI}_2$ in H_2O at 22.9° contains 8.66% K, 22.49% Hg and 52.48% I, corresponding to 0.22 mol. K, 0.11 mol. Hg and 0.45 mol. I. (Duboin, C. R. 1905, 141. 385.)

Sp. gr. at 16°/4° of aqueous solution containing 12.2875% salt = 1.10148; containing 12.2371% = 1.1038; containing 7.9843% = 1.06491. (Schönrock, Z. phys. Ch. 1893, 11. 782.)

Sol. in methyl acetate. (Bezold, Dissert. 1906.)

Sol. in ethyl acetate. (Alexander, Dissert. 1899; Hamers, Dissert. 1906.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4328.)

Sol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3601.)

$+ 2\text{H}_2\text{O}$. Sol. in alcohol, ether and acetone; decomp. by H_2O . (Pawlow, C. C. 1901, I. 363.)

Solubility determinations show that KHgI_2 and $\text{KHgI}_2 + \text{H}_2\text{O}$ are the only double salts formed at 20°–30°. See $\text{HgI}_2 + \text{KI}$ under HgI_2 . (Dunningham, Chem. Soc. 1914, 105. 368.)

Mercuric rubidium iodide, HgI_2, RbI .

Sol. in alcohol; decomp. by H_2O .

$\text{HgI}_2, 2\text{RbI}$. Very easily sol. in H_2O . (Grossmann, B. 1904, 37. 1258.)

Very sol. in acetic acid and alcohol; decomp. by H_2O . Stable in aq. solution in the presence of an excess of RbI . (Erdmann, Arch. Pharm. 1894, 232. 30.)

Mercuric silver iodide, $\text{HgI}_2, 2\text{AgI}$.

(Wegelius and Kilpi, Z. anorg. 1909, 61. 416.)

Mercuric sodium iodide, HgI_2, NaI .

Deliquescent, and decomp. by much H_2O . (v. Bonsdorff, Pogg. 17. 266.)

Sol. in alcohol; decomp. by H_2O .

$\text{HgI}_2, 2\text{NaI}$. Deliquescent; sol. in H_2O and alcohol. (Boullay.)

Sat. solution of $\text{NaI} + \text{HgI}_2$ in H_2O at 24.75° contains 4.59% Na, 25% Hg, and 58.25% I, corresponding to 0.20 mol. Na,

0.12 mol. Hg, and 0.45 mol. I. (Duboin, C. R. 1905, 141. 385.)

+4H₂O. Extremely deliquescent. (Duboin, C. R. 1906, 143. 314.)

Mercuric strontium iodide, HgI₂, SrI₂ (?).

Sol. in H₂O without decomp. (Boullay.)

+8H₂O. As Ca salt. (Duboin, C. R. 1906, 142. 573.)

2HgI₂, Srl₂ (?). Decomp. by much H₂O into sol. HgI₂, Srl₂ and insol. HgI₂. (Boullay.)

Mercuric thorium iodide, 5HgI₂, ThI₄+18H₂O.

Very deliquescent. Easily decomp. by H₂O. (Duboin, A. ch. 1909, (8) 16. 282.)

5HgI₂, 2ThI₄+21H₂O. (Duboin.)

2HgI₂, ThI₄+12H₂O. (Duboin.)

Mercuric zinc iodide.

Deliquescent. Decomp. by H₂O. (v. Bonsdorff.)

Mercuric iodide ammonia, HgI₂, 2NH₃.

Decomp. by NH₃ giving NH₄I and NH₄I. (François, C. R. 1900, 130. 333.)

Stable only in the presence of excess of ammonia. Gives off NH₃ in the air. (François, J. Pharm. 1897, (6) 5. 388; C. C. 1897, I. 1088.)

Mercuric iodide hydrazine, HgI₂, N₂H₄.

Decomp. by H₂O. (Hofmann and Marburg, A. 1899, 305. 215.)

Mercuric iodide rubidium bromide,

HgI₂, 2RbBr.

Decomp. by H₂O.

Sol. in alcohol without decomp. (Grosman, B. 1903, 36. 1603.)

Mercuric iodide silver chloride, HgI₂, 2AgCl.

Insol. in H₂O. (Lea, Sill. Am. J. (3) 7. 34.)

Mercury iodoantimonide, Hg₃Sb₄2HgI₂.

Sol. in HNO₃, aqua regia and hot H₂SO₄; insol. in HCl. (Granger, C. R. 1901, 132. 1116.)

Mercury nitride, Hg₃N₂.

Gradually decomp. by H₂O. Decomp. by conc. HNO₃, or HCl+Aq. (Hirzel, J. B. 1852. 419.)

Not attacked by cold, but decomp. by hot dil. H₂SO₄.

Sol. in acids+Aq.

Sol. in ammoniacal solutions of ammonium salts.

Insol. in excess of KNH₃. (Franklin, Z. anorg. 1905, 46. 18.)

Sol. in ammonia solutions of ammonium salts and in aq. acid solutions.

Very explosive. (Franklin, J. An Soc. 1905, 27. 835.)

HgN₂.

See Mercurous azoimide.

HgN₄.

See Mercuric azoimide.

Mercurous oxide, Hg₂O.

Insol. in H₂O. Insol. in dil. HCl +Aq. Sol. in warm conc. HC₂H₃O₂. Sol. in 150,000 pts. H₂O. (Bh anorg. 1897, 13. 410.)

Decomp. by H₂O or weak bases (NH₄)₂CO₃+Aq (Wittstein), K (Rose), KI+Aq (Berthelot), NH₄Cl+Aq (Pagenstecher) into Hg, or HgCl₂, etc.

Sl. decomp. by alkali chlorides formation of HgCl₂, which (Miahle.)

Sl. sol. in alkali cyanides+Aq

Insol. in KOH, and NaOH+Aq

Insol. in liquid NH₃. (Franklin J. 1898, 20. 829.)

Insol. in alcohol and ether.

Mercuric oxide, HgO.

Sol. in 20,000 to 30,000 pts. H₂O C. R. 41. 509.)

Sol. in 200,000 pts. H₂O. (W Gaz. 1858. 345.)

Ordinary coarse HgO is sol. to the extent of 50 mg. per l. at 25° finely powdered the solubility is 150 mg. per l. (Hulett, Z. phys. 37. 406.)

Red modification is:—

Sol. in 19,500 pts. H₂O at 25°; in H₂O at 100°. (Schick, Z. phys. 42. 172.)

1 l. H₂O dissolves 50 mg. red n of HgO at 25°. (Hulett, Z. phys. 37. 406.)

Yellow modification is:—

Sol. in 19,300 pts. H₂O at 25°; in at 100°. (Schick, Z. phys. Ch. 172.)

Sol. in acids. Insol in H₃PO₄, or Aq. (Haack, A. 262. 190.)

Scarcely attacked by H₂C₂O₄, + lon, A. ch. (3) 18. 352.)

Solubility of HgO in HF at
Hg = g.-atoms Hg in 1 l. of the s

HF normal	Hg
0.12	0.012
0.24	0.024
0.57	0.062
1.11	0.116
2.17	0.268

(Jaeger, Z. anorg. 1901, 27.

Solubility of HgO in HF is decreased by the addition of KF, which proves the non-existence of complex fluorides. (Jaeger.)

Insol. in H_3AsO_4 , H_3PO_4 , and in primary and secondary alkali salts of these acids. (Haack, A. 1891, 262. 190.)

Sol. in hot $NH_4Cl + Aq$, less in $NH_4NO_3 + Aq$. (Brett.)

Insol. in KOH, or NaOH + Aq.

Decomp. by alkali chlorides + Aq into $HgCl_2$, which dissolves. (Miahle, A. ch. 1887, 177.)

Sol. in $Fe(NO_3)_3$, and $Bi(NO_3)_3 + Aq$ with evolution of oxides. **Sol.** in KI + Aq. (Persoz.)

Very sol. in acid sulphites + Aq. (Barth, phys. Ch. 1892, 9. 192.)

Completely sol. in conc. $CaCl_2$, $BaCl_2$, $LiCl$, and $SrCl_2 + Aq$. (André, C. R. 1887, 431.)

Solubility in Ag salts + Aq. 100 g. Ag_2SO_4 in aqueous solution dissolve 13 g. HgO. **Solubility in $AgNO_3 + Aq$** is 15.6 : 100; in $Ag acetate + Aq$ is 1.137 : 100. (Finci, Gazz. ch. it. 1911, 41. (2) 545.)

Much less sol. in KCl and NaCl + Aq than in H_2O . (Schoch.)

Sol. in $U(NO_3)_6$, $Al(NO_3)_3$, and $Fe(NO_3)_3 + Aq$. (Mailhe, A. ch. 1902, (7) 27. 373.)

Very sl. sol. in cold $Hg(CN)_2 + Aq$, abundantly sol. at 75° with evolution of HCN. (Barthe, J. Pharm. 1896, (6) 3. 183.)

Sol. in cold or hot alcoholic NH_4SCN in large amounts. (Fleischer, A. 1875, 179. 225.)

Completely sol. in KI + Aq. (Jehn, Arch. Pharm. 1873, 201. 97.)

Solubility of red or yellow modification in N/50 KCl + Aq is about 25% greater than in pure H_2O . (Schick, Z. phys. Ch. 1903, 42. 168.)

Insol. in liquid HF. (Franklin, Z. anorg. 1905, 46. 2.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Sol. in alcoholic solution of hydroxylamine hydrobromide below 0°. (Adams, Am. Ch. J. 1902, 28. 216.)

Insol. in alcohol.

Sol. in trichloroacetic acid + Aq. (Brand, J. pr. 1913, (2) 88. 342.)

Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

When freshly pptd., is insol. in acetone + Aq. even on warming, but easily sol. if liquid is made alkaline by NaOH. **Insol.** in acetophenone even after long warming at 100°. **Sol.** in acetaldehyde and much H_2O and a little NaOH. (Auld and Hantzsch, B. 1905, 38. 2680.)

Sol. in formamide. (Fischer, Arch. Pharm. 1894, 232. 329.)

Very sol. in ethylene diamine. For 1 mol.

HgO, 7-10 mols. ethylene diamine are necessary. (Traube and Löwe, B. 1914, 47. 1910.)

Easily sol. in benzamide. (Dessaigues, A. ch. 1852, (3) 34. 146.)

When freshly pptd., is sol. in picric acid + Aq. (Varet, C. R. 1894, 119. 560.)

Sol. in alkaline solution of phenol disulphonic acid. (Lumière and Chevrotier, C. R. 1901, 132. 145.)

Sol. in nucleic acid + Aq when freshly pptd. (Schweckerath, Pat. 1899.)

Sol. in gum arabic + Aq. (Peschier, J. Pharm. 1896, (6) 3. 509.)

Mercuric oxybromide, $HgBr_2$, HgO.

(André, A. ch. (6) 3. 123.)

$HgBr_2$, 2HgO. (André.)

$HgBr_2$, 3HgO. (a) Yellow. **Insol.** in cold, sl. sol. in hot H_2O . **Easily sol.** in alcohol. (Löwig.)

(b) Brown. **Insol.** in alcohol. (Rammelsberg, Pogg. 55. 248.)

$HgBr_2$, 4HgO. (André.)

Insol. in ord. solvents. **Decomp.** by alkalies and acids. (Fischer and von Wartenburg, Ch. Z. 1902, 26. 894.)

2HgBr₂, 7HgO. **Readily decomp.** by acids and alkalies. (Fischer and von Wartenburg.)

Mercurous oxychloride, Hg_2O , 2HgCl.

Min. *Eglestonite*.

Decomp. by hot HCl and by HNO_3 . (Moses, Am. J. Sci, 1903, (4) 16. 253.)

Mercuric oxychloride,

HgO, $HgCl_2$. **Less sol.** than $HgCl_2$, but not isolated. (Thümmel.) **Decomp.** by cold H_2O . (André, A. ch. (6) 3. 118.)

HgO, 2HgCl₂. **Decomp.** by warm H_2O or cold alcohol into 2HgO, $HgCl_2$. (Thümmel, Arch. Pharm. (3) 27. 589.)

Decomp. by H_2O . **Not decomp.** by alcohol. (Arctowski, Z. anorg. 1895, 9. 178.)

2HgO, $HgCl_2$. *Two modifications.*

A. Red. **Insol.** in H_2O ; **decomp.** by alkali carbonates, or chlorides + Aq into 4HgO, $HgCl_2$.

Acted upon by cold alkali carbonates and alkali chlorides + Aq. (Schoch, Am. Ch. J. 1903, 29. 335.)

Not decomp. by H_2O at ord. temp. (Thümmel.)

Very sl. sol. in cold, **completely sol.** in hot H_2O . (Haack, A. 1891, 262. 189.)

A small amt. of HNO_3 converts it into a white powder; more HNO_3 dissolves it. (Haack, A. 1891, 262. 189.)

B. Black. **Not decomp.** by alkali chlorides, or carbonates + Aq. (Thümmel.)

Not affected by boiling alkali carbonates or alkali chlorides + Aq. (Schoch, Am. Ch. J. 1901, 29. 335.)

Insol. in cold and hot H_2O and alcohol.

Sol. in acid. (Van Nest, Dissert. 1909.)

Not changed by H_2O . (Blaas, Miner. Mitt. (2) 2. 177.)

$\text{H}_2\text{SO}_4 + 66.7\%$ 4-N HNO_3 , and H_2SO_4 alone. (Moore, J. Am. Chem. Soc. 11, 33. 1094.)

H_2SO_4 does not attack red or white, but they are attacked by hot acid. (Schneider, Pogg. 127. 488.)

ch. 1898, (7) 14. 198.)
d. HgS is insol. in dil. $\text{KCN} + \text{H}_2\text{O}$.

+Aq, but readily only in presence of alkali. (Brunner, Pogg. 15. 596.)
ng $\text{KOH} + \text{Aq}$.

[or $\text{NaSH} + \text{Aq}$. Very sl. sol. in $(\text{NH}_4)_2\text{S} + \text{Aq}$. Insol. in $\text{KCN} + \text{H}_2\text{O}$. (Fresenius.)

in conc. Na_2S or $\text{K}_2\text{S} + \text{Aq}$, or in conc. KOH or NaOH . Insol. in H_2O . Sol. in CaS , BaS , or $\text{SrS} + \text{Aq}$.

or $\text{KSH} + \text{Aq}$. (de Koninck, 1891. 51.)

1 NaSH is very small in comparison with that in $\text{Na}_2\text{S} + \text{Aq}$. (Knox, Trans. Faraday Soc. 1908, 4. 30.)

1 BaS is practically equal to that in $\text{Na}_2\text{S} + \text{Aq}$. (Knox.)

modifications are sol. in conc. $\text{Na}_2\text{S} + \text{Aq}$. (Allen and Smith, J. Sci. 1912, (4) 34. 368.)

potassium thiocarbonate + Aq. (Smith, J. Sci. 1912, (4) 34. 368.)

anal. 26. 15.)
ali sulpho-molybdates, -tungstates, -arsenates, -antimonates, -vanadates. (Storch, B. 16. 2015.)

+Aq containing 50 g. Ba dissolves in the cold, but 50-60 g. at 100°C.

liquid NH_3 . (Gore, Am. Ch. J. 1899, 21. 109.)

stone. (Eidmann, C. C. 1899, 1. 109.)

pyridine. (Schroeder, Dissert. 1906, 1. 109.)

ethyl acetate. (Naumann, B. 1899, 1. 109.)

colloidal state, sol. in H_2O . (Bull. Soc. (2) 49. 452.)

bar. Insol. in H_2O , alcohol, or alkaline solutions.

y hot dil. $\text{HNO}_3 + \text{Aq}$. Not sol. in $\text{HCl} + \text{Aq}$, but easily by hot aqua regia. Easily sol. in $\text{CuCl}_2 + \text{Aq}$.

.)
mixture of Na_2S and NaOH when proportion of $\text{HgS} : 2\text{Na}_2\text{S}$.

or $\text{Na}_2\text{S} + \text{Aq}$ or in mixtures of H_2S and NaOH .

Insol. in cold $\text{NaSH} + \text{Aq}$, but on warming with evolution of H_2S . (Sill. Am. J. (3) 33. 199.)

stone. (Krug and M'Elroy.)

easily sol. in 20% $\text{HBr} + \text{Aq}$. (Smith, J. Am. Chem. Soc. 1896, 18. 109.)

er. (Smith, J. Am. Chem. Soc. 1896, 18. 109.)

er. (Smith, J. Am. Chem. Soc. 1896, 18. 109.)

mercuric sulphide.

mercuric sulphate, mercuric.

Mercuric potassium sulphide, $\text{K}_2\text{S}, 2\text{HgS}$.

Decomp. into its constituents by H_2O ; decomp. by HCl , and $\text{HNO}_3 + \text{Aq}$, and by hot KOH , and $\text{NH}_4\text{OH} + \text{Aq}$. (Schneider, Pogg. 127. 488.)

$\text{K}_2\text{S}, \text{HgS} + 5\text{H}_2\text{O}$. Decomp. by H_2O or alkalis. (Weber, Pogg. 97. 76.)

+ H_2O . (Ditte.)
+ $7\text{H}_2\text{O}$. Sol. in $\text{K}_2\text{S} + \text{Aq}$. (Ditte, C. R. 98. 1271.)

$\text{K}_2\text{S}, 5\text{HgS} + 5\text{H}_2\text{O}$. Easily decomp. by H_2O . (Ditte.)

Mercuric sodium sulphide, $\text{HgS}, \text{Na}_2\text{S} + 8\text{H}_2\text{O}$.

Decomp. by H_2O or alkalis.
 $5\text{HgS}, 2\text{Na}_2\text{S} + 3\text{H}_2\text{O}$. Decomp. by H_2O . (Knox, Trans. Faraday Soc. 1908, 4. 36.)

(Knox, Trans. Faraday Soc. 1908, 4. 36.)

Mercuric sulphobromide, $2\text{HgS}, \text{HgBr}_2$.

Insol. in H_2O . Not attacked by boiling HNO_3 or H_2SO_4 . (Rose.)

Mercuric sulphochloride, $2\text{HgS}, \text{HgCl}_2$.

Insol. in H_2O , cold or hot, dil. or conc. HNO_3 , H_2SO_4 , or $\text{HCl} + \text{Aq}$. (Rose, Pogg. 13. 59.)

Decomp. by hot aqua regia.
By boiling with dil. HNO_3 , H_2SO_4 , and HCl , Hg and Cl go into solution. (Hamers, Dissert. 1906.)

Insol. in H_2O and H_2SO_4 . Partly sol. in HCl and HNO_3 ; easily sol. in aqua regia. (Alexander, Dissert. 1899.)

Sol. in aqua regia. (Denigès, Bull. Soc. 1915, (4) 17. 356.)

$3\text{HgS}, \text{HgCl}_2$. Properties as the above comp. (Poleck and Goercki, B. 21. 2415.)

$4\text{HgS}, \text{HgCl}_2$. As above. (P. and G.)
 $5\text{HgS}, \text{HgCl}_2$. As above. (P. and G.)

Insol. in alkali sulphides and in fuming HNO_3 ; decomp. by $\text{NaOBr} + \text{Aq}$ and by KOH . (Bodroux, C. R. 1900, 130. 1399.)

Sl. sol. in solutions of alkali sulphides unless heated. (Berzelius.)

Easily sol. in alkali sulphides + Aq; slowly sol. in alkalis or alkali hydrosulphides + Aq. (Atterberg, J. B. 1873. 258.)

Mercurous sulphotetrachloride, Hg_2SCl_4 .

Decomp. by H_2O with separation of S , HgCl_2 going into solution. (Capitaine, J. Pharm. 25. 525.)

Mercuric sulphofluoride, $2\text{HgS}, \text{HgF}_2$.

Decomp. by boiling H_2O . Not decomp. by hot HCl or $\text{HNO}_3 + \text{Aq}$, but gives HF with hot $\text{H}_2\text{SO}_4 + \text{Aq}$. (Rose, Pogg. 13. 66.)

Mercury sulphodithionide, $\text{HgN}_2\text{S}, \text{NH}_3$.

Ppt. (Ruff, B. 1904, 37. 1585.)

Mercuric sulphoiodide, HgS, HgI_2 .

Ppt. (Rammelsberg, Pogg. 48. 175.)
 $2\text{HgS}, \text{HgI}_2$. (Palm, C. C. 1863. 121.)

Insol. in min. acids with exception of aqua regia. (Hamers, Dissert. 1906.)

Mercuric sulphoiodide ammonia, 2HgS , HgI_2 , NH_3 .
(Foerster, Ch. Z. 1895, 19. 1895.)

Mercuric telluride, HgTe .

Min. *Coloradoite*. Sol. in boiling HNO_3 + Aq with separation of H_2TeO_3 .

Metastannic acid.

See Stannic acid.

Molybdatoiodic acid.

See Molybdoiodic acid.

Molybdenum, Mo.

Not attacked by HCl , HF , or dil. H_2SO_4 + Aq. Sol. in conc. H_2SO_4 . Very easily sol. in aqua regia. Oxidised by HNO_3 + Aq either to molybdenum oxide, which dissolves in HNO_3 , or, if HNO_3 is in excess, to molybdic acid, which remains undissolved.

Attacked by HNO_3 + Aq containing 3–70% HNO_3 , but only slowly by 70% acid, with formation of insol. white powder; much more vigorously by 50% acid, in which case a clear solution is formed. (Montemartini, Gazz. ch. it. 22. 384.)

Not attacked by alkalis + Aq. (Bucholz, Scher. J. 9. 485.)

With a sp. gr. 9.01, the metal is malleable and sol. in a mixture of HF and HNO_3 ; sol. in fused KClO_3 . (Moissan, Bull. Soc. 1895, (3) 13. 966.)

Ductile Mo is moderately quickly attacked by HNO_3 , H_2SO_4 and HCl . (Fink, Met. Chem. Eng. 1910, 8. 341.)

Not immediately attacked by cold dil. HNO_3 . Not attacked by dil. and conc. H_2SO_4 . Boiling dil. HCl + Aq does not attack; conc. dissolves traces by long heating. Sol. in aqua regia. (Lederer, Dissert. 1911.)

Dil. HCl dissolves 20.3% Mo at 110° in 18 hrs. More slowly sol. in HCl (sp. gr. 1.15).

Insol. in dil. H_2SO_4 at 110° . Slowly sol. in conc. H_2SO_4 (sp. gr. 1.82) at 110° , rapidly sol. at 200° – 250° .

Slowly sol. in conc. HNO_3 (sp. gr. 1.40), rapidly sol. in dil. HNO_3 (sp. gr. 1.15).

Rapidly sol. in hot aqua regia. Insol. in hot or cold HF . (Ruder, J. Am. Chem. Soc. 1912, 34. 388.)

Insol. in KOH + Aq. Sol. in fused KOH . (Ruder, J. Am. Chem. Soc. 1912, 34. 389.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Molybdenum acichloride.

See Molybdenyl chloride.

Molybdenum amide, $\text{OH}.\text{MoO}_3.\text{NH}_3$.

Very unstable. Insol. in aba. (Fleck, Z. anorg. 1894, 7. 353.)

Molybdenum amide nitride, Mo_2N_2 , 4MoN_3 , $\text{Mo}(\text{NH}_2)_3$.

Not attacked by HCl , or dil. HN (Uhrlaub.)

Molybdenum amidochloride, $\text{Mo}_2(\text{NH})$

Insol. in H_2O and dil. acids. (Roe Z. anorg. 1905, 46. 317.)

Molybdenum amidochloride ammoni
 $\text{Mo}_2(\text{NH}_2)_2\text{Cl}_2$, 10NH_3 .

Unstable in the air. (Rosenheim, Z. 1905, 46. 319.)

Molybdenum boride, Mo_2B_4 .

Moderately attacked by hot con and vigorously by hot aqua regia. and Moody, Chem. Soc. 1902, 81. 17

Molybdenum dibromide, $\text{MoBr}_2 = \text{Mo}$

See Bromomolybdenum bromide.

Molybdenum tribromide, MoBr_3 .

Not decomp. by H_2O . Boiling co and cold dil. HNO_3 + Aq do not attack. Dil. alkalies act slowly, but with separation of Mo_2O_3 on boiling. strand, J. pr. 82. 435.)

Molybdenum tetrabromide, MoBr_4 .

Rapidly deliquescent, and easily H_2O . (Blomstrand, J. pr. 82. 433.)

Molybdenum bromochloride, etc.

See Bromomolybdenum chloride, (

Molybdenum bronze.

See Molybdate molybdenum oxide

Molybdenum carbide, Mo_2C .

Insol. in HNO_3 . (Moissan, B 1895, (3) 13. 967.)

MoC . Does not decomp. H_2O 500–600°. Slowly attacked by hot and hot conc. H_2SO_4 . Easily dec HNO_3 . Not attacked by NaOH KOH + Aq. (Moissan and Hoffman 1904, 138. 1559.)

Molybdenum carbonyl, $\text{Mo}(\text{CO})_6$.

Quickly attacked by bromine. ether or benzene. (Mond, Hirts and Chem. Soc. 1910, 97. 808.)

Molybdenum dichloride, $\text{MoCl}_2 = \text{M}$

See Chloromolybdenum chloride.

Molybdenum trichloride, MoCl_3 .

Insol. in H_2O or boiling conc. $\text{HCl} + \text{Aq.}$ Slightly sol., especially when heated, in HNO_3 . Aq. Sol. in H_2SO_4 . Decomp. by NH_4OH , KOH , or $\text{NaOH} + \text{Aq.}$

Sl. sol. in alcohol. (Leichti and Kempe.)

Practically insol. in alcohol and ether. (Hampe, Ch. Z. 1888, 12. 5.)

Molybdenum tetrachloride, MoCl_4 .

Deliquescent. Hisses with little H_2O , but partly sol. in more H_2O . Only sl. sol. in conc. $\text{HCl} + \text{Aq.}$ Sol. in H_2SO_4 or $\text{HNO}_3 + \text{Aq.}$ Partly sol. in alcohol and ether. (Liechti and Kempe.)

Molybdenum pentachloride, MoCl_5 .

Very deliquescent. Sol. in H_2O with extreme evolution of heat. Sol. in HCl , HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq.}$

When freshly prepared, is incompletely sol. in H_2O , but after standing is easily sol. with hissing. (Kalischer, Dissert. 1902.)

Sol. in a small amt. of conc. HCl . (Hampe, Ch. Z. 1888, 12. 5.)

Sol. in absolute alcohol or ether. (Liechti and Kempe.)

Sol. in CHCl_3 and in CCl_4 . Sol. with hissing in many organic solvents (ethers, alcohols, ketones, aldehydes, acids, acid esters, acid anhydrides, amines). Sol. in cinnamic aldehyde. (Kalischer, Dissert. 1902.)

Molybdenum hydroxyl chloride, $\text{Mo}(\text{OH})_2\text{Cl}_2$.

Easily sol. in H_2O . (Debray, C. R. 46. 1101.)

Molybdenum tetrachloride phosphorus pentachloride, $\text{MoCl}_4, \text{PCl}_5$.

Sol. in H_2O .

$\text{MoCl}_4, 2\text{PCl}_5$. Sol. in H_2O . (Cronander, Bull. Soc. (2) 19. 500.)

Molybdenum phosphorus pentachloride, $\text{MoCl}_5, \text{PCl}_5$.

Easily decomp. (Smith and Sargent, Z. anorg. 1894, 6. 385.)

Molybdenum phosphoryl chloride, $\text{MoCl}_5, \text{POCl}_3$.

Decomp. by H_2O ; insol. in CS_2 ; sol. in C_6H_6 and CHCl_3 .

Molybdenum trichloride potassium chloride.

Efflorescent. Decomp. with H_2O . (Berzelius.)

$\text{MoCl}_3, 3\text{KCl}$. Very sol. in H_2O . Nearly insol. in alcohol and ether. (Chilesotti, C. C. 1903, II. 652.)

$+2\text{H}_2\text{O}$. Fairly easily sol. in cold H_2O without any apparent decomp. Decomp. in aqueous solution, slowly in the cold but rapidly on boiling. This decomp. is prevented by the presence of HCl .

Sl. sol. in conc. HCl . (Henderson, Proc. Chem. Soc. 1903, 19. 245.)

Molybdenum rubidium chloride, $\text{Rb}_2\text{MoCl}_6 + \text{H}_2\text{O}$.

Sol. in H_2O . Nearly insol. in alcohol and ether. (Chilesotti, C. C. 1903, II. 652.)

Molybdenum pentachloride nitrogen sulphide, $\text{MoCl}_5, \text{N}_4\text{S}_4$.

Decomp. in moist air. (Davis, Chem. Soc. 1906, 89, (2) 1575.)

Molybdenum hexafluoride, MoF_6 .

Decomp. by a little H_2O with separation of blue oxide. Sol. in large amount of H_2O forming a colorless solution.

Absorbed by alkalies and $\text{NH}_4\text{OH} + \text{Aq.}$ (Ruff, B. 1907, 40. 2930.)

Molybdenum fluoride with MF.

See Fluomolybdate, M.

Molybdenum potassium trifluoride (?).

Precipitate. Sol. in $\text{HCl} + \text{Aq.}$

Molybdenum potassium tetrafluoride (?).

Sl. sol. in H_2O . (Berzelius.)

Molybdenum sesquihydroxide, $\text{Mo}_2\text{O}_5\text{H}_3$.

Difficultly sol. in acids. Insol. in KOH , NaOH , NH_4OH , or $\text{K}_2\text{CO}_3 + \text{Aq.}$ Somewhat sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ but pptd. on boiling. (Berzelius.)

Molybdenum hydroxide, $\text{Mo}_2\text{O}_5, 5\text{H}_2\text{O}$.

Easily sol. in H_2O . Insol. in CaCl_2 , NH_4Cl , or $\text{NaCl} + \text{Aq.}$ Sl. sol. in alcohol. (Berzelius.)

Molybdenum dihydroxide, $\text{MoO}_3, x\text{H}_2\text{O}$.

Slowly and not abundantly sol. in H_2O , from which it is precipitated by NH_4Cl and other salts. Gelatinises by standing in closed vessels or by evaporating on the air. Sol. in the ordinary acids. Insol. in KOH , or $\text{NaOH} + \text{Aq.}$ Sol. in alkali carbonates $+ \text{Aq.}$

Molybdenum diiodide, MoI_2 .

Insol. in H_2O and alcohol. Sl. attacked by cold H_2SO_4 or HNO_3 . (Guichard, A. ch. 1901, (7) 23. 567.)

Sl. decomp. H_2O at ordinary temp. Slowly sol. in H_2SO_4 and HNO_3 . (Guichard, C. R. 1896, 123. 822.)

Molybdenum tetraiodide (?).

Completely sol. in water. (Berzelius.)

Molybdenum nitride, Mo_3N_3 , and Mo_3N_4 .

(Uhrlaub.)

See Molybdenum amide.

Mo_3N_2 . (Rosenheim, Z. anorg. 1905, 46. 317.)

Molybdenum monoxide, MoO.

Known only as hydroxide. (Blomstrand, J. pr. 77. 90.)

Molybdenum sesquioxide, Mo₂O₃.

Insol. in acids or alkalies.

See **Molybdenum sesquihydroxide**.

Molybdenum dioxide, MoO₂.

Insol. in HCl or HF + Aq. Sl. sol. in conc. H₂SO₄. HNO₃ oxidises to MoO₃. Not attacked by KOH + Aq. (Ullik, A. 144. 227.)

Sl. sol. in KHC₄H₄O₆ + Aq.

Molybdenum trioxide, MoO₃.

Sol. in 500 pts. cold, and much less hot H₂O. (Bucholz.)

Sol. in 960 pts. hot H₂O. (Hatchett.)

Sol. in 570 pts. cold, and much less hot H₂O. (Dumas.)

Sol. in acids before ignition. Insol. in acids, but sl. sol. in acid potassium tartrate + Aq after ignition. Sol. in alkalies or alkali carbonates + Aq.

Sol. in NH₄OH + Aq.

See also **Molybdic acid**.

Min, *Molybite*. Sol. in HCl + Aq.

Molybdenum oxide, Mo₂O₅.

Sol. in H₂SO₄ and HCl; only sl. sol. in H₂SO₄. (Klason, B. 1901, 34. 151.)

+ 3H₂O. Sl. sol. in H₂O (2 g. in 1 l.). Insol. in NH₄Cl + Aq. Insol. in caustic alkalies, somewhat sol. in NH₄OH. Much more sol. in M₂CO₃ + Aq and in (NH₄)₂CO₃ + Aq. (Klason, B. 1901, 34. 150.)

Mo₄O₁₀ + 3H₂O. (Smith and Oberholtzer, Z. anorg. 1893, 4. 243.)

Mo₄O₁₁ + 6H₂O. Sol. in H₂O. (Bailhache, C. R. 1901, 133. 1212.)

Mo₄O₁₄ + 6H₂O. Very sol. in H₂O. (Guichard, C. R. 1900, 131. 419.)

Mo₇O₂₀. Sol. in H₂O. (Junius, Z. anorg. 1905, 46. 447.)

Mo₃₀O₄₁ + 21H₂O = Mo₂O₅, 18MoO₃ + 21H₂O. Easily sol. in H₂O. Insol. in NH₄Cl + Aq. (Klason, B. 1901, 34. 160.)

Mo₂₈O₇₇ + 24H₂O = Mo₂O₅, 24MoO₃ + 24H₂O. (Klason, B. 1901, 34. 159.)

3Mo₂O₃, 2Mo₇O₂₄ + 18H₂O. Sol. in H₂O. (B.)

It is probable that the five blue oxides of molybdenum described by Klason (B. 34, 148, 158) and Bailhache are either the blue oxide Mo₅O₁₄ prepared by the author or mixtures of this compd. with molybdenum trioxide. (Guichard, C. R. 1902, 134. 173.)

Mo₅O₁₂. Not attacked by ammonia; easily oxidised by HNO₃ + Aq. Not attacked by HCl or H₂SO₄ + Aq. (Wöhler, A. 110. 275.)

Formula is Mo₅O₈, according to Wöhler, but Muthmann (A. 238. 108) has shown that correct formula is Mo₅O₁₂.

Not attacked by boiling alkalies, HCl, or dil. H₂SO₄ + Aq. Sol. in conc. H₂SO₄, with

subsequent decomp. Sol. in aqua regia Cl₂ + Aq. (Muthmann.)

Mo₃O₈. Sol. in H₂O. (Muthmann, 238. 108.)

Min. *Ilsemannite* (?).

+ 5H₂O. Moderately sol. in H₂O. chetti, Z. anorg. 1899, 19. 393.)

Mo₅O₇. (v. d. Pfordten, B. 15. 192)

Molybdenum trioxide ammonia, MoO₃.

Unstable in air. Very sol. in H₂O. evolution of ammonia. (Rosenh. anorg. 1906, 50. 303.)

3MoO₃, NH₃ + 1/2 H₂O. True comp. of commercial molybdic acid. (Klason, 1901, 34. 156.)

NH₄H₂Mo₃O₁₂. Very sl. sol. in cold sol. in hot H₂O with partial decomp. (B. 1901, 34. 156.)

3MoO₃, 3NH₃ + 7H₂O = (NH₄)₃Mo₃O₁₂ + 4H₂O. True composition of Ramme's salt. (Klason, 1901, 34. 155.)

4MoO₃, NH₃ + 6H₂O. Very sl. sol. very easily sol. in hot H₂O. An indication with less H₂O gradually c. (Mylius, B. 1903, 36. 639.)

4MoO₃, 2NH₃ + 3H₂O. (Klason, 34. 156.)

6MoO₃, 3NH₃ + 5H₂O. Very sl. sol. cold, more easily sol. in hot H₂O, with decomp. (Klason, B. 1901, 34. 156)

12MoO₃, 3NH₃ + 12H₂O. (Klason, 1901, 34. 158.)

12MoO₃, 3NH₃, 12H₂O + 3MoO₃ = 15MoO₃, 3NH₃ + 6H₂O. Insol. (Klason.)

4MoO₃, MoO₃, 2NH₃ + 7H₂O. S. in H₂O; fairly stable, gradually decomp. in dil. acids. (Hofmann, Z. anorg. 280.)

Molybdenum trioxide ammonia peroxide, 18MoO₃, 14NH₃, 18H₂O.

Sol. in H₂O. Sp. gr. of sat. sol. 1.486 at 17.4°. (Baerwald, B. 1884, 10. 100.)

Molybdenum oxybromide.

See **Molybdenyl bromide**.

Molybdenum oxychloride.

See **Molybdenyl chloride**.

Molybdenum oxyfluoride.

See **Molybdenyl fluoride**.

Molybdenum oxyfluoride with MF₃.

See **Fluoxymolybdate, M, and Flumolybdate, M.**

Molybdenum phosphide, Mo₂P₃.

Gradually sol. in hot HNO₃ + Aq. (Rautenberg, A. 100. 374.)

selenide, MoSe_2 .

Prepared pure. (Uelsmann, A. 116.)

silicide.

only very sl. sol. in other acids. (N. 1898, 78. 319.)

sol. in all min. acids; sol. in a mixture of $\text{HF} + \text{HNO}_3$. (Defacqz, 44. 1425.)

insol. in acids; sol. in $\text{HF} + \text{HNO}_3$ by 10–20% $\text{KOH} + \text{Aq.}$ Decomposed by NaOH . (Hönigschmid, M. 1.)

Decomposed by boiling HNO_3 , aqua regia attacks, Trans. Am. Electrochem. 106.)

Figouroux, C. R. 129. 1238.)

disulphide, MoS_2 .

Insol. in H_2O . Easily sol. in aqua regia. Decomposed by HNO_3 . Sol. in boiling water attacked by $\text{KOH} + \text{Aq.}$ (Ber-

molybdenite. Sol. in $\text{HNO}_3 + \text{Aq.}$, decomposition of MoO_3 ; sol. in aqua regia; insol. in H_2SO_4 .

trisulphide, MoS_3 .

sol. in H_2O , especially if hot, but insol. in cold. Difficultly sol. except when $\text{KOH} + \text{Aq.}$ Sl. sol. in solutions of alkalis unless heated. (Berzelius.) Insol. in alkali sulphides + Aq. ; slowly decomposed by alkali hydrosulphides + Aq. (B. 1873. 258.)

tetrasulphide, MoS_4 .

Prepared by hot H_2O or acids. Insol. in cold alkali sulphides + Aq. , but decomposed by boiling. (Berzelius.) Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1.)

sesquisulphide, Mo_2S_3 .

Insol. in Cl and H_2SO_4 ; sol. in hot conc. aqua regia. (Guichard, C. R. 3.)

sulphide with MS .

molybdate, M .

molybdophosphate, $\text{Mo}_6\text{S}_8\text{Cl}_2$.

Insol. in H_2O and alkalies. Slowly sol. in aqua regia. (Smith and Oberholtzer, Z. 1. 67.)

monamide, NH_4MoO_4 or $(\text{NH}_4)_2\text{MoO}_7$.

Prepared by H_2 + H_2O . (N. 1898, 78. 318.)

Molybdenyl bromide, MoO_2Br_2 .

Deliquescent, and sol. in H_2O with slight evolution of heat.

$\text{Mo}_2\text{O}_5\text{Br}_4$. Unstable in air. (Smith and Oberholtzer, Z. anorg. 4. 236.)

Molybdenyl potassium bromide, $\text{MoOBr}_3, 2\text{KBr}$.

(Weinland, Z. anorg. 1905, 44. 109.)

$\text{MoOBr}_3, \text{KBr} + 2\text{H}_2\text{O}$. (Weinland, Z. anorg. 1905, 44. 110.)

Molybdenyl rubidium bromide, $\text{MoOBr}_3, 2\text{RbBr}$.

(Weinland, Z. anorg. 1905, 44. 108.)

Molybdenyl chloride, MoO_2Cl_2 .

Sol. in H_2O and alcohol.

Abundantly sol. in abs. alcohol. Not very sol. in abs. ether. (Hampe, Ch. Z. 1888, 12. 23.)

+ H_2O . Composition settled by mol. wt. determinations. Dissociates in alcohol and in H_2O . (Vaudenberghe, Z. anorg. 1895, 10. 52.)

Very hygroscopic. Sol. in acetone, ether and alcohol. (Vaudenberghe, l. c.)

MoOCl_4 . Deliquescent. Sol. in little H_2O with violent action. More H_2O decomposes. (Püttbach, A. 201. 123.)

Formula is $\text{Mo}_2\text{O}_5\text{Cl}_2$, according to Blomstrand (J. pr. 71. 460).

$\text{Mo}_2\text{O}_5\text{Cl}_4$. (Püttbach, l. c.)

$\text{Mo}_2\text{O}_5\text{Cl}_6$. Deliquescent. Sol. in H_2O with very slight evolution of heat and subsequent formation of precipitate. (Blomstrand.)

Sol. in acids. (Püttbach, A. 201. 129.)

$\text{Mo}_2\text{O}_5\text{Cl}_8$. Deliquescent, and sol. in H_2O . (Blomstrand.)

$\text{Mo}_2\text{O}_5\text{Cl}_{10}$. Insol. in HCl and cold H_2SO_4 . Sol. in hot H_2SO_4 and HNO_3 . (Püttbach, A. 201. 123.)

$\text{Mo}_2\text{O}_5\text{Cl}_{12}$. Difficultly sol. in HCl . Easily sol. in HNO_3 , and alkalies + Aq. (Püttbach.)

Molybdenyl potassium chloride, $\text{MoO}_2\text{Cl}_2, \text{KCl} + \text{H}_2\text{O}$.

(Weinland, Z. anorg. 1905, 44. 97.)

+ $2\text{H}_2\text{O}$. (Weinland, Z. anorg. 1905, 44. 96.)

$6\text{MoO}_2\text{Cl}_2, 2\text{KCl} + 6\text{H}_2\text{O}$. (Weinland, Z. anorg. 1905, 44. 97.)

$\text{MoOCl}_3, 2\text{KCl} + 2\text{H}_2\text{O}$. Sol. in H_2O . (Nordenskjöld, B. 1901, 34. 1573.)

Ppt. (Henderson, Proc. Chem. Soc. 1903, 19. 245.)

Molybdenyl rubidium chloride, $\text{MoO}_2\text{Cl}_2, \text{RbCl} + \text{H}_2\text{O}$.

$\text{MoO}_2\text{Cl}_2, 2\text{RbCl}$. (Weinland, Z. anorg. 1905, 44. 95.)

$\text{MoOCl}_3, 2\text{RbCl}$. Sl. sol. in H_2O . Less sol. than K salt. (Nordenskjöld, B. 1901, 34. 1573.)

Molybdenyl fluoride, MoO_2F_2 .

Decomp. rapidly in moist air. (Schulze, J. pr. (2) 21. 442.)

Very hygroscopic. Sol. in a little H_2O giving a blue solution; in more H_2O giving a colorless solution.

Sol. in AsCl_3 , SiCl_4 , SO_2Cl_2 and PCl_3 . On warming these solutions, gas is evolved.

Insol. in toluene. Nearly insol. in ether, CHCl_3 , CCl_4 , and CS_2 . Sol. in warm pyridine and in ethyl and methyl alcohol. (Ruff, B. 1907, 40. 2934.)

MoOF_4 . Very hygroscopic. Decomp. by H_2O and conc. H_2SO_4 . Decomp. by alcohol. Sol. in ether and CHCl_3 with evolution of gas. Insol. in toluene. Very sl. sol. in benzene and CS_2 . (Ruff, B. 1907, 40. 2932.)

$\text{Mo}_2\text{O}_7\text{F}_4$. Deliquescent. Easily sol. in $\text{HF} + \text{Aq}$, not in H_2O . (Smith and Oberholtzer.)

Molybdenyl fluoride with MF.

See Fluoxymolybdate, M, and Fluoxyhypomolybdate, M.

Molybdenyl hydroxide, $\text{MoO}(\text{OH})_2$.

2 g. are sol. in 1000 cc. H_2O ; insol. in $\text{H}_2\text{O} + \text{NH}_4\text{Cl}$; only sl. sol. in NH_4OH and alkali carbonates + Aq. (Klason, B. 1901, 34. 151.)

Molybdic acid, H_2MoO_4 .

(Ullik, A. 144. 217.)

Nearly insol. in H_2O . (Vivier, C. R. 106. 601.)

Very sparingly sol. in cold H_2O , more sol. in hot H_2O . (Rosenheim and Berthelm, Z. anorg. 1903, 34. 435.)

 α -modification.

Solubility of MoO_3 , H_2O (α -modification) in H_2O at t° .

1000 g. H_2O dissolve g. MoO_3 .

t°	G. MoO_3	t°	G. MoO_3
14.8	2.117	42.0	3.446
15.2	2.131	45.0	3.661
24.6	2.619	52.0	4.184
25.6	2.689	60.0	4.685
30.3	2.973	70.0	4.231
36.0	3.085	80.0	5.212
36.8	3.295

(Rosenheim and Davidsohn, Z. anorg. 1903, 37. 318.)

(β modification), MoO_3 , H_2O . From MoO_3 , $2\text{H}_2\text{O}$ at 60° – 70° . (Rosenheim and Davidsohn.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Easily sol. in H_2SO_4 . (Ruegenberg and Smith, J. Am. Chem. Soc. 1900, 22. 772.)

H_4MoO_4 . Sol. in H_2O and acids. (Millingk.)

Very sol. in H_2O . (Mylius, B. 19 638.)

Solubility of MoO_3 , $2\text{H}_2\text{O}$ in H_2O
1000 g. H_2O dissolve g. MoO_3 at

t°	G. MoO_3	t°	G.
18	1.066	59	11
23	1.856	60	12
30	2.638	66	17
40	4.761	70	20
48	6.360	74.4	20
50.2	6.873	75	21
54	7.855	79	21

(Rosenheim and Berthelm, Z. anorg. 430.)

Solubility of MoO_3 , $2\text{H}_2\text{O}$ in ammon
+ Aq at t° .

1000 g. of the solvent dissolve g.

Solvent	t°	G
10% $(\text{NH}_4)_2\text{SO}_4$	29.6	
10% NH_4HSO_4	31.5	
"	41.8	
"	49.7	

(Rosenheim and Davidsohn, Z. anorg. 37. 315.)

H_4MoO_4 (?). Known only in sol. $\text{H}_2\text{Mo}_2\text{O}_7$. Easily sol. in H_2O . $\text{H}_2\text{Mo}_4\text{O}_{13}$. Easily sol. in H_2O . $\text{H}_2\text{Mo}_5\text{O}_{18}$. Easily sol. in H_2O . Molybdic acid also exists in a modification, sol. in H_2O . (Graham 59. 174.)

Molybdates.

The normal molybdates of the all are easily sol. in H_2O , while the sl. sol. or insol. therein.

The trimolybdates are sl. sol. in very easily sol. in hot H_2O .

The tetramolybdates are easily sol.

Aluminum molybdate, $\text{Al}_2(\text{MoO}_4)_3$.

Precipitate. (Gentile, J. pr. 81. Contains aluminum hydroxide phate. (Struve, J. pr. 61. 441.)

Aluminum ammonium molybdate.

See Aluminicomolybdate, ammonium

Aluminum barium molybdate.

See Aluminicomolybdate, barium

Ammonium molybdate, $(\text{NH}_4)_2\text{MoO}_4$.

Efflorescent through loss of NH_3 by H_2O into acid salt. (Svanberg 59. 174.)

Insol. in liquid NH_3 . (Franklin J. 1898, 20. 828.)

O₇. Sol. in H₂O.
I₄HMoO₄. Sol. in H₂O. Sol.
 I₂O. (Brandes; Mauro, Gazz.
 .)
 O₂₄+4H₂O. (Commercial am-
 bdate.)
 cent. Sol. in H₂O. (Delafor-
 1. Sc. ph. nat. 23. 17.)
 to Struve and Berlin=
 O₁₇+3H₂O.
 to Marignac and Delffs=
 . The true composition of
 ammonium molybdate is
 O₄₁. (Junius, Z. anorg. 1905,
 .
 More sol. than the above.
 3, Pogg. 127. 298.)
 etone. (Krug and M'Elroy, J.
 h. 6. 184.)
 O₄₁. True formula for com-
 onium molybdate (Sand and
 anorg. 1907, 52. 68.)
 Junius, Z. anorg. 1905, 46. 428.)
 O₁₇+H₂O. (Jean, C. R. 78.
 .
 O₁₉+H₂O. Very difficultly sol.
 sol. in hot H₂O. (Berlin, J. pr.
 .
 in NH₄OH+Aq. (Kämmerer,
 58.)
 4MoO₃. Practically insol. in
 hot H₂O. (Westphal, Dissert.
 .
 Very difficultly sol. in cold,
 sol. in hot H₂O. (Berlin.)
) dissolve 3.5200 g. at 15°; sp.
 711 g. at 18°; sp. gr.=1.04;
 12°; sp. gr.=1.05. (Wempe, Z.
 78. 258.)
 (Junius, Z. anorg. 1905, 46.
 .
 3MoO₃+13H₂O. (Rosenheim,
 7, 15. 188.)
 9MoO₃+17H₂O. (Westphal,
)
molybdenum trioxide ammonia.
 .
barium molybdate,
), 3BaO, 14MoO₃+12H₂O.
 Dissert. 1895.)
 .
bismuth molybdate,
 10O₄)₂.
 J. Am. Chem. Soc. 1903, 25.
 .
cadmium molybdate ammonia,
 1(MoO₄)₂, 2NH₃.
 y H₂O.
 NH₄OH+Aq. (Briggs, Chem.
 . 674.)
 .
cerium molybdate,
 2Mo₁₄O₄₈+24H₂O.
 . (Barbieri, C. A. 1909. 293.)

Ammonium chromic molybdate.

See **Chromicomolybdate, ammonium.**

Ammonium cobaltous molybdate,

3(NH₄)₂O, 7MoO₃, 3CoO, 7MoO₃+xH₂O.
 5[3(NH₄)₂O, 7MoO₃], 7[3CoO, 7MoO₃]+
 xH₂O.
 2[3(NH₄)₂O, 7MoO₃], 3[3CoO, 7MoO₃]
 +xH₂O.
 3[3(NH₄)₂O, 7MoO₃], 5[3CoO, 7MoO₃]
 +xH₂O.
 3(NH₄)₂O, 7MoO₃, 5[3CoO, 7MoO₃]+
 xH₂O.
 9[2(NH₄)₂O, 5MoO₃], 5[2CoO, 5MoO₃]+
 118 H₂O.
 4(NH₄)₂O, 2CoO, 15MoO₃+20H₂O.
 (Marckwald, Dissert. 1895.)

Ammonium cobaltous molybdate ammonia,
 (NH₄)₂Co(MoO₄)₂, 2NH₃.

Decomp. by H₂O.
 Sol. in dil. NH₄OH+Aq. (Briggs, Chem.
 Soc. 1904, 85. 674.)

Ammonium cobaltic molybdate.

See **Cobaltimolybdate, ammonium.**

Ammonium cupric molybdate, (NH₄)₂O, CuO,
 5MoO₃+9H₂O.

Sl. sol. in cold, sol. in boiling H₂O without
 decomp. (Struve.)

Ammonium cupric molybdate ammonia,
 (NH₄)₂Cu(MoO₄)₂, 2NH₃.

Sol. in dil. NH₄OH+Aq.
 Decomp. by H₂O. (Briggs, Chem. Soc.
 1904, 85. 673.)

Ammonium ferric molybdate, 3(NH₄)₂Mo₂O₇,
 Fe₂(MoO₄)₃+20H₂O.

Sol. in H₂O. (Struve.)
 See also **Ferricomolybdate, ammonium.**

Ammonium lanthanum molybdate,
 (NH₄)₆La₂Mo₁₄O₄₈+24H₂O.

Sol. in H₂O. (Barbieri, C. A. 1909. 293.)

Ammonium lithium molybdate, NH₄LiMoO₄
 +H₂O.

(Traube, N. Jahrb. Miner. 1894, I. 194.)

Ammonium magnesium molybdate, (NH₄)₂O,
 MgO, 2MoO₃+2H₂O=(NH₄)₂MoO₄,
 MgMoO₄+2H₂O.

Easily sol. in H₂O. (Ullik, A. 144. 344.)

Ammonium manganous molybdate,
 2(NH₄)₂O, MnO, 3MoO₃+5H₂O.

Decomp. by boiling H₂O. (Marckwald,
 Dissert. 1895.)
 (NH₄)₂O, 2MnO, 6MoO₃+16H₂O. De-
 comp. by boiling H₂O. (Marckwald, Dissert.
 1895.)

$(\text{NH}_4)_2\text{O}$, 3MnO , $6\text{MoO}_3 + 16\text{H}_2\text{O}$. Decomp. by boiling H_2O . (Marckwald, Dissert. 1895.)

$3(\text{NH}_4)_2\text{O}$, 2MnO , $12\text{MoO}_3 + 22\text{H}_2\text{O}$. (Marckwald, Dissert. 1895.)

Ammonium manganic molybdate.

See Permanganomolybdate ammonium.

Ammonium mercuric molybdate.

Sol. in $\text{HCl} + \text{Aq}$. Sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$, separating out on cooling. Sol. in hot $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Hirzel.)

Ammonium molybdenum molybdate,
 $(\text{NH}_4)_2\text{O}$, 2MoO_3 , $4\text{MoO}_3 + 9\text{H}_2\text{O}$.

Easily sol. in H_2O , but the solution soon becomes cloudy. (Rammelsberg, Pogg. 127. 291.)

Ammonium neodymium molybdate,
 $(\text{NH}_4)_3\text{NdMoO}_{14} + 12\text{H}_2\text{O}$.

Ppt. (Barbieri, C. C. 1911, I. 1043.)

Ammonium nickel molybdate,
 $(\text{NH}_4)_2\text{O}$, 3NiO , $9\text{MoO}_3 + 25\text{H}_2\text{O}$.

Very sl. sol. in cold, sol. in hot H_2O without decomp. (Marckwald, Dissert. 1895.)

$3(\text{NH}_4)_2\text{O}$, 2NiO , $10\text{MoO}_3 + 14\text{H}_2\text{O}$. Very sl. sol. in cold, sol. in hot H_2O without decomp. (Marckwald, Dissert. 1895.)

$5(\text{NH}_4)_2\text{O}$, 3NiO , $16\text{MoO}_3 + 16\text{H}_2\text{O}$. (Hall, J. Am. Chem. Soc. 1907, 29, 702.)

$6(\text{NH}_4)_2\text{O}$, 3NiO , $16\text{MoO}_3 + 29\text{H}_2\text{O}$. Very sl. sol. in cold, sol. in hot H_2O without decomp. (Marckwald, Dissert. 1895.)

$8(\text{NH}_4)_2\text{O}$, 6NiO , $31\text{MoO}_3 + 63\text{H}_2\text{O}$. Very sl. sol. in cold, sol. in hot H_2O without decomp. (Marckwald, Dissert. 1895.)

$3(\text{NH}_4)_2\text{O}$, 9NiO , $34\text{MoO}_3 + 120\text{H}_2\text{O}$. Very sl. sol. in cold, easily sol. in hot H_2O without decomp. (Marckwald, Dissert. 1895.)

Ammonium nickelic molybdate.

See Nickelomolybdate, ammonium.

Ammonium nickel hydrogen molybdate,
 $(\text{NH}_4)_4\text{H}_2[\text{Ni}(\text{MoO}_4)_6] + 5\text{H}_2\text{O}$.

See Nickelomolybdate, ammonium hydrogen.

Ammonium praseodymium molybdate,
 $(\text{NH}_4)_3\text{PrMoO}_{14} + 12\text{H}_2\text{O}$.

Ppt. (Barbieri, C. A. 1911. 1884.)

Ammonium samarium molybdate,
 $(\text{NH}_4)_3\text{SmMoO}_{14} + 12\text{H}_2\text{O}$.

Ppt. (Barbieri, C. A. 1911. 1884.)

Ammonium sodium molybdate, $7(\text{NH}_4)_2\text{O}$, $2\text{Na}_2\text{O}$, $21\text{MoO}_3 + 15\text{H}_2\text{O}$ (?).

Easily sol. in H_2O . (Delafontaine, J. pr. 95. 136.)

$7(\text{NH}_4)_2\text{O}$, $3\text{Na}_2\text{O}$, $25\text{MoO}_3 + 30\text{H}_2\text{O}$ (?). (Delafontaine.)

$(\text{NH}_4, \text{Na})_2\text{O}$, $3\text{MoO}_3 + \text{H}_2\text{O}$. Sol. (Mauro, Gazz. ch. it. 11. 214.)

Ammonium thorium molybdate.

See Thoromolybdate, ammonium.

Ammonium titanium molybdate.

See Titanomolybdate, ammonium.

Ammonium vanadium molybdate.

See Vanadiomolybdate, ammonium.

Ammonium zinc molybdate.

Sol. in H_2O . (Berzelius.)

Ammonium zirconium molybdate.

See Zirconomolybdate, ammonium.

Ammonium molybdate hydrogen
 18MoO_3 , $7(\text{NH}_4)_2\text{O}$, $3\text{H}_2\text{O} + 1$

Sol. in H_2O . (Bärwald, B. 17. 120)

Barium molybdate, basic, 2BaO , H_2O (?).

Insol. in H_2O . Sol. in dil. HCl $\text{HNO}_3 + \text{Aq}$. (Heine, J. pr. 9. 204.)

Barium molybdate, BaMoO_4 .

Difficultly sol. in H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Svanberg and Struve) Sol. in 17,200 pts. H_2O at 23° . More sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$ than in H_2O . (See Bradbury, B. 24. 2930.)

$+ 3\text{H}_2\text{O}$. (Westphal, Dissert. 1895) $\text{BaMo}_3\text{O}_{11} + 3\text{H}_2\text{O}$. Sl. sol. in H_2O . $\text{Ba}_2\text{MoO}_{11} + 9\text{H}_2\text{O}$. Appreciably sol. in H_2O . (Jørgensen.)

According to Svanberg and $\text{Ba}_2\text{Mo}_3\text{O}_{11} + 6\text{H}_2\text{O}$.

$+ 12\text{H}_2\text{O}$ or 5BaO , 12MoO_3 , (Junius, Z. anorg. 1905, 46. 433.)

$+ 22\text{H}_2\text{O}$. Ppt. (Westphal, I. 1895.)

BaO , $4\text{MoO}_3 + 3\frac{1}{2}\text{H}_2\text{O}$. Ppt. (W. anorg. 1912, 78. 320.)

$+ 12\text{H}_2\text{O}$. Ppt. (Rosenheim, I. 1913, 79. 299.)

$\text{BaMo}_3\text{O}_{11} + 4\text{H}_2\text{O}$. Insol. in cold H_2O or $\text{HNO}_3 + \text{Aq}$. Extremely easily comp. by H_2SO_4 , or $\text{H}_2\text{SO}_4 + \text{HNO}_3$, Aq . (Svanberg and Struve.)

Barium paramolybdate, 5BaO , 1 $10\text{H}_2\text{O}$.

Ppt. Sol. in excess of $\text{BaCl}_2 + \text{Aq}$. Z. anorg. 1905, 46. 433.)

Barium tetramolybdate, $\text{BaH}_2(\text{M}$ $17\text{H}_2\text{O}$.

Insol. in cold, apparently decomp. in H_2O , a small part dissolving, and forming an insol. residue. (Ullrich, 336.)

Insol. in cold and hot H_2O .
Z. anorg. 1912, 78. 320.)
 $MoO_3 + 17H_2O$. (Felix, Dissert.)

romic molybdate.

micomolybdate, barium.

baltic molybdate.

altimolybdate, barium.

nganic molybdate.

nanganomolybdate, barium.

kelic molybdate.

relimolybdate, barium.

ckel hydrogen molybdate,
 $[Ni(MoO_4)_2] + 10H_2O$.

relomolybdate, barium hydrogen.

nadium molybdate.

adiomolybdate, barium.

olybdate hydrogen dioxide, $8BaO$,
 O_2 , $2H_2O_2 + 13H_2O$.
ate. (Bärwald.)

nolybdate, Bi_2O_3 , $3MoO_3$.

at sol. in H_2O . Sol. in 500 pts.
n the stronger acids. (Richter.)

ybdenum molybdate.

er Bromomolybdenum comps.

molybdate, $CdMoO_4$.

H_2O ; sol. in $NH_4OH + Aq$, $KCN +$
ds. (Smith and Bradbury, B. 24.

MoO_3 , $8MoO_3 + 6H_2O$. Decomp. by
th H_2O . (Wempe, Z. anorg. 1912,

nolybdate, Cs_2O , $3MoO_3 + H_2O$.
m and Herschinkel, Z. anorg.
770.)

$5MoO_3 + 3H_2O$. (Ephraim and
tel, Z. anorg. 1909, 64. 270.)

O . Very sl. sol. in cold, easily sol.
). (Wempe, Dissert. 1911.)

$5MoO_3 + 5H_2O$. (Ephraim and
tel, Z. anorg. 1909, 64. 271.)

$10MoO_3 + 3H_2O$. (Ephraim and
tel, Z. anorg. 1909, 64. 271.)

$16MoO_3 + 8H_2O$. (Ephraim and
tel, Z. anorg. 1909, 64. 271.)

$10MoO_3 + 3H_2O$. (Ephraim and
tel, Z. anorg. 1909, 64. 271.)

stramolybdate, Cs_2O , $4MoO_3$.

. sol. in H_2O . (Muthmann, B.
1841.)

. Sl. sol. in H_2O . (Muthmann,
1. 1841.)

$+3H_2O$. Easily sol. in cold or hot H_2O .
(Wempe, Z. anorg. 1912, 78. 317.)

$+5H_2O$. Very sol. in cold and hot H_2O .
(Wempe, Dissert. 1911.)

Cs_2O , MoO_3 , Cs_2O , $3MoO_3 + 4.5H_2O$. Sol.
in H_2O . (Wempe, Z. anorg. 1912, 78. 317.)

Cæsium paramolybdate, $5Cs_2O$, $12MoO_3 +$
 $11H_2O$.

Efflorescent. Easily sol. in H_2O . (Wempe,
Z. anorg. 1912, 78. 317.)

Calcium molybdate, $CaMoO_4$.

Insol. precipitate. (Ullik.)

Sl. sol. in H_2O ; insol. in alcohol. (Smith
and Bradbury, B. 24. 2930.)

$+H_2O$. (Westphal, Dissert. 1895.)

$+2H_2O$. (Westphal, Dissert. 1895.)

$+6H_2O$. Difficultly sol. in cold, easily in
hot H_2O . (Ullik, A. 144. 231.)

$CaMo_4O_{13} + 9H_2O$. Easily sol. in cold H_2O .

CaO , $2H_2O$, $12MoO_3 + 21H_2O$. Efflores-
cent. Sl. sol. in cold, easily sol. in hot H_2O .
(Wempe.)

Calcium hydrogen tetramolybdate,

$CaH_2(Mo_4O_{13})_2 + 17H_2O$.

Sl. sol. in cold, easily sol. in hot H_2O with
decomp. (Ullik.)

$+16H_2O$. Insol. in cold, difficultly sol. in
hot H_2O . (Wempe, Z. anorg. 1912, 78. 318.)

Cerium molybdate, $Ce_2(MoO_4)_3$.

Precipitate. Insol. in H_2O ; sol. in acids.
(Cossa, B. 19. 536 R.)

Chromic molybdate.

Insol. in H_2O , but sol. in acids. Sol. in NH_4
molybdate + Aq. (Berzelius.)

See also Chromicomolybdic acid.

Chromic molybdate, with M. molybdate.

See Chromicomolybdate. M.

Cobaltous molybdate, $CoMoO_4$.

Decomp. by alkalis and strong acids.
(Berzelius.)

$+H_2O$. Sl. sol. in pure, easily sol. in acidi-
fied H_2O . (Coloriano, Bull. Soc. (2) 50.
451.)

CoO , $2MoO_3 + 2H_2O$. (Marckwald, Dis-
sert. 1895.)

$6\frac{1}{2}H_2O$. Sl. sol. in H_2O . (Marckwald.)

$CoMo_2O_{10} + 10H_2O$. Very sl. sol. in cold,
but very easily sol. in hot H_2O . (Ullik, W.
A. B. 55, 2. 767.)

Cobaltic potassium molybdate.

See Cobaltimolybdate, potassium.

Cobaltous sodium molybdate,

Na_2O , $2CoO$, $6MoO_3 + 18H_2O$.

(Marckwald, Dissert. 1895.)

$2Na_2O$, CoO , $7MoO_3 + 20H_2O$. Sol. in
cold H_2O without decomp. Decomp. on
heating. (Marckwald.)

$3\text{Na}_2\text{O}, 2\text{CoO}, 12\text{MoO}_3 + 27\text{H}_2\text{O}$. (Marckwald.)

$3\text{Na}_2\text{O}, 3\text{CoO}, 14\text{MoO}_3 + 50\text{H}_2\text{O}$. Sol. in much cold H_2O . (Marckwald.)

$4\text{Na}_2\text{O}, 6\text{CoO}, 25\text{MoO}_3 + 68\text{H}_2\text{O}$. (Marckwald.)

Cobaltous molybdate ammonia, $\text{CoMoO}_4, 2\text{NH}_3 + \text{H}_2\text{O}$.

Sol. in H_2O . (Sonnenschein, J. pr. 53. 340.)

Cupric molybdate, basic, $4\text{CuO}, 3\text{MoO}_3 + 5\text{H}_2\text{O}$.

Insol. in H_2O . (Struve, J. B. 1854. 350.)

Cupric molybdate, CuMoO_4 .

Sl. sol. in H_2O ; decomp. by acids and alkaline solutions.

$\text{CuMoO}_4 + 6\frac{1}{2}\text{H}_2\text{O}$. Easily sol. in cold H_2O . (Ullik, A. 144. 233.)

$+9\text{H}_2\text{O}$. Very sl. sol. in cold, and extraordinarily easily sol. in hot H_2O . (Ullik.)

Cupric molybdate ammonia, $\text{CuMoO}_4, 2\text{NH}_3 + \text{H}_2\text{O}$.

Gives off NH_3 at ord. temp. Decomp. by H_2O .

Sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$ from which it can be cryst. (Briggs, Chem. Soc. 1904, 85. 674.)

$\text{CuMoO}_4, 4\text{NH}_3$. Decomp. by H_2O . Sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Jørgensen, Ch. Z. Repert. 1896, 20. 225.)

Didymium molybdate, $\text{Di}_2(\text{MoO}_4)_3$.

Ppt. Insol. in H_2O . (Cossa, B. 19. 536R.)

$\text{Di}_2\text{O}_3, 6\text{MoO}_3 + 3\text{H}_2\text{O}$ (?). Precipitate. (Smith.)

Glucinum molybdate, basic, $2\text{GfO}, \text{MoO}_3 + 3\text{H}_2\text{O}$.

Nearly insol. in H_2O . (Atterberg, J. B. 1873. 258.)

Glucinum molybdate, $\text{GfO}, \text{MoO}_3 + 2\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Rosenheim, Z. anorg. 1897, 15. 307.)

$\text{GfMoO}_4, \text{MoO}_3 + x\text{H}_2\text{O}$. Easily sol. in H_2O . (Atterberg.)

Gold (auric) molybdate (?).

Sl. sol. in H_2O . Sol. in HCl , and $\text{HNO}_3 + \text{Aq}$. (Richter.)

Hydroxylamine potassium molybdate.

$\text{MoO}_4\text{H}_2(\text{NH}_2\text{O})_3(\text{NH}_2\text{OK})$.

Easily sol. in H_2O ; pptd. by alcohol. (Hofmann, A. 1899, 309. 324.)

Indium molybdate, $\text{In}_2(\text{MoO}_4)_3 + 2\text{H}_2\text{O}$.

Ppt. Insol. in H_2O .

Easily sol. in HCl . (Renz, B. 1901, 34. 2765.)

Iron (ferrous) molybdate, FeMoO_4 .

Insol. in H_2O . (Schultze, A. 126.)

Iron (ferric) molybdate, $\text{Fe}_2\text{O}_3, 4\text{H}_2\text{O}$.

Nearly insol. in H_2O . Slowly sol. easily in hot HCl , or $\text{HNO}_3 + \text{Aq}$. It gradually dissolve out Fe_2O_3 in t. When ignited, difficultly sol. in all a (Steinacker.)

$\text{Fe}_2\text{O}_3, 5\text{MoO}_3 + 16\text{H}_2\text{O}$. Very sl. H_2O . (Struve, J. B. 1854. 346.)

$2\text{Fe}_2\text{O}_3, 7\text{MoO}_3 + 34\text{H}_2\text{O}$. Ppt. Am. Chem. Soc. 1907, 29. 704.)

Ferric potassium molybdate, Fe_2O_3 .

$12\text{MoO}_3 + 20\text{H}_2\text{O} = 3\text{K}_2\text{Mo}_2\text{O}_7, \text{Fe}_2(\text{Mo}_2\text{O}_7)_3 + 20\text{H}_2\text{O}$.

Sol. in H_2O . (Struve.)

Lanthanum molybdate, $\text{LaH}_3(\text{MoO}_4)_3$, $\text{La}_2\text{O}_3, \text{MoO}_3 + 3\text{H}_2\text{O}$. (?)

Precipitate. (Smith.)

Lead molybdate, PbMoO_4 .

Insol. in H_2O . Sol. in warm H_2SO_4 ; decomp. by H_2SO_4 ; sol. in conc. $\text{HCl} + \text{KOH} + \text{Aq}$.

Min. *Wulfenite*. As above.

Lithium molybdate, Li_2MoO_4 .

Moderately sol. in cold, and only sol. in hot H_2O . (Ephraim, Z. anorg. 1912, 78. 309.)

$+2\frac{1}{2}\text{H}_2\text{O}$. Easily sol. in H_2O .

$5\text{Li}_2\text{O}, 5\text{MoO}_3 + 2\text{H}_2\text{O}$. 46.13 g. a in 100 ccm. of the aqueous solution and sp. gr. of the solution = 1.44. (Z. anorg. 1912, 78. 309.)

$\text{Li}_2\text{O}, 2\text{MoO}_3 + 5\text{H}_2\text{O}$. Sol. in cold sol. in hot H_2O . (Ephraim, Z. anorg. 1912, 78. 309.)

$\text{Li}_2\text{O}, 3\text{MoO}_3 + \text{H}_2\text{O}$. Easily sol. H_2O . (Wempe, Dissert. 1911.)

$+4\text{H}_2\text{O}$. (Wempe.)

$+4\frac{1}{4}\text{H}_2\text{O}$. (Wempe.)

$+7\text{H}_2\text{O}$. Nearly insol. in cold hot H_2O . (Ephraim, Z. anorg. 1912, 78. 309.)

$2\text{Li}_2\text{O}, 3\text{MoO}_3$. Sl. sol. in H_2O . (Z. anorg. 1909, 64. 258.)

Lithium paramolybdate, $3\text{Li}_2\text{O}, 12\text{H}_2\text{O}$.

Sol. in H_2O . (Rosenheim, Z. anorg. 1909, 64. 258.)

$+28\text{H}_2\text{O}$. Easily sol. in cold and (Ephraim, Z. anorg. 1909, 64. 258.)

Lithium tetramolybdate, $\text{Li}_2\text{O}, 4\text{MoO}_3$.

Sol. in cold H_2O . (Ephraim, Z. anorg. 1909, 64. 258.)

$\text{Li}_2\text{O}, \text{H}_2\text{O}, 8\text{MoO}_3 + 10\text{H}_2\text{O}$. E

O. (Wempe, Z. anorg. 1912, 78.)

H_2O , $16\text{MoO}_3 + 6\frac{1}{2}\text{H}_2\text{O}$. Easily
in H_2O . (Wempe, Z. anorg. 1912,

potassium molybdate, $\text{KLiMoO}_4 +$

, N. Jahrb. Miner, 1894, I. 194.)

n molybdate, MgMoO_4 .

elonesia.

$\text{HCl} + \text{Aq.}$ (Scacchi, Zeit. Kryst.
123.)

Easily sol. in cold, but still more
 H_2O . (Delafontaine.)

2-15 pts. cold H_2O . (Brandes.)

Easily sol. in hot or cold H_2O .

$\text{O}_{11} + 10\text{H}_2\text{O}$. Difficultly sol. in
easily in hot H_2O . (Ullik.)

n paramolybdate, $\text{Mg}_3\text{Mo}_7\text{O}_{24} +$
1.

l. in cold, more easily in hot H_2O .

n tetramolybdate,
 H_2O , $8\text{MoO}_3 + 19\text{H}_2\text{O}$.

n hydrogen tetramolybdate,
 $(\text{Mo}_4\text{O}_{11})_2 + 19\text{H}_2\text{O}$.

ol. in cold H_2O . (Ullik, A. 144.)

in cold, easily sol. in hot H_2O .
Dissert. 1911.)

l. Ppt. (Wempe, Z. anorg. 1912,

n hydrogen octomolybdate,
 $(\text{Mo}_8\text{O}_{11})_2 + 29\text{H}_2\text{O}$.

icully sol. in cold, very easily sol.
l. (Ullik, W. A. B. 60, 2. 314.)

n potassium molybdate, MgMoO_4 ,
 $\text{O}_4 + 2\text{H}_2\text{O}$.

sol. in cold, easily in hot H_2O .
44. 343.)

s molybdate, $\text{MnMoO}_4 + \text{H}_2\text{O}$.

H_2O . Sl. sol. in pure, easily sol.
1 H_2O . Decomp. by alkalis or
nates + Aq. (Coloriano, Bull. Soc.
.)

l. (Marckwald, Dissert. 1895.)

l. (Marckwald.)

potassium molybdate.

manganomolybdate, potassium.

silver molybdate.

nanganomolybdate, silver.

molybdate, $\text{Hg}_2\text{Mo}_2\text{O}_7$.

. by H_2O . (Struve, J. B. 1754.

Sol. in 500-600 pts. H_2O ; decomp. by
 $\text{HNO}_3 + \text{Aq.}$ (Hatchett.)

Molybdenum molybdate.

See Molybdenum oxides, Mo_2O_7 , Mo_4O_{11} ,
etc.

Neodymium molybdate, $\text{Nd}_2(\text{MoO}_4)_3$.

Very sl. sol. in H_2O .

1 pt. is sol. in 53790 pts. H_2O at 28° .

1 " " " " 32466 " H_2O " 75° .

(Hitchcock, J. Am. Chem. Soc. 1895, 17. 532.)

Nickel molybdate, $\text{NiMoO}_4 + \frac{1}{2}\text{H}_2\text{O} + \frac{1}{4}\text{H}_2\text{O}$
and $+5\text{H}_2\text{O}$.

(Marckwald, Dissert. 1895.)

NiO , $3\text{MoO}_3 + 18\text{H}_2\text{O}$. Sl. sol. in cold; easily
sol. in hot H_2O . (Marckwald.)

5NiO , $14\text{MoO}_3 + 57\text{H}_2\text{O}$, and $+70\text{H}_2\text{O}$.
Sl. sol. in cold; easily sol. in hot H_2O . (Marck-
wald.)

Nickel potassium molybdate, 3NiO , $5\text{K}_2\text{O}$
 $16\text{MoO}_3 + 21\text{H}_2\text{O}$.

Can be cryst. from H_2O . (Hall, J. Am.
Chem. Soc. 1907, 29. 701.)

Nickelic potassium molybdate.

See Nickelimolybdate, potassium.

Nickel potassium hydrogen molybdate,
 $\text{K}_4\text{H}_6[\text{Ni}(\text{MoO}_4)_6] + 5\text{H}_2\text{O}$.

See Nickelomolybdate, potassium hydro-
gen.

Nickel sodium molybdate, 2NiO , Na_2O ,
 $6\text{MoO}_3 + 17\text{H}_2\text{O}$.

Sol. in cold H_2O without decomp. but de-
comp. on warming. (Marckwald, Dissert.
1895.)

Nickel molybdate ammonia, NiMoO_4 , 2NH_3
 $+ \text{H}_2\text{O}$.

Decomp. by H_2O . (Sonnenschein, J. pr.
53. 341.)

Potassium molybdate, K_2MoO_4 .

Deliquescent in moist air. Very sol. in
 H_2O . Insol. in alcohol. (Svanberg and
Struve, J. pr. 44. 265.)

184.6 grams are sol. in 100 grams H_2O at
 25° . (Amadori, C. A. 1912. 2878.)

Solubility of $\text{K}_2\text{MoO}_4 + \text{K}_2\text{SO}_4$ at 25° .

G. per 100 g. H_2O		G. per 100 g. H_2O	
K_2SO_4	K_2MoO_4	K_2SO_4	K_2MoO_4
0	184.6	1.50	99.49
0.46	180.7	2.13	45.89
0.72	177.	3.95	17.48
0.98	127.2	8.55	4.73
1.27	107.5	12.10	0

(Amadori, Att. acc. Linc. 1912, 21, I. 467,
667.)

+ $\frac{3}{4}$ H₂O. Easily sol. in H₂O. (Wempe, Dissert. 1911.)

K₂O, 8MoO₃+13H₂O. Easily sol. in warm H₂O. (Wempe, Dissert. 1911.)

K₂O, 10MoO₃+9H₂O. Nearly insol in hot and cold H₂O. 100 g. H₂O dissolve 0.682 g. at 100°. (Felix, Dissert. 1912.)

+15H₂O. Sol. in H₂O. (Felix.)

5K₂O, 12MoO₃+8H₂O. Sl. sol in cold H₂O. (Junius, Z. anorg. 1905, 46. 439.)

Potassium trimolybdate, K₂Mo₃O₁₀.

Difficultly sol. in cold, but much more easily in hot H₂O. When ignited is absolutely insol. in H₂O. (Svanberg and Struve.)

+2H₂O. (Junius, Z. anorg. 1905, 46. 439.)

Sl. sol. in cold, easily sol. in hot H₂O (Wempe, Dissert. 1911.)

+2 $\frac{1}{4}$ H₂O. Easily sol. in H₂O. (Wempe, Dissert. 1911.)

+3H₂O. Very sl. sol. in cold, more easily sol. in hot H₂O. (Wempe, Dissert. 1911.)

+11H₂O. Practically insol. in H₂O. (Westphal, Dissert. 1895.)

Potassium hydrogen tetramolybdate,

K₄H₄(H₄(Mo₄O₇)₄)+18H₂O.

Sl. sol in cold H₂O. Decomp. by boiling H₂O. (Rosenheim, Z. anorg. 1913, 79. 298.)

KHMo₄O₁₁+6H₂O. Decomp. by H₂O. (Ullik.)

Potassium paramolybdate, K₄Mo₇O₂₄+4H₂O.

Decomp. even by cold H₂O. (Delafontaine.)

Formula is K₄Mo₇O₂₁+6H₂O, according to Svanberg and Struve (?).

Potassium selenium molybdate.

See Selenomolybdate, potassium.

Potassium sodium molybdate, K₂MoO₄, 2Na₂MoO₄+14H₂O.

Very easily sol. in cold, still more easily in hot H₂O. (Delafontaine.)

Potassium vanadium molybdate.

See Vanadiomolybdate, potassium.

Potassium zinc molybdate.

Sol. in H₂O. (Berzelius.)

Potassium molybdate hydrogen dioxide,
6K₂O, 16MoO₃, 4H₂O₂+13H₂O.

Sol. in H₂O. (Bärwald, C. C. 1885. 424.)

Potassium molybdate sulphocyanide, KSCN,
K₂Mo₃O₁₀+4H₂O.

Decomp. by H₂O. Sol. in dil. HCl+Ag (Péchar, C. R. 1894, 118. 806.)

Praseodymium molybdate, Pr₂(MoO₄)₃

Very sl. sol. in H₂O.

1 pt. is sol. in 65820 pta. H₂O at 21°

1 " " " " 69800 " " " " (Hitchcock, J. Am. Chem. Soc. 1909, 31. 530.)

Rubidium molybdate, Rb₂O, MoO₃.

Hygroscopic. (Ephraim, Z. anorg. 1909, 64. 263.)

Rb₂O, 2MoO₃+2H₂O. Easily sol. (Ephraim, Z. anorg. 1909, 64. 263.)

Rb₂Mo₇O₂₄+4H₂O. Very sl. sol. much more easily sol. in hot H₂O

fontaine, N. Arch. Sc. phys. nat. 30 (Ephraim, Z. anorg. 1909, 64. 26)

+4 $\frac{1}{2}$ H₂O. (Wempe, Dissert. 1911)

2Rb₂O, 7MoO₃+5H₂O. Very s. cold, very easily sol. in hot H₂O.

5Rb₂O, 7MoO₃+14H₂O. (Ephr. Herschfinkel, Z. anorg. 1909, 64. 26)

3Rb₂O, 8MoO₃+6H₂O. (Ephr. Herschfinkel, Z. anorg. 1909, 64. 26)

5Rb₂O, 12MoO₃+H₂O. 100 cc. solve 1.941 g. at 24°. (Wempe, 1912, 78. 258.)

Rb₂O, 3MoO₃. Insol. in H₂O. mann, B. 1898, 31. 1839.)

+H₂O. (Muthmann, B. 1898, 31. 1839.)

+3H₂O. Sl. sol in cold, easily s. H₂O. (Wempe, Dissert. 1911.)

6 $\frac{1}{2}$ H₂O. (Ephraim and Herschf. anorg. 1909, 64. 269.)

2Rb₂O, 3MoO₃+4H₂O. Sl. sol. easily in hot H₂O. (Wempe, Dissert. 1911.)

Rb₂O, 4MoO₃. Difficultly sol. easily in hot H₂O. (Wempe, Z. anorg. 1909, 64. 263.)

+ $\frac{1}{2}$ H₂O. Practically insol. in H₂O. sol. by addition of NH₃. (Ephr. Herschfinkel, Z. anorg. 1909, 64. 26)

+2.5H₂O. Insol. in H₂O. (Ephr. anorg. 1909, 64. 263.)

+4H₂O. Sol. in cold, more easily in hot H₂O. (Wempe, Z. anorg. 1912, 78. 312.)

Rb₂O, MoO₃, Rb₂O, 3MoO₃+5H₂O. Sol. in cold or hot H₂O. (Wempe, Z. anorg. 1912, 78. 312.)

Rb₂O, H₂O, 8MoO₃+3H₂O. Sol. in cold, easily in hot H₂O. (V. anorg. 1912, 78. 312.)

Rb₂O, 11MoO₃+5.5H₂O. Ppt. (Z. anorg. 1909, 64. 263.)

Rb₂O, 13MoO₃+4H₂O. Ppt. (Z. anorg. 1909, 64. 263.)

Rb₂O, 18MoO₃. Ppt. (Ephraim, Z. anorg. 1909, 64. 263.)

Samarium molybdate, Sm₂(MoO₄)₃.

Insol. in H₂O. (Cleve.)

Samarium sodium molybdate,

Na₂Sm₂(MoO₄)₆.

Insol. in H₂O. Easily sol. in HNO₃+Aq. (Cleve.)

tous) molybdate, Ag_2O , 2MoO_3 .
 $\text{VO}_3 + \text{Aq}$. $\text{KOH} + \text{Aq}$ dissolves
 ZrO separates out. Not decomp.
 $3\text{H} + \text{Aq}$. (Wöhler and Rauten-
 119.)
 exist. (Muthmann, B. 20. 983.)

tic) molybdate, Ag_2MoO_4 .
 sol. in H_2O ; less when HNO_3 is
 character.)
 sol. in pure H_2O ; easily sol. in
 acid with HNO_3 . (Struve and

N or $\text{NaOH} + \text{Aq}$. (Smith and
 O_3 . Sl. sol. in H_2O . Sol. in
 (Junius, Dissert. 1905.)
 10O_3 . Somewhat sol. in H_2O .
 id Struve, J. B. 1847-48. 412.)
 $\text{O}_3 + 6\text{H}_2\text{O}$. Sl. sol. in H_2O with
 Wempe, Z. anorg. 1912, 78. 322.)

n molybdate.

molybdate, silver.

late ammonia, Ag_2MoO_4 , 4NH_3 .
 O with rapid decomposition.
 Bull. Soc. (2) 20. 64.)

date hydrogen dioxide, $13\text{Ag}_2\text{O}$,
 2MoO_3 .
 wald, B. 17. 1206.)

bdate, Na_2MoO_4 .

. Easily and completely sol. in

sol. in H_2O .

Efflorescent.

solubility in H_2O at t° .

	t°	Per cent of anhydrous salt	Mols. H_2O to 1 mol of anhydrous salt	Mols. of anhydrous salt to 100 mols H_2O
H_2O	0	30.63	25.92	3.86
	4	33.83	22.38	4.47
	6	35.58	20.72	4.83
	90	38.16	18.54	5.39
I_2O	10	39.28	17.70	5.65
	15	39.27	17.70	5.65
	32	39.82	17.30	5.78
	51	41.27	16.28	6.14
	100	45.57	13.67	7.32

nk, B. 1900, 33. 3699.)

ethyl acetate. (Naumann, B.
 J.)

After ignition, very difficultly
 and very slowly sol. in hot H_2O .
 id Struve.)
 easily sol. in H_2O .

$+3\frac{1}{2}\text{H}_2\text{O}$. Easily sol. in cold or hot H_2O .
 (Wempe, Dissert. 1911.)

$+4\text{H}_2\text{O}$. Easily and completely sol. in
 cold H_2O . (Ullik.)

$+6\frac{1}{2}\text{H}_2\text{O}$. Sl. sol. in cold, very easily sol
 in hot H_2O . (Wempe, Dissert. 1911.)

$+7\text{H}_2\text{O}$. Difficultly sol. in cold H_2O , but
 more easily than the corresponding K salt.
 100 pts. H_2O dissolve 3.878 pts. at 20° and
 13.7 pts. at 100° . (Ullik, A. 144. 244.)

$+9\text{H}_2\text{O}$. Easily sol. in cold, very easily
 sol. in hot H_2O . (Wempe.)

$+11\text{H}_2\text{O}$. (Junius, Z. anorg. 1905, 46.
 437.)

$3\text{Na}_2\text{O}$, 7MoO_3 . Easily sol. in cold, very
 easily sol. in hot H_2O . (Ott, Dissert. 1911.)

$+20\text{H}_2\text{O}$. (Westphal, Dissert. 1895.)

$+22\text{H}_2\text{O}$. Efflorescent. Easily sol. in
 H_2O . (Ullik, A. 144. 219.)

Na_2O , $8\text{MoO}_3 + \frac{1}{2}\text{H}_2\text{O}$. Very sol. in cold
 or hot H_2O . (Wempe, Dissert. 1911.)

$+4\text{H}_2\text{O}$. Insol. in H_2O . (Ullik, W. A. B.
 60, 2. 312.)

$+15\text{H}_2\text{O}$. (Rosenheim, Z. anorg. 1897, 15.
 188.)

Na_2O , $10\text{MoO}_3 + 6\text{H}_2\text{O}$. Very sl. sol. in
 H_2O . 100 g. H_2O dissolve 0.842 g. at 100° .
 (Felix, Dissert. 1912.)

$+7\text{H}_2\text{O}$. (Felix.) Nearly insol. in hot and
 cold H_2O . (Rosenheim, Z. anorg. 1903, 37.
 323.)

$+12\text{H}_2\text{O}$. Difficultly sol. in H_2O .

$+21\text{H}_2\text{O}$. Abundantly but slowly sol. in
 cold H_2O . $= \text{NaHNa}_2\text{O}_{10} + 10\text{H}_2\text{O}$. (Ullik.)

$5\text{Na}_2\text{O}$, $12\text{MoO}_3 + 8\text{H}_2\text{O}$. Sl. sol. in cold,
 easily sol. in hot H_2O . (Wempe, Dissert.
 1911.)

$+20\text{H}_2\text{O}$. Sl. sol. in cold, easily sol. in
 hot H_2O . (Wempe, Dissert. 1911.)

$+36\text{H}_2\text{O}$. (Junius, Z. anorg. 1905, 46. 436.)

$+44\text{H}_2\text{O}$. Sl. sol. in cold, easily sol. in hot
 H_2O . (Wempe, Dissert. 1911.)

Sodium tetramolybdate, $\text{Na}_2\text{Mo}_4\text{O}_{13} + 6\text{H}_2\text{O}$.

Difficultly sol. in cold, easily in hot H_2O .
 (Ullik.)

100 cc. H_2O dissolve at 21° , 28.39 g.
 of the salt. Sp. gr. of the solution = 1.47.
 (Wempe, Z. anorg. 1912, 78. 306.)

$+17\text{H}_2\text{O}$. (Felix, Dissert. 1912.)

$\text{Na}_2\text{H}_4[\text{H}_2(\text{Mo}_2\text{O}_7)_2] + 21\text{H}_2\text{O}$. Slowly sol.
 in cold, easily sol. in hot H_2O . (Rosenheim,
 Z. anorg. 1913, 79. 298.)

$\text{NaHMo}_4\text{O}_{13} + 8\text{H}_2\text{O}$. Very sol. in hot or
 cold H_2O . (Ullik, A. 144. 333.)

$\text{NaHMo}_6\text{O}_{21} + 4\text{H}_2\text{O}$. Insol. in H_2O .
 (Ullik.)

Sodium manganous molybdate, $2\text{Na}_2\text{O}$, MnO ,
 $6\text{MoO}_3 + 19\text{H}_2\text{O}$.

(Marckwald, Dissert. 1895.)

Sodium molybdate molybdenum oxide,

$\text{Na}_2\text{Mo}_6\text{O}_{18}$.

Insol. in H_2O . Sol. in HNO_3 and aqua
 regia. Insol. in HCl and in H_2SO_4 . Sol. in

alkalies. (Stavenhagen and Engels, B. 1895, 28. 2280.)

Strontium molybdate, SrMoO_4 .

Sl. sol. in H_2O . (Schultze.)

Sol. in 9600 pts. H_2O at 17° . (Smith and Bradbury, B. 24. 2930.)

SrO , $3\text{MoO}_3 + \frac{1}{2}\text{H}_2\text{O}$. Scarcely sol. in cold, easily in hot H_2O . (Wempe, Dissert. 1911.)

SrO , H_2O , $8\text{MoO}_3 + 6\text{H}_2\text{O}$. Scarcely sol. in cold, easily in hot H_2O . (Wempe, Dissert. 1911.)

2SrO , $3\text{H}_2\text{O}$, $20\text{MoO}_3 + 21\text{H}_2\text{O}$. Ppt. (Wempe, Z. anorg. 1912, 78. 321.)

Thallous molybdate, Tl_2MoO_4 .

Insol. in H_2O . Sol. in alkalies. Insol. in alcohol. (Oettinger, J. B. 1864, 254.)

Sl. sol. in hot or cold H_2O . (Ullik, J. B. 1867, 234.)

$8\text{Tl}_2\text{O}$, 11MoO_3 . Sol. in hot H_2O . (Fleming, J. B. 1863, 250.)

$3\text{Tl}_2\text{O}$, 8MoO_3 . (Fleming.)

Thallous tetramolybdate, Tl_2O , $4\text{MoO}_3 + \text{H}_2\text{O}$.

Sl. sol. in H_2O with decomp. (Wempe, Z. anorg. 1912, 78. 322.)

Thallous paramolybdate, $5\text{Tl}_2\text{O}$, 12MoO_3 .

Insol. in H_2O . Easily sol. in mineral acids and in alkali hydroxides and carbonates. (Junius, Z. anorg. 1905, 46. 432.)

Tin (stannic) molybdate.

Insol. in H_2O . Sol. in dil. or conc. $\text{HCl} + \text{Aq}$, or in $\text{KOH} + \text{Aq}$. Not decomp. by $\text{HNO}_3 + \text{Aq}$. (Berzelius.)

Uranous molybdate.

Precipitate. Sol. in $\text{HCl} + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$.

$\text{U}(\text{MoO}_4)_2$. (Lancien, C. C. 1908, I. 1763.)

Uranyl molybdate, $(\text{UO}_2)\text{MoO}_4$.

Insol. in H_2O , methyl and ethyl alcohol, ether, acetic acid, CHCl_3 , C_6H_6 and C_7H_8 . Sol. in mineral acids. (Lancien, C. C. 1907, I. 784.)

2UO_3 , 3MoO_3 (?). Insol. in H_2O . Sol. in strong acids and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Berzelius.)

3UO_3 , 7MoO_3 . Insol. in hot and cold H_2O . Insol. in NaOH , KOH , and $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in all min. acids and decomp. by an excess of H_2O . Insol. in acetic acid. (Lancien, C. C. 1908, I. 1763.)

UO_3 , 8MoO_3 . (Lancien.)

$+13\text{H}_2\text{O}$. Insol. in HNO_3 . (Lancien.)

Ytterbium molybdate, Yb_2O_3 , $7\text{MoO}_3 + 6\text{H}_2\text{O}$.

Insol. in hot H_2O . (Cleve, Z. anorg. 1902, 32. 152.)

$2\text{Yb}_2\text{O}_3$, MoO_3 . Ppt. (Cleve.)

Yttrium molybdate.

Insol. in H_2O . Sol. in $\text{HNO}_3 + \text{A lin.}$)

Zinc molybdate, ZnMoO_4 .

Difficultly sol. in H_2O ; easily (Schultze, A. 126. 49.)

$+ \text{H}_2\text{O}$. Sl. sol. in H_2O . Easily dil. acids. (Coloriano, Bull. Soc. (2)

$\text{ZnMo}_3\text{O}_{11} + 10\text{H}_2\text{O}$. Very difficult in cold, but extraordinarily easily in H_2O . (Ullik, W. A. B. 55, 2. 767.)

Zinc tetramolybdate, $\text{ZnMo}_4\text{O}_{11} + 8\text{H}_2\text{O}$.

Easily sol. in cold H_2O . (Ullik.)

ZnO , H_2O , $8\text{MoO}_3 + 14\text{H}_2\text{O}$. Ppt. Z. anorg. 1912, 78. 324.)

Zinc molybdate ammonia, $\text{ZnMoO}_4 \cdot \text{H}_2\text{O}$.

(Sonnenschein, J. pr. 53. 339.)

Permolybdic acid.

See Permolybdic acid.

Molybdic sulphuric acid, MoO_3 .

Deliquescent. (Schultz-Sellack, I)

Very deliquescent. Very sol. (Muthmann, A. 1886, 238. 126.)

MoO_3 , $3\text{SO}_3 + 2\text{H}_2\text{O}$ (?).

Molybdocyanhydric acid, $\text{H}_4\text{Mo}(\text{CN})_6 \cdot 6\text{H}_2\text{O}$.

Easily sol. in H_2O and abs. alcohol. Solutions are stable at ord. temp. (Rosenheim and Garfunkel, Z. anorg. 1910, 65.)

Cadmium molybdocyanide, $\text{Cd}_2\text{M}(\text{CN})_6 \cdot 8\text{H}_2\text{O}$.

Insol. in H_2O . (Rosenheim.)

Cadmium molybdocyanide ammonia, $\text{Cd}_2\text{Mo}(\text{CN})_6 \cdot 4\text{NH}_3 + 2\text{H}_2\text{O}$.

(Rosenheim.)

Cupric molybdocyanide ammonia, $\text{Cu}_2\text{Mo}(\text{CN})_6 \cdot 4\text{NH}_3 + 7\text{H}_2\text{O}$.

(Rosenheim.)

Potassium molybdocyanide, $\text{K}_4\text{M}(\text{CN})_6 \cdot 2\text{H}_2\text{O}$.

Very sol. in H_2O . (Rosenheim.)

Thallous molybdocyanide, $\text{Tl}_4\text{Mo}(\text{CN})_6$.

Very sl. sol. in H_2O . (Rosenheim.)

Molybdoiodic acid, HIO_3 , H_2MoO_4 .

Easily sol. in H_2O . (Blomstrand, 40. 320.)

I_2O_5 , $2\text{MoO}_3 + 2\text{H}_2\text{O}$. Very sol. Insol. in cold, sol. in hot HNO_3 . Insol. in alcohol. (Chrétien, A. ch. 1898, (7) 1)

Ammonium molybdoiodate, NH_4IO_3 , O_4 .

at more sol. than K salt. (Blom-

strand, I_2O_5 , 2MoO_3 . Very sl. sol. in H_2O . More sol. in hot H_2O . (Rosen-Liebknicht, A. 1899, 308. 50.)

1 l. H_2O dissolves 5.39 g. salt at 100° . More sol. in dil. HNO_3 . (Chrétien, A. ch. 1898, (7) 15. 409.)

O , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_4 + 6\text{H}_2\text{O}$. (Chrétien.)

Barium molybdoiodate, BaO , I_2O_5 , $2\text{MoO}_3 +$

are sol. in 1 l. H_2O at ord. temp. (Chrétien.)

Cadmium molybdoiodate, acid, 3CdO , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3 + 16\text{H}_2\text{O}$.

sl. sol. in H_2O . (Chrétien.)

Calcium molybdoiodate, CaO , I_2O_5 , $2\text{MoO}_3 +$

O dissolves 7.8 g. of the salt at 90° . (Chrétien.)

Cobalt molybdoiodate, CoO , I_2O_5 , $2\text{MoO}_3 +$

are sol. in 1 l. H_2O at 15° ; 22.27 g. (Chrétien.)

Cobalt molybdoiodate acid, CoO , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3 + 18\text{H}_2\text{O}$.

sl. sol. in H_2O . (Chrétien.)

Copper molybdoiodate, CuO , I_2O_5 , $2\text{MoO}_3 +$

O dissolves 10.63 g. of the salt at 100° . (Chrétien.)

Lithium molybdoiodate, Li_2O , I_2O_5 , $2\text{MoO}_3 +$

are sol. in 1 l. H_2O at 15° . Sol. in $\text{HNO}_3 + \text{Aq}$. (Chrétien.)

Magnesium molybdoiodate, MgO , I_2O_5 , $2\text{MoO}_3 + 6\text{H}_2\text{O}$.

O dissolves 3.85 g. of the salt at 100° . (Chrétien.)

Manganese molybdoiodate, 3MnO , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3 + 9\text{H}_2\text{O}$.

O dissolves 17.05 g. of the salt at 100° . (Chrétien.)

Nickel molybdoiodate, NiO , I_2O_5 , $2\text{MoO}_3 +$

are sol. in 1 l. H_2O at 15° ; 21.8 g. at 100° . (Chrétien.)

I_2O_5 , $3\text{MoO}_3 + 23\text{H}_2\text{O}$. Easily sol. Not decomp. by acids. (Maass, 1901.)

Nickel molybdoiodate, acid, 2NiO , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3 + 15\text{H}_2\text{O}$.

Very sol. in H_2O . (Chrétien.)

Potassium molybdoiodate,

KHO , IO_3 , MoO_3OH , or KIO_3 , $\text{MoO}_3 + 2\text{H}_2\text{O}$.

Ppt. Sl. sol. in H_2O . (Blomstrand, J. pr. (2) 40. 320.)

K_2O , I_2O_5 , 2MoO_3 . Only sl. sol. in cold H_2O ; sol. on long boiling. 4.48 grs. are sol. in 1 l. H_2O at 12° . (Compare Blomstrand: not identical.) (Rosenheim, A. 1899, 308. 50.)

$+ \text{H}_2\text{O}$. Sl. sol. in H_2O . 3.45 g. are sol. in 1 l. H_2O at 15° ; 28.38 g. at 100° . More sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Chrétien, A. ch. 1898, (7) 15. 404.)

Potassium molybdoiodate, acid.

$(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3, 4\text{K}_2\text{O} + 7\text{H}_2\text{O}$.

$(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3, 2\text{K}_2\text{O} + 13\text{H}_2\text{O}$.

$(\text{I}_2\text{O}_5, 2\text{MoO}_3)_2, \text{K}_2\text{O} + 4\text{H}_2\text{O}$.

$(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3, \text{K}_2\text{O} + 7\text{H}_2\text{O}$.

$(\text{I}_2\text{O}_5, 2\text{MoO}_3)_4, \text{K}_2\text{O} + 5\text{H}_2\text{O}$.

(Chrétien.)

Silver molybdoiodate, Ag_2O , I_2O_5 , $2\text{MoO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O .

$4\text{Ag}_2\text{O}$, $4\text{I}_2\text{O}_5$, 3MoO_3 . Sol. in H_2O containing HNO_3 . (Chrétien.)

Sodium molybdoiodate, Na_2O , I_2O_5 , $2\text{MoO}_3 + \text{H}_2\text{O}$.

Sl. sol. in H_2O . Sol. in HNO_3 with decomp. (Chrétien, C. R. 1896, 123. 178.)

1 l. H_2O dissolves 6.97 g. of the salt at 15° ; 22.75 g. at 90° .

1 l. $\text{HNO}_3 + \text{Aq}$ (1 : 10) dissolves 23.78 g. of the salt at ord. temp. (Chrétien, A. ch. 1898, (7) 15. 410.)

$+ 2\text{H}_2\text{O}$. Only sl. sol. in cold H_2O ; sol. on long boiling. 3.35 grams are sol. in 1 l. H_2O at 12° . (Rosenheim, A. 1899, 308. 50.)

Strontium molybdoiodate, SrO , I_2O_5 , $2\text{MoO}_3 + 3\text{H}_2\text{O}$.

Very sol. in H_2O . (Chrétien, A. ch. 1898, (7) 15. 415.)

Strontium molybdoiodate, acid, 3SrO , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_3 + 15\text{H}_2\text{O}$.

1 l. H_2O dissolves 2.94 g. of the salt at 15° ; 13.64 g. at 100° . (Chrétien.)

Uranyl molybdoiodate, 2UO_3 , $4\text{I}_2\text{O}_5$, $3\text{MoO}_3 + 3\text{H}_2\text{O}$. (Chrétien.)

Zinc molybdoiodate, ZnO , I_2O_5 , $2\text{MoO}_3 + 5\text{H}_2\text{O}$.

1 l. H_2O dissolves 4.08 g. of the salt at 15° ; 16.25 g. at 100° . (Chrétien.)

Zinc molybdoiodate acid, ZnO , $(\text{I}_2\text{O}_5, 2\text{MoO}_3)_2$, $+16\text{H}_2\text{O}$.

Very sol. in H_2O . (Chrétien.)

Molybdoperiodic acid.

Ammonium molybdoperiodate, $5(\text{NH}_4)_2\text{O}$, I_2O_7 , $12\text{MoO}_3 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Blomstrand, Sv. V. A. H. Bih. 1892. No. 6.)

$4(\text{NH}_4)_2\text{O}$, I_2O_7 , $8\text{MoO}_3 + 7\text{H}_2\text{O}$. Very sl. sol. in cold H_2O . (Blomstrand.)

Ammonium sodium —, $2(\text{NH}_4)_2\text{O}$, Na_2O , I_2O_7 , $2\text{MoO}_3 + 10\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (B.)

Barium sodium —, 9BaO , Na_2O , $2\text{I}_2\text{O}_7$, $24\text{MoO}_3 + 28\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (B.)

Calcium —, 5CaO , I_2O_7 , $12\text{MoO}_3 + 26\text{H}_2\text{O}$.

Extremely sol. in H_2O . (Blomstrand.)

4CaO , I_2O_7 , $12\text{MoO}_3 + 21\text{H}_2\text{O}$. Less sol. in H_2O than above salt.

Lithium —, $5\text{Li}_2\text{O}$, I_2O_7 , $12\text{MoO}_3 + 30\text{H}_2\text{O}$.

Not so efflorescent as Na salt. Sol. in H_2O . (B.)

$+18\text{H}_2\text{O}$. (B.)

Manganous sodium —, 2MnO , $3\text{Na}_2\text{O}$, I_2O_7 , $12\text{MoO}_3 + 32\text{H}_2\text{O}$.

Sol. in H_2O . (B.)

Potassium —, $5\text{K}_2\text{O}$, I_2O_7 , $12\text{MoO}_3 + 12\text{H}_2\text{O}$.

Not efflorescent. (Blomstrand.)

Sodium —, $5\text{Na}_2\text{O}$, I_2O_7 , $12\text{MoO}_3 + 34\text{H}_2\text{O}$.

Efflorescent. Very sol. in H_2O . (Blomstrand, Sv. V. A. H. Bih. 1892. No. 6. 24.)

$+26\text{H}_2\text{O}$. Not efflorescent. Very sol. in H_2O . (Blomstrand.)

Sodium strontium —, Na_2O , 4SrO , I_2O_7 , $12\text{MoO}_3 + 20\text{H}_2\text{O}$.

Sol. in H_2O . (B.)

Molybdophosphoric acid.

See Phosphomolybdic acid.

Molybdosubphosphoric acid.

Sodium molybdosubphosphate, $\text{Na}_2[\text{P}(\text{Mo}_2\text{O}_7)_3] + 8\text{H}_2\text{O}$.

Ppt. (Rosenheim, Z. anorg. 1913, 84. 222.)

Molybdophosphorous acid.

Potassium molybdophosphite, $\text{K}_2[\text{HP}(\text{Mo}_2\text{O}_7)_3] + 11\text{H}_2\text{O}$.

Difficultly sol. in cold H_2O . (Rosenheim, Z. anorg. 1913, 84. 219.)

Sodium molybdophosphite, $\text{Na}_2[\text{HP}(\text{Mo}_2\text{O}_7)_3] + 11\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Rosenheim, Z. 1913, 84. 218.)

Molybdophosphovanadic acid.

See Phosphovanadiomolybdic acid.

Molybdoselenious acid.

Ammonium molybdoselenite, $4(\text{NH}_4)_2\text{O}$, 3SeO_2 , $10\text{MoO}_3 + 4\text{H}_2\text{O}$.

More sol. in hot than cold H_2O ; alcohol. (Péchar, A. ch. (6) 30. 40.)

Ammonium potassium molybdoselenite, $2(\text{NH}_4)_2\text{O}$, $2\text{K}_2\text{O}$, 3SeO_2 , $10\text{MoO}_3 + 5\text{H}_2\text{O}$.

Very sol. in H_2O ; insol. in alcohol and.)

Barium molybdoselenite, 4BaO , 3SeO_2 , $10\text{MoO}_3 + 3\text{H}_2\text{O}$.

Sl. sol. in cold, easily in war (Péchar.)

Potassium molybdoselenite, $4\text{K}_2\text{O}$, 3SeO_2 , $10\text{MoO}_3 + 5\text{H}_2\text{O}$.

Very sol. in H_2O ; insol. in alcohol and.)

Sodium molybdoselenite, $4\text{Na}_2\text{O}$, 3SeO_2 , $10\text{MoO}_3 + 15\text{H}_2\text{O}$.

Very efflorescent, and sol. in H_2O ; alcohol. (Péchar.)

Molybdosilicic acid.

See Silicomolybdic acid.

Molybdosilicovanadic acid.

See Silicovanadiomolybdic acid.

Molybdosulphuric acid.

Appreciably sol. in H_2O . (F. Dissert. 1903.)

Ammonium molybdosulphate, $(\text{NH}_4)_2\text{MoO}_4$, $\text{SO}_3 + 4\text{H}_2\text{O}$, and $+9\text{H}_2\text{O}$.

Decomp. by H_2O . (Weinland, 1907, 54. 261.)

$(\text{NH}_4)_2\text{O}$, 2MoO_3 , $3\text{SO}_3 + 10\text{H}_2\text{O}$ land.)

Ammonium molybdenyl molybdate, $(\text{NH}_4)_2\text{O}$, MoO_3 , 7MoO_3 , SO_3 .

$1\frac{1}{2}(\text{NH}_4)_2\text{O}$, MoO_3 , 7MoO_3 , SO_3 , $2(\text{NH}_4)_2\text{O}$, MoO_3 , 7MoO_3 , SO_3 .

(Hoffmann, Dissert. 1903.)

3NH_3 , MoO_3 , 7MoO_3 , $\text{SO}_3 + 10\text{H}_2\text{O}$ sol. in H_2O . Very sl. sol. in NH_4OH .

Very stable toward alkali + Aq. C. R. 1893, 116. 1441.)

$\text{oO}_3, 7\text{MoO}_3, \text{SO}_3 + 8\text{H}_2\text{O}$. (Pé-
1893, 116. 1441.)

molybdosulphate, $\text{K}_2\text{O}, 2\text{MoO}_3,$
 H_2O .

$\text{O}_3, \text{SO}_3 + 6\text{H}_2\text{O}$.
 $\text{O}_3, 3\text{SO}_3 + 6\text{H}_2\text{O}$. (Weinland, Z.
54. 260.)

molybdenyl molybdosulphate,
 $\text{oO}_3, 7\text{MoO}_3, \text{SO}_3 + 8\text{H}_2\text{O}$.
1, Dissert. 1903.)

sulphurous acid.

molybdosulphite, $4(\text{NH}_4)_2\text{O},$
 $10\text{MoO}_3 + 6\text{H}_2\text{O}$.

cold, more easily in hot H_2O .
alcohol. (Péchar, A. ch. (6) 30.

$8\text{MoO}_3, 2\text{SO}_2 + 5\text{H}_2\text{O}$. Sl. sol.
ly sol. in warm H_2O . Easily de-
 H_2O , and can be recryst. only in
an excess of sulphurous acid.
Z. anorg. 1894, 7. 177.)

potassium molybdosulphite,
 $\text{O}, 2\text{K}_2\text{O}, 3\text{SO}_2, 10\text{MoO}_3 + 9\text{H}_2\text{O}$.
cold H_2O . Decomp. on warming.

molybdosulphite, $2\text{BaO}, 5\text{MoO}_3,$
 $10\text{H}_2\text{O}$.

m, Z. anorg. 1897, 15. 185.)

molybdosulphite, $2\text{Cs}_2\text{O}, 5\text{MoO}_3,$
 $10\text{H}_2\text{O}$.

As K salt. (Rosenheim.)

molybdosulphite, $4\text{K}_2\text{O}, 3\text{SO}_2,$
 $10\text{H}_2\text{O}$.

l. in H_2O , but decomp. on warm-
rd.)

$\text{O}_3, 2\text{SO}_2 + \text{H}_2\text{O}$. (Rosenheim.)

molybdosulphite, $2\text{Rb}_2\text{O}, 5\text{MoO}_3,$
 $10\text{H}_2\text{O}$.

(Rosenheim.)

molybdosulphite, $4\text{Na}_2\text{O}, 3\text{SO}_2,$
 $10\text{H}_2\text{O}$.

in cold H_2O ; insol. in alcohol.

Very efflorescent. (Péchar.)
 $4\text{MoO}_3, 2\text{SO}_2 + 8\text{H}_2\text{O}$. In dry state
gives off SO_2 and soon effloresces.
)

molybdosulphite, $2\text{SrO}, 5\text{MoO}_3,$
 $10\text{H}_2\text{O}$.

m.)

anic acid.

molybdic acid.

Molybdous acid.

Magnesium molybdite, $\text{Mg}_2\text{Mo}_3\text{O}_8 = 2\text{MgO},$
 3MoO_3 .

Not attacked by KOH , and $\text{HCl} + \text{Aq}$.
(Muthmann, A. 238. 108.)

Zinc molybdite, $\text{Zn}_2\text{Mo}_3\text{O}_8 = 2\text{ZnO}, 3\text{MoO}_3$.

Easily sol. in aqua regia. (Muthmann, A.
238. 108.)

Molybdovanadates.

See Vanadiomolybdates.

Neodymicotungstic acid.

Ammonium neodymicotungstate, $3(\text{NH}_4)_2\text{O},$
 $\text{Nd}_2\text{O}_3, 16\text{WO}_3 + 20\text{H}_2\text{O}$.

Difficultly sol. in H_2O . (E. F. Smith, J.
Am. Chem. Soc. 1904, 26. 1480.)

Barium neodymicotungstate, $6\text{BaO}, \text{Nd}_2\text{O}_3,$
 $16\text{WO}_3 + 17\text{H}_2\text{O}$.

Insol. in H_2O . (E. F. Smith.)

Neodymium.

See also under Didymium.

Neodymium bromide, NdBr_3 .

(Matignon, C. R. 1905, 140. 1638.)

Neodymium carbide, NdC_2 .

Decomp. by H_2O ; insol. in conc. HNO_3 ;
decomp. by dil. HNO_3 . (Moissan, C. R.
1900, 131. 597.)

Neodymium chloride, NdCl_3 .

100 g. H_2O dissolve 98.68 g. NdCl_3 at 13° ;
140.4 g. at 100° .

Sp. gr. at $15^\circ/4^\circ$ of the solution sat. at $13^\circ =$
1.74. (Matignon, A. ch. 1906, (8) 8. 249.)

44.5 g. are sol. in 100 g. abs. alcohol at 20° .
1.8 g. " " " " " pyridine at 15° .

Insol. in ether, CHCl_3 , quinoline, toluidine,
etc. Sl. sol. in aniline and in phenylhydra-
zine. (Matignon, A. ch. 1906, (8) 8. 266.)

+ $6\text{H}_2\text{O}$. Deliquescent.

At 13° , 100 pts. H_2O dissolve 246.2 pts. of
the hydrated salt.

At 100° , 100 pts. H_2O dissolve 511 pts. of
hydrated salt.

Sat. solution at 13° has a sp. gr. $15^\circ/4^\circ =$
1.741. (Matignon, C. R. 1901, 133. 289.)

Neodymium chloride ammonia, $\text{NdCl}_3,$
 12NH_3 .

Decomposes on heating into $\text{NdCl}_3 + \text{NH}_3$;
+ 2NH_3 ; + 4NH_3 ; + 5NH_3 ; + 8NH_3 ; and
+ 11NH_3 . (Matignon, C. R. 1906, 142. 1043.)

Neodymium hydroxide.

Sol. in citric acid. (Baskerville, J. Am.
Chem. Soc. 1904, 26. 49.)

Solubility in glycerine+Aq containing about 60% by vol. of glycerine. 100 ccm. of the solution contain 4.5 g. neodymium oxide. (Müller, Z. anorg. 1905, **43**. 322.)

Neodymium hydride, NdH₂ (?).

Slowly attacked by boiling H₂O. Sol. in acids with violent evolution of H₂. (Muthmann, A. 1904, **331**. 58.)

Neodymium iodide, NdI₃.

(Matignon, C. R. 1905, **140**. 1638.)

Neodymium nitride, NdN.

Decomp. in moist air with evolution of NH₃. (Muthmann, A. 1904, **331**. 59.)

Neodymium oxide, Nd₂O₃.

Easily sol. in acids. (v. Welsbach, M. 6. 477.)

Neodymium oxychloride, NdOCl.

(Matignon, C. R. 1905, **140**. 1638.)

Neon, Ne.

Less sol. than argon in H₂O; sol. in liquid oxygen. (Ramsay, B. 1898, **31**. 3118.)

Absorption by H₂O at t°.

t°	Coefficient of absorption
0	0.0114
10	0.0118
20	0.0147
30	0.0158
40	0.0203
50	0.0317

(Antropoff, Roy. Soc. Proc. 1910, **83**. A, 480.)

Nickel, Ni.

Not attacked by H₂O. Very slowly sol. in dilute H₃PO₄, H₂SO₄, or HCl+Aq. (Tupputi, A. ch. 78. 133.)

Very easily attacked by HNO₃+Aq, and difficultly by hot H₂SO₄. When pure, is converted into passive condition by conc. HNO₃. (Nicklès, C. R. **38**. 284.)

Very sl. attacked by cold acids, except HNO₃+Aq. (Tissier, C. R. **50**. 106.)

Not attacked by NaOH+Aq. (Venator, Dingl. **261**. 133.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, **20**. 828.)

Nickel amide, Ni(NH₂)₂.

Decomp. by H₂O; slowly sol. in min. acids. Insol. in liquid NH₃. (Bohart, J. phys. Chem. 1915, **19**. 560.)

Nickel antimonide, NiSb.

Insol. in HCl+Aq; easily sol. in HNO₃+Aq. (Christofle, 1863.)

Min. *Breithauptite*. Insol. in acid sol. in aqua regia.

Ni₂Sb₂. (Christofle.)

Nickel antimonide sulphide, NiSb, NiSbS.

Min. *Nickel glance*, *Ullmannite*.

Decomp. by HNO₃+Aq; compl in aqua regia with separation of S.

Nickel arsenide, NiAs.

Min. *Niccolite*. Sol. in conc. H with separation of As₂O₃; more eas aqua regia.

NiAs₂. Min. *Chloanthite*, *Ram* Sol. in HNO₃+Aq.

Ni₃As₂. Sol. in HNO₃ and in a Readily attacked by fused alkali. C. R. 1900, **130**. 915.)

Nickel arsenide sulphide, NiAs₂, N

Min. *Gersdorffite*. Partly sol. in Aq with separation of S and As₂O₃ tacked by KOH+Aq.

Nickel azoimide, basic, Ni(OH)N₂

Insol. in H₂O. (Curtius, J. pr. **58**. 300.)

Nickel azoimide, NiN₂+H₂O.

Sol. in H₂O; insol. in alcohol (Curtius, J. pr. 1900, (2) **61**. 418.)

Nickel potassium azoimide, Ni(N₂)₂

Sol. in H₂O. (Curtius, J. pr. 18 302.)

Nickel boride, Ni₂B.

Attacked by HNO₃. Slowly a HCl. (Jassoneix, C. R. 1907, **145**.

NiB. Decomp. by moist air and nitrates, chlorates, hydroxides an ates; decomp. by steam at red h attacked by HCl. Easily attacked and aqua regia; by H₂SO₄ only o (Moissan, C. R. 1896, **122**. 425.)

NiB₂. (Jassoneix, C. R. 1907, **14**

Nickel bromide, NiBr₂.

Deliquescent. Slowly sol. in H₂ Sat. NiBr₂+Aq contains at:

—21°	—6°	+19°	38°
47.1	51.7	56.6	58.9% Ni

58°	77°	98°	100°	140
60.5	60.3	61.0	61.0	60.7

(Étard, A. ch. 1894, (7) **2**. 5

Somewhat hygroscopic. Nearly cold H₂O but begins to dissolve a at 50°, and somewhat more rapid but even at that temp. 1 g. requires for solution. HNO₃ does not a hasten solution. (Richards and Z. anorg. 1898, **16**. 169.)

Nickel chloride, NiCl_2 .
 Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)
 Sol. in quinoline. (Beckmann and Gabel, anorg. 1906, 51. 236.)
 $+3\text{H}_2\text{O}$. Deliquescent. Very sol. in H_2O , $\text{HCl}+\text{Aq}$, $\text{NH}_4\text{OH}+\text{Aq}$, alcohol, and ether. (Berthelot, A. ch. 44. 389.)
 $+6\text{H}_2\text{O}$. (Bolschakoff, C. C. 1897, II. 726 and 726.)
 $+9\text{H}_2\text{O}$. (Bolschakoff, C. C. 1897, II. 726 and 331.)

Nickel stannic bromide.

See Bromostannate, nickel.

Nickel bromide ammonia, $\text{NiBr}_2 \cdot 6\text{NH}_3$.

Sol. in little H_2O , but decomp. by more. (Rammelsberg, Pogg. 55. 243.)
 Sol. in warm conc. $\text{NH}_4\text{OH}+\text{Aq}$; insol. in cold. (Richards and Cushmann, Z. anorg. 1898, 16. 175.)

Nickel bromide cupric oxide, $\text{NiBr}_2 \cdot 3\text{CuO} + 4\text{H}_2\text{O}$.

Not decomp. by H_2O . (Mailhe, A. ch. 1902, (7) 27. 377.)

Nickel bromide hydrazine, $\text{NiBr}_2 \cdot 2\text{N}_2\text{H}_4$.

Easily sol. in dil. acids and $\text{NH}_4\text{OH}+\text{Aq}$.
 $\text{NiBr}_2 \cdot 3\text{N}_2\text{H}_4$. Sol. in dil. acids. (Franzen, Z. anorg. 1908, 60. 263-4.)

Nickel carbonyl, $\text{Ni}(\text{CO})_4$.

Insol. in H_2O ; not attacked by dil. acids or alkalis or conc. $\text{HCl}+\text{Aq}$. Easily sol. in conc. HNO_3+Aq and in aqua regia. Sol. in alcohol, benzene, and chloroform. (Mond, Langer, and Quincke, Chem. Soc. 57. 749.)

Sol. in hydrocarbons, especially oil of turpentine. (Berthelot, C. R. 1891, 112. 1346.)

Sol. in acetone, toluene, methyl and ethyl alcohol, etc. (Lenher and Loos, J. Am. Chem. Soc. 1900, 22. 114.)

Nickel chloride, NiCl_2 .

Anhydrous. Not immediately sol. in H_2O , but gradually dissolves on boiling or by addition of $\text{HCl}+\text{Aq}$. Deliquesces on air, and is then easily sol. in H_2O . Sol. in $\text{NH}_4\text{OH}+\text{Aq}$. Sol. in alcohol. Sol. in hot $\text{HCl}+\text{Aq}$ only slowly.

Sp. gr. of NiCl_2+Aq containing:

5	10	15	20	25% NiCl_2
1.0493	1.0995	1.1578	1.2245	1.3000

(B. Franz, J. pr. (2) 5. 285.)

Sp. gr. of NiCl_2+Aq containing, in 1000 grms. H_2O , g. $\text{NiCl}_2+7\text{H}_2\text{O}$ at 23.1° :

128 g. ($=\frac{1}{2}$ mol.)	256	384	512
1.057	1.107	1.149	1.187
640	768	896	1024
1.220	1.249	1.276	1.301

Containing g. NiCl_2 (anhydrous):

65 g. ($=\frac{1}{2}$ mol.)	130	195	260	325	390
1.061	1.119	1.176	1.230	1.284	1.335

(Gerlach, Z. anal. 28. 468.)

Sp. gr. of NiCl_2+Aq at room temp. containing:

11.449	22.69	30.40% NiCl_2
1.1093	1.2264	1.3371

(Wagner, W. Ann. 1883, 18. 269.)

Sp. gr. of NiCl_2+Aq at 25° .

Concentration of NiCl_2+Aq	Sp. gr.
1-normal	1.0591
$\frac{1}{2}$ " "	1.0308
$\frac{1}{4}$ " "	1.0144
$\frac{1}{8}$ " "	1.0067

(Wagner, Z. phys. Ch. 1890, 5. 39.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

100 pts. absolute alcohol dissolve at room temperature 10.05 pts. NiCl_2 . (Bödtker, Z. phys. Ch. 1897, 22. 511.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Solubility in glycol = 16.1-16.3%. (de Coninck, C. C. 1905, II. 1234.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Anhydrous NiCl_2 is insol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

$+ \text{H}_2\text{O}$. (Baubigny.)

1 l. sat. $\text{HCl}+\text{Aq}$ at 12° contains 40 g. NiCl_2 dissolved from $\text{NiCl}_2, \text{H}_2\text{O}$. (Ditte.)

$+2\text{H}_2\text{O}$. (Sabatier, Bull. Soc. (3) 1. 88.)

$+6\text{H}_2\text{O}$. Deliquescent in moist, efflorescent in dry air; sol. in H_2O with evolution of heat. Sol. in 1.5 to 2 pts. H_2O . Easily sol. in alcohol. (Tupputi.)

1 l. H_2O dissolves 600 g. $\text{NiCl}_2+6\text{H}_2\text{O}$. (Ditte, A. ch. 1879, (5) 22. 551.)

Sat. aq. solution contains at:

-17°	-16°	$+10^\circ$	18°
29.7	31.0	37.3	38.5% NiCl_2

38°	59°	78°	96°
41.9	45.0	46.6	46.7% NiCl_2

(Étard, A. ch. 1894, (7) 2. 539.)

Solubility of $\text{NiCl}_2+6\text{H}_2\text{O}=37.53\%$ NiCl_2 at 25° . (Foote, J. Am. Chem. Soc. 1912, 34. 882.)

100 pts. absolute alcohol dissolve at room temperature 53.71 pts. $\text{NiCl}_2+6\text{H}_2\text{O}$. (Bödtker, Z. phys. Ch. 1897, 22. 511.)

+7H₂O. 100 g. absolute alcohol dissolve 2.16 g. NiCl₂+7H₂O at 17° and 1.4 g. at 3°. (de Bruyn, R. t. c. 1892, 11. 156.)

Nickel hydrogen chloride, 3NiCl₂, 2HCl+1½H₂O.
(Reitzenstein, Z. anorg. 1898, 18. 270.)

Nickel rubidium chloride, NiCl₂, 2RbCl.
Easily sol. in H₂O and HCl+Aq. (Godefroy, B. 8. 9.)

Nickel thallic chloride, NiCl₂, 2TlCl₃+8H₂O.
Deliquescent. Can be cryst. from H₂O. (Gewecke, A. 1909, 366. 221.)

Nickel tin (stannous) chloride, NiCl₂, SnCl₂+6H₂O.
Sol. in H₂O. (Jørgensen.)

Nickel tin (stannic) chloride.
See Chlorostannate, nickel.

Nickel chloride ammonia, NiCl₂, 2NH₃.
Sol. in H₂O, decomp. on boiling; insol. in alcohol.

NiCl₂, 3NH₃+3H₂O. (André, C. R. 1888, 106. 937.)

NiCl₂, 6NH₃. Sol. in cold H₂O without decomp. Insol. in alcohol. Very sl. sol. in conc. NH₄OH+Aq.

Nearly insol. in a sat. solution of NH₄Cl in NH₄OH+Aq. (Sørensen, Z. anorg. 1894, 5. 363.)

Nickel chloride cupric oxide, NiCl₂, 3CuO+4H₂O.

Not decomp. by H₂O. (Mailhe, A. ch. 1902, (7) 27. 377.)

Nickel chloride hydrazine, NiCl₂, 2N₂H₄.
Sol. in dil. acids and NH₃+Aq. (Franzen, Z. anorg. 1908, 60. 262.)

NiCl₂, 3N₂H₄. Sol. in dil. acids. (F.)

Nickel fluoride, NiF₂.

Sol. in about 5000 pts. H₂O; insol. in alcohol and ether. Not attacked by HCl, HNO₃, or H₂SO₄ even when hot. (Poulenc, C. R. 114. 1426.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 828.)

+2H₂O. Decomp. by pure H₂O. Sol. in H₂O acidulated with HF. (Berzelius.)

+3H₂O. (Clarke, Sill. Am. J. (3) 13. 291.)

Nickel hydrogen fluoride, NiF₂, 5HF+6H₂O.

Easily sol. in H₂O and dil. acids. Sol. in NH₄OH+Aq with decomp. (Böhm, Z. anorg. 1905, 43. 330.)

Nickel potassium fluoride, NiF₂, KF.

+H₂O. Sol. in H₂O. (Wagner, B. 19. 896.)

NiF₂, 2KF. Sl. sol. in H₂O. Scarce in methyl or ethyl alcohol or br (Poulenc, C. R. 114. 747.)

Nickel potassium zirconium fluoride.

See Fluozirconate, nickel potassium.

Nickel manganic fluoride.

See Fluomanganate, nickel.

Nickel sodium fluoride, NiF₂, NaF+1
Sol. in H₂O. (Wagner, B. 19. 896.)

Nickel stannic fluoride.

See Fluostannate, nickel.

Nickel titanium fluoride.

See Fluotitanate, nickel.

Nickel tungstyl fluoride.

See Fluoxytungstate, nickel.

Nickel vanadium fluoride.

See Fluovanadate, nickel.

Nickel zirconium fluoride.

See Fluozirconate, nickel.

Nickel fluoride ammonia, 5NiF₂, 8H₂O.

Insol. in cold H₂O. Decomp. by h Easily sol. in dil. acids. (Böhm, Z. 1905, 43. 334.)

Nickelous hydroxide, 4NiO₂H₂, H₂O.

Very sl. sol. in H₂O. Sol. in acids in KOH or NaOH+Aq. Somewh cultly sol. in (NH₄)₂CO₃ or NH₄OH+ easily sol. in presence of NH₄ salts. NH₄ salts+Aq. Sol. in KCN+Aq gers, 1834.)

Sol. in boiling NH₄Cl+Aq.

NiO₂H₂.

Solubility in NH₄OH+Aq at 2

NH ₃ norm.	G. Ni per l.	G. NiO ₂
1	0.084	0.0
2	0.170	0.0
3	0.257	0.0
4	0.360	0.0
4.911	2.580	0.0
3.900	1.780	0.0
2.101	0.835	0.0
0.602	0.158	0.0

The non-agreement of the results i the formation of different modifica NiO₂H₂.

(Bonsdorff, Z. anorg. 1904, 41. 1

y in $\text{NH}_4\text{OH} + \text{Aq.}$

$\text{Ni} = 0.014\text{N}$ in $1\text{N } \text{NH}_4\text{OH} + \text{Aq.}$

" $= 0.036\text{N}$ " $2\text{N } \text{NH}_4\text{OH} + \text{Aq.}$

B. 1903, **36**. 3840.)

not $\text{NH}_4\text{F} + \text{Aq.}$ (von Helmholtz, Z. 3, **3**. 133.)

methyl or amyl amine. (Wurtz.)

td. in presence of Na citrate.

d. in presence of a large number of the organic substances, particularly (Rose.)

Nickelic hydroxide, $\text{Ni}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$.

acids; insol. in H_2O and alkalies. Am. Chem. Soc. 1896, **18**. 901.)

Hydroxide, $\text{Ni}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$ (?).

ke, Pogg. **141**. 122.)

H_2O (?). Sol. in acids as nickelous not attacked by boiling KOH or q. Slowly sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ H_2OH , and NH_4 salts + Aq. (Ode-

ide, NiI_2 .

scant and sol. in H_2O . (Erdmann, 54.)

$2 + \text{Aq}$ contains at:

—6°	+11°	16°
54.3	57.8	59.0% NiI_2 ,

80°	85°	90°
65.0	65.2	65.7% NiI_2 .

ard, A. ch. 1894, (7) **2**. 546.)

. Deliquescent. Easily sol. in dmann.)

Nickel ammonia, $\text{NiI}_2 \cdot 4\text{NH}_3$.

elsberg, Pogg. **48**. 119.)

NH_3 . Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Very sl. sol. in conc. Aq. (Erdmann.)

Nickel hydrazine, $\text{NiI}_2(\text{N}_2\text{H}_4)_2$.

a H_2O . Sol. in acids. (Franzen, 1911, **70**. 150.)

Nickel oxide, $\text{Ni}_2\text{O}_3 + \text{H}_2\text{O}$.

a H_2O ; sol. in HCl and H_2SO_4 and so in $\text{KCN} + \text{Aq.}$ (Moore, C. N. 31.)

Nickel oxide, NiO .

a H_2O . Sol. in conc. acids, except talline, when it is scarcely attacked (Ebelmen, C. R. **33**. 256.)

sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq.}$ (De-

rwly sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Insol. in l $\text{NaOH} + \text{Aq.}$

min. acids, especially $\text{HCl} + \text{Aq.}$, med; insol. in $\text{HC}_2\text{H}_3\text{O}_2$, NH_4Cl ,

and $\text{NH}_4\text{SCN} + \text{Aq.}$ Insol. in conc. $\text{NaOH} + \text{Aq.}$ (Zimmerman, A. **232**. 324.)

1 l. solution containing 418.6 g. sugar and 34.3 g. CaO dissolves 0.29 g. NiO . (Bodenbender, J. B. 1865. 600.)

Min. *Bunsenite*.

Nickelonickelic oxide, Ni_2O_4 .

Sol. in acids. (Baubigny, C. R. **87**. 1082.) + $2\text{H}_2\text{O}$. Insol. in H_2O , and in alkalies + Aq. Sol. in acids. (Dudley, J. Am. Chem. Soc. 1896, **18**. 901.)

6NiO , $\text{Ni}_2\text{O}_3 + \text{H}_2\text{O}$. (Schönbein, J. pr. **93**. 35.)

Nickelic oxide, Ni_2O_3 .

Sol. in HNO_3 , H_2SO_4 , or $\text{HCl} + \text{Aq}$ with decomp., also in NH_4OH and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ (Winkelblech, A. **13**. 259.)

Nickel peroxide, Ni_2O_4 (?).

(Bayley, C. N. **39**. 81.)

Correct composition is Ni_2O_3 . (Carnot, C. R. **108**. 610.)

Ni_4O_7 (?). (Wicke, Zeit. Ch. 1865. 303.)

NiO_4 . (Hollard, C. R. 1903, **136**. 230.)

Nickel oxychloride.

Sl. sol. in H_2O . (Berzelius.)

NiCl_2 , $8\text{NiO} + 13\text{H}_2\text{O}$. (Raoult, C. R. **69**. 826.)

Nickel oxyiodide, NiI_2 , $9\text{NiO} + 15\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$ or acetic acid. Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Alcohol dissolves out NiI_2 . (Erdmann.)

Nickel oxyselenide.

Almost insol. in boiling HCl; decomp. by HNO_3 . (Fonzes-Diacon, C. R. 1900, **131**. 557.)

Nickel phosphide, Ni_2P .

Sol. in $\text{HNO}_3 + \text{Aq}$ and aqua regia; insol. in $\text{HCl} + \text{Aq.}$ (Struve, J. pr. **79**. 321.)

Sol. in aqua regia and in HNO_3 ; sol. in fused alkali. (Granger, Bull. Soc. 1896, (3) **15**. 1089.)

Easily sol. in HNO_3 . (Granger, C. N. 1898, **77**. 229.)

When prepared by heating phosphorus, copper and nickel in electric furnace, is insol. in all acids except a mixture of HNO_3 and HF. (Maronneau, C. R. 1900, **130**. 657.)

NiP_2 . Sol. in HNO_3 ; decomp. by fused NaOH . (Jolibois, C. R. 1910, **150**. 107.)

NiP_3 . Sol. in HNO_3 ; decomp. by fused NaOH . (J.)

Ni_2P_3 . Insol. in HNO_3 , HCl and aqua regia; stable in the air even when heated (Granger, Bull. Soc. 1896, (3) **15**. 1086.)

Ni_3P_2 . Not attacked by HCl. Easily attacked by HNO_3 . (Rose, Pogg. 1832, **24**. 232.)

Ni_3P_2 . Sol. in HNO_3 , aqua regia and in fused alkali. (Granger, C. R. 1896, 123. 177.)

Nickel phosphosulphide, Ni_3PS_2 .

Decomp. by hot H_2O or by aqua regia. Sl. attacked by HNO_3 . (Ferrand, A. ch. 1899, (7) 17. 417.)

Nickel semiselenide, Ni_2Se .

Almost insol. in boiling HCl ; decomp. by HNO_3 . (Fonzes-Diacon, C. R. 1900, 131. 557.)

Nickel selenide, NiSe .

Insol. in H_2O , dil. or conc. $\text{HCl} + \text{Aq}$; slowly sol. in $\text{HNO}_3 + \text{Aq}$; easily in aqua regia. (Little, A. 112. 211.)

Almost insol. in boiling HCl ; decomp. by HNO_3 . (Fonzes-Diacon, C. R. 1900, 131. 557.)

NiSe_2 . (Fonzes-Diacon.)

Ni_2Se_3
 Ni_3Se_4 } Almost insol. in boiling HCl ; decomp. by HNO_3 . (Fonzes-Diacon.)

Nickel silicide, Ni_3Si .

Sol. in HF and aqua regia; insol. in cold H_2O ; decomp. by steam at red heat; sol. in fused alkali carbonates. (Vigouroux, C. R. 1895, 121. 687.)

Nickel semisulphide, Ni_2S .

Sol. in $\text{HNO}_3 + \text{Aq}$, with residue of S. Difficultly sol. in conc. $\text{HCl} + \text{Aq}$; insol. in dil. $\text{HCl} + \text{Aq}$. (Arfvedson, Pogg. 1. 65; Gautier, C. R. 108. 1111.)

Does not exist. (Bornemann, C. A. 1908. 1686.)

Nickel monosulphide, NiS .

Anhydrous. Insol. in H_2O , HCl , or $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in $\text{HNO}_3 + \text{Aq}$ or aqua regia.

Min. *Millerite*.

$+x\text{H}_2\text{O}$. Insol. in H_2O , but decomp. by H_2O in contact with the air (Clermont and Guiot, C. R. 84. 714), or by boiling with H_2O . (Geitner, A. 139. 354.)

When pptd. with $(\text{NH}_4)_2\text{S}$, is somewhat sol. in H_2O . 1 l. H_2O dissolves 39.87×10^{-6} moles NiS at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

Very sl. sol. in dil. $\text{HCl} + \text{Aq}$, and still less in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Fresenius.)

More sol. in $\text{HNO}_3 + \text{Aq}$, and easily in aqua regia.

Somewhat sol. in $\text{NH}_4\text{OH} + \text{Aq}$ or solutions of alkali sulphides. Insol. in $\text{NH}_4\text{SH} + \text{Aq}$. (Fresenius.)

Sol. at moment of formation in Na_2S but not in $(\text{NH}_4)_2\text{S} + \text{Aq}$. (Villiers, C. R. 1894, 119. 1264.)

Sol. while yet moist in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Berthier.)

When recently pptd., sol. in $\text{KCN} + \text{Aq}$. (Haidlen.)

Pptd. in presence of non-volatile substances as tartaric acid, etc. (Ro

Sol. in potassium thiocarbonate- (Rosenbladt, Z. anal. 26. 15.)

Exists in a colloidal form in a solution. (Winnsinger, Bull. Soc. (2) a modification:

Very sol. in $2\text{N-HCl} + \text{Aq}$ sat. with β modification:

0.033 g. is sol. in 1 l. 2N-HCl with H_2S ; very sol. in $2\text{N-HCl} + \text{Aq}$ γ modification:

Insol. in $2\text{N-HCl} + \text{Aq}$ sat. w 0.013 g. is sol. in $2\text{N-HCl} + \text{Aq}$. C. C. 1914, I. 19.)

Nickel sulphide, Ni_3S_2 .

(Bornemann, C. A. 1908. 1686.)

Ni_2S_4 . (Bornemann.)

Ni_3S_5 . (Bornemann.)

Ni_4S_6 . Min. *Polydymite*. Insol. Aq. Sol. in $\text{HNO}_3 + \text{Aq}$ with separa Ni_5S_7 . Min. *Beyrichite*. Sol. in l

Nickel disulphide, NiS_2 .

(Fellenberg, Pogg. 50. 75.)

Does not exist. (Bellucci, C. A. 1

Nickel potassium sulphide, $3\text{NiS}, \text{K}$

Insol. in H_2O . (Schneider, J. 1 209.)

$\text{K}_2\text{Ni}_{11}\text{S}_{10}$. Not attacked by hot slowly attacked by HCl or cold aq quickly by hot aqua regia. HF at dissolve only on heating. Insol. in acids, alkalies and 12% HCl , also AgNO_3 or $\text{CuSO}_4 + \text{Aq}$. (Milbauer, 1904, 42. 447.)

Nickel telluride, Ni_2Te_3 .

Min. *Melonite*. Sol. in $\text{HNO}_3 + \text{A}$ NiTe . (Fabre, C. R. 105. 277.)

Nickelicotungstic acid.

Ammonium nickelicotungstate, 2
 $2\text{Ni}_2\text{O}_3, 8\text{WO}_3 + 14\text{H}_2\text{O}$.

(Rogers and Smith, J. Am. Ch 1904, 26. 1476.)

$3(\text{NH}_4)_2\text{O}, \text{Ni}_2\text{O}_3, 16\text{WO}_3 + 22\text{H}_2\text{O}$ sl. sol. in H_2O . (Rogers and Smith.

Barium nickelicotungstate, 19BaO
 16WO_3 .

Ppt. Very insol. in H_2O . (E. F.

Nickelimolybdic acid.

Barium nickelimolybdate, 3BaO ,
 $9\text{MoO}_3 + 12\text{H}_2\text{O}$.

Ppt. (Hall, J. Am. Chem. Soc. 702.)

nickelomolybdate, $3K_2O, NiO_2, 3H_2O + 6\frac{1}{2}H_2O$.

sol. even in hot H_2O . (Hall.)

molybdic acid.

nickelomolybdate, $[Ni(MoO_4)_2] + 5H_2O$.

in H_2O , easily in dil. acids. (Barbieri. 1915. 897.)

hydrogen nickelomolybdate, $[Ni(MoO_4)_2] + 10H_2O$.

(Barbieri.)

hydrogen nickelomolybdate, $[Ni(MoO_4)_2] + 5H_2O$.

in H_2O , easily in acids. (Barbieri.)

hydrogen nickelomolybdate, $[Ni(MoO_4)_2] + 3H_2O$.

in H_2O ; sol. in NH_4OH , or HNO_3 + (Barbieri.)

nickelous acid.

nickelonickelite, $K_2Ni_2O_4$ or K_2O, NiO_2 .

Ann and Hiendlmaier, B. 1906, 39.

nickelonickelite, $Na_2Ni_2O_4 = Na_2O, 2NiO_2$.

Sci and Rubegni, C. C. 1907, I. 794.)

is acid.

nickelite, $BaO, 2NiO_2$.

decomp. by cold H_2O ; slowly rapidly by hot H_2O . (Dufau, C. R. 1896.)

Nb.

niobium and its compounds, see columbium, and the corresponding compounds.

nitrate, NH_4NO_3 .

decomp. by conc. H_2SO_4 . Easily sol. in alcohol, ether and acetone. Less sol. in benzene. Almost insol. in ligroin. (Thiele and Mann, A. 1895, 288. 297.)

insol. in petroleum ether. Stable; decomp. by hot H_2O . (Thiele and Mann, B. 1894, 27. 1909.)

chloroplatinamine comps.

chloronitratoplatinamine comps.

cobalt octamine comps.

nitratooctamine cobaltic comps.

Nitratooctamine cobaltic carbonate,

$(NO_3)_2Co_2(NH_3)_8(CO_3)_2 + H_2O$.

Less sol. than other octamine carbonates. (Vortmann and Blasberg, B. 22. 2650.)

— **chloride**, $(NO_3)_2Co_2(NH_3)_8Cl_4 + 4H_2O$.

(Vortmann and Blasberg, B. 22. 2652.)

— **iodide**, $(NO_3)_2Co_2(NH_3)_8I_4 + 2H_2O$.

(Vortmann and Blasberg.)

— **nitrate**.

See Octamine cobaltic nitrate.

— **sulphate**, $(NO_3)_2Co_2(NH_3)_8(SO_4)_2 + 2H_2O$.

+ $4H_2O$. (Vortmann and Blasberg, B. 22. 2652.)

Nitratoplatinamine nitrate,

$(NO_3)_2Pt(NH_3NO_2)_2$.

Sl. sol. in cold, more easily in hot H_2O ; easily sol. in dil. HNO_3 + Aq. (Cleve.)

— **nitrite**, $(NO_3)_2Pt(NH_3NO_2)_2$.

Easily sol. in H_2O . (Cleve.)

Nitratoplatindiamine chloride,

$(NO_3)_2Pt(N_2H_4Cl)_2 + H_2O$.

Moderately sol. in cold, very easily in hot H_2O .

— **chloroplatinate**, $(NO_3)_2Pt(N_2H_4Cl)_2, PtCl_4 + 2H_2O$.

Ppt.

— **chromate**, $(NO_3)_2Pt(N_2H_4)_2CrO_4$.

Nearly insol. in H_2O . (Cleve.)

— **dichromate**, $(NO_3)_2Pt(N_2H_4)_2Cr_2O_7$.

Sl. sol. in H_2O .

— **nitrate**, $(NO_3)_2Pt(N_2H_4NO_3)_2$.

Sol. in H_2O . Insol. in HNO_3 + Aq.

— **phosphate**, $NO_3Pt(N_2H_4)_2 + H_2O$.

$\backslash /$
 PO_4

Very sl. sol. in H_2O . (Cleve.)

Nitratodiplatindiamine nitrate,

$(NO_3)_2Pt_2(N_2H_4)_4(NO_3)_4$.

Sol. in H_2O with decomp.

Nitratopurpureocobaltic bromide,

$Co(NO_3)(NH_3)_6Br_2$.

Resembles the chloride in its properties. (Jorgensen, J. pr. (2) 23. 227.)

Nitratopurpureocobaltic carbonate,
 $\text{Co}(\text{NO}_3)(\text{NH}_3)_5(\text{CO}_3) + \text{H}_2\text{O}$.

Less sol. in H_2O than other purpureocarbonates. (Vortmann and Blasberg, B. 22. 2648.)

— **chloride,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{Cl}_2$.

Sl. sol. in cold H_2O , but more than nitrate; more easily sol. in hot H_2O , but is converted into roseo salt. Insol. in $\text{HCl} + \text{Aq}$ or alcohol. (Jørgensen, J. pr. (2) 23. 227.)

— **mercuric chloride,**
 $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{Cl}_2, \text{HgCl}_2$.

Not wholly insol. in H_2O . (Jørgensen.)

— **chloroplatinate,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{Cl}_2, \text{PtCl}_4$.

Ppt. Nearly insol. in cold H_2O . (Jørgensen.)

— **chromate,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{CrO}_4$.

Nearly insol. in H_2O . (Jørgensen.)

— **dichromate.**

Sl. sol. in H_2O , but more easily than the neutral salt. (Jørgensen.)

— **dithionate,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{S}_2\text{O}_8$.

Very sl. sol. in cold, more easily in hot H_2O . (Jørgensen.)

— **nitrate,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5(\text{NO}_3)_2$.

Sol. in 273 pts. H_2O at 16° . Much more sol. in hot H_2O containing HNO_3 . (Jørgensen, J. pr. (2) 23. 227.)

— **cobaltic nitrite,** $3\text{Co}(\text{NO}_3)(\text{NH}_3)_5, 2\text{Co}(\text{NO}_2)_4 + 2\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (Jørgensen, Z. anorg. 5. 176.)

— **diamine cobaltic nitrite,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5, (\text{NO}_2)_4\text{Co}(\text{NH}_3)_2$.

Ppt. (Jørgensen.)

— **oxalate,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{C}_2\text{O}_4$.

Ppt.

— **sulphate,** $\text{Co}(\text{NO}_3)(\text{NH}_3)_5\text{SO}_4 + \text{H}_2\text{O}$.

Rather difficultly sol. in cold H_2O . (Jørgensen.)

Nitratopurpureorhodium chloride,
 $(\text{NO}_3)\text{Rh}(\text{NH}_3)_5\text{Cl}_2$.

Sl. sol. in cold H_2O , but more easily than the nitrate. (Jørgensen, J. pr. (2) 34. 394.)

— **dithionate,** $(\text{NO}_3)\text{Rh}(\text{NH}_3)_5\text{S}_2\text{O}_8 + \text{H}_2\text{O}$.

Nearly insol. in cold H_2O . (Jørgensen.)

Nitratopurpureorhodium nitrate,
 $(\text{NO}_3)\text{Rh}(\text{NH}_3)_5(\text{NO}_3)_2$.

Very sl. sol. in cold H_2O . Insol. in alcohol. (Jørgensen.)

Nitric acid, HNO_3 .

Miscible with H_2O . When $\text{HNO}_3 + \text{Aq}$ is distilled at 760 mm. pressure, an acid containing 68% HNO_3 is formed, which boils at 120.5° under 735 mm. pressure. By distilling at 150 mm. pressure the acid contains 67% HNO_3 ; at 70 mm. (b.-pt. $65-70^\circ$) the acid contains 66.7% HNO_3 . The percentage of HNO_3 in the liquid obtained by passing air into $\text{HNO}_3 + \text{Aq}$ containing 64-68% HNO_3 varies with the temp.; the higher the temp. the greater the percentage of HNO_3 . (Royal Chem. Soc. 13. 150.)

$\text{HNO}_3 + \text{Aq}$ of 1.51	sp. gr. contains 67% HNO_3
1.42	54
1.35	44.4
1.315	38.6

(Dalton.)

$\text{HNO}_3 + \text{Aq}$ of 1.52	sp. gr. contains 86% HNO_3
1.522	86.17
1.4	41

(Mitscherlich.)

$\text{HNO}_3 + \text{Aq}$ of 1.298 sp. gr. contains 31.75% HNO_3 (Kirwan.)

$\text{HNO}_3 + \text{Aq}$ of 1.298 sp. gr. contains 18% HNO_3 . (Davy.)

$\text{HNO}_3 + \text{Aq}$ of 1.298 sp. gr. contains 32-33% HNO_3 . (Berthollet.)

For Ure's table of sp. gr. of $\text{HNO}_3 + \text{Aq}$ see Watt's Dict.. 1st ed.

Sp. gr. of $\text{HNO}_3 + \text{Aq}$ at 0° and 15° .

% HNO_3	% N_2O_5	Sp. gr. at 0°	Sp. gr. at 15°
100.00	85.71	1.559	1.530
99.84	85.57	1.559	1.530
99.72	85.47	1.558	1.530
99.52	85.30	1.557	1.529
97.89	83.90	1.551	1.523
97.00	83.14	1.548	1.520
96.00	82.28	1.544	1.516
95.27	81.66	1.542	1.514
94.00	80.57	1.537	1.509
93.01	79.72	1.533	1.506
92.00	78.85	1.529	1.503
91.00	78.00	1.526	1.499
90.00	77.15	1.522	1.495
89.56	76.77	1.521	1.494
88.00	75.43	1.514	1.488
87.45	74.95	1.513	1.486
86.17	73.86	1.507	1.482
85.00	72.86	1.503	1.478
84.00	72.00	1.499	1.474
83.00	71.14	1.495	1.470
82.00	70.28	1.492	1.467
80.96	69.39	1.488	1.463
80.00	68.77	1.484	1.460
79.00	67.71	1.481	1.456
77.66	66.56	1.476	1.451
76.00	65.14	1.469	1.445
75.00	64.28	1.465	1.442
74.01	63.44	1.462	1.438

f HNO₃, etc.—Continued.

$\frac{\%}{\text{N}_2\text{O}_5}$	Sp. gr. at 0°	Sp. gr. at 15°
2.57	1.457	1.435
2.05	1.455	1.432
1.06	1.450	1.429
0.00	1.444	1.423
9.31	1.441	1.419
8.29	1.435	1.414
7.43	1.430	1.410
6.57	1.425	1.405
5.77	1.420	1.400
4.84	1.415	1.395
4.50	1.413	1.393
3.14	1.404	1.386
2.46	1.400	1.381
1.43	1.393	1.374
1.08	1.391	1.372
0.47	1.387	1.368
9.71	1.382	1.363
8.86	1.376	1.358
8.08	1.371	1.353
7.14	1.365	1.346
6.29	1.359	1.341
6.12	1.358	1.339
5.40	1.353	1.335
4.85	1.349	1.331
3.70	1.341	1.323
2.83	1.334	1.317
2.00	1.328	1.312
1.14	1.321	1.307
0.44	1.315	1.308
9.97	1.312	1.295
8.57	1.300	1.284
7.31	1.291	1.274
6.00	1.280	1.264
5.14	1.274	1.257
4.28	1.267	1.251
3.43	1.260	1.244
2.53	1.253	1.237
0.86	1.240	1.225
9.99	1.234	1.218
9.02	1.226	1.211
7.43	1.214	1.198
6.57	1.207	1.192
5.71	1.200	1.185
4.85	1.194	1.179
4.00	1.187	1.172
3.14	1.180	1.166
2.04	1.171	1.157
9.71	1.163	1.138
7.14	1.132	1.120
4.97	1.115	1.105
2.85	1.099	1.089
1.14	1.085	1.077
9.77	1.075	1.067
8.62	1.050	1.045
3.42	1.026	1.022
1.71	1.013	1.010
0.00	1.000	0.999

Sp. gr. of HNO₃+Aq at 15° a = %; b = sp. gr. if % is N₂O₅; c = sp. gr. if % is HNO₃.

a	b	c	a	b	c
1	1.007	1.006	51	1.372	1.323
2	1.014	1.012	52	1.378	1.329
3	1.021	1.018	53	1.385	1.335
4	1.027	1.024	54	1.390	1.341
5	1.034	1.029	55	1.396	1.346
6	1.040	1.035	56	1.401	1.356
7	1.047	1.040	57	1.407	1.358
8	1.053	1.045	58	1.413	1.363
9	1.061	1.051	59	1.418	1.369
10	1.069	1.057	60	1.423	1.374
11	1.076	1.064	61	1.427	1.380
12	1.083	1.070	62	1.432	1.386
13	1.091	1.077	63	1.436	1.390
14	1.098	1.083	64	1.440	1.395
15	1.104	1.089	65	1.445	1.400
16	1.112	1.095	66	1.449	1.405
17	1.120	1.100	67	1.452	1.410
18	1.126	1.106	68	1.457	1.414
19	1.134	1.112	69	1.461	1.419
20	1.141	1.120	70	1.466	1.422
21	1.149	1.126	71	1.470	1.427
22	1.156	1.132	72	1.474	1.430
23	1.165	1.138	73	1.478	1.435
24	1.172	1.145	74	1.482	1.439
25	1.180	1.151	75	1.486	1.442
26	1.187	1.159	76	1.490	1.445
27	1.195	1.166	77	1.494	1.449
28	1.202	1.172	78	1.499	1.452
29	1.211	1.179	79	1.503	1.456
30	1.218	1.185	80	1.507	1.460
31	1.225	1.192	81	1.511	1.463
32	1.232	1.198	82	1.515	1.467
33	1.240	1.204	83	1.519	1.470
34	1.248	1.210	84	1.523	1.474
35	1.255	1.218	85	1.527	1.478
36	1.264	1.225	86	1.530	1.481
37	1.271	1.230	87		1.484
38	1.280	1.236	88		1.488
39	1.286	1.244	89		1.491
40	1.295	1.251	90		1.495
41	1.304	1.257	91		1.499
42	1.312	1.264	92		1.503
43	1.318	1.270	93		1.506
44	1.325	1.276	94		1.509
45	1.332	1.284	95		1.512
46	1.340	1.290	96		1.516
47	1.346	1.298	97		1.520
48	1.352	1.304	98		1.523
49	1.360	1.312	99		1.526
50	1.366	1.316	100		1.530

(Kolb, calculated by Gerlach, Z. anal. 8. 292.)

Sp. gr. of HNO₃+Aq at 17.5°.

$\frac{\%}{\text{N}_2\text{O}_5}$	Sp. gr.	$\frac{\%}{\text{N}_2\text{O}_5}$	Sp. gr.	$\frac{\%}{\text{N}_2\text{O}_5}$	Sp. gr.
■	1.032	9	1.060	13	1.069
6	1.038	10	1.068	14	1.096
7	1.045	11	1.075	15	1.104
8	1.053	12	1.082	■	1.111

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Sp. gr. of HNO_3 , etc.—Continued.

% N_2O_5	Sp. gr.	% N_2O_5	Sp. gr.	% N_2O_5	Sp. gr.
17	1.118	40	1.294	63	1.434
18	1.125	41	1.301	64	1.438
19	1.132	42	1.308	65	1.442
20	1.140	43	1.315	66	1.447
21	1.147	44	1.323	67	1.451
22	1.115	45	1.330	68	1.456
23	1.163	46	1.338	69	1.460
24	1.170	47	1.345	70	1.465
25	1.178	48	1.352	71	1.469
26	1.186	49	1.358	72	1.472
27	1.194	50	1.364	73	1.476
28	1.201	51	1.371	74	1.480
29	1.209	52	1.377	75	1.484
30	1.217	53	1.383	76	1.488
31	1.224	54	1.389	77	1.492
32	1.232	55	1.394	78	1.496
33	1.239	56	1.400	79	1.500
34	1.247	57	1.406	80	1.504
35	1.255	58	1.412	81	1.508
36	1.263	59	1.416	82	1.512
37	1.271	60	1.421	83	1.516
38	1.279	61	1.426	84	1.519
39	1.287	62	1.430	85	1.523

(Hager, Comm. 1883.)

Sp. gr. HNO_3 + Aq at 17.5° .

% N_2O_5	Sp. gr.	% N_2O_5	Sp. gr.	% N_2O_5	Sp. gr.
10	1.068	50	1.293	70	1.465
15	1.104	55	1.361	80	1.500
20	1.140	60	1.417	85	1.514
30	1.217				

(Hager, Adjumenta varia, Leipzig, 1876.)

Sp. gr. of HNO_3 + Aq at 15° .

% HNO_3	Sp. gr.	% HNO_3	Sp. gr.
1	1.00581	26	1.15869
2	1.01136	27	1.16660
3	1.01713	28	1.17371
4	1.02286	29	1.18073
5	1.02851	30	1.18830
6	1.03439	31	1.19552
7	1.04019	32	1.20276
8	1.04592	33	1.20635
9	1.05234	34	1.21300
10	1.05746	35	1.22013
11	1.06330	36	1.22675
12	1.06951	37	1.23347
13	1.07581	38	1.23980
14	1.08126	39	1.24510
15	1.08843	40	1.25235
16	1.09500	41	1.25850
17	1.10102	42	1.26475
18	1.10725	43	1.27125
19	1.11321	44	1.28895
20	1.12024	45	1.28450
21	1.12714	46	1.29110
22	1.13349	47	1.29780
23	1.13890	48	1.30443
24	1.14460	49	1.31101
25	1.15164	50	1.31722

(Squire, Pharm. Era, Jan. 1891.)

Most accurate table.

Sp. gr. of HNO_3 + Aq at 15° ; H_2

Sp. gr.	% N_2O_5	% HNO_3
1.000	0.08	0.10
1.005	0.85	1.00
1.010	1.62	1.90
1.015	2.39	2.80
1.020	3.17	3.70
1.025	3.94	4.60
1.030	4.71	5.50
1.035	5.47	6.38
1.040	6.22	7.26
1.045	6.97	8.13
1.050	7.71	8.99
1.055	8.43	9.84
1.060	9.15	10.68
1.065	9.87	11.51
1.070	10.57	12.33
1.075	11.27	13.15
1.080	11.96	13.95
1.085	12.64	14.74
1.090	13.31	15.53
1.095	13.99	16.32
1.100	14.67	17.11
1.105	15.34	17.89
1.110	16.00	18.67
1.115	16.67	19.45
1.120	17.34	20.23
1.125	18.00	21.00
1.130	18.66	21.77
1.135	19.32	22.54
1.140	19.98	23.31
1.145	20.64	24.08
1.150	21.29	24.84
1.155	21.94	25.60
1.160	22.60	26.36
1.165	23.25	27.12
1.170	23.90	27.88
1.175	24.54	28.63
1.180	25.18	29.38
1.185	25.83	30.13
1.190	26.47	30.88
1.195	27.10	31.62
1.200	27.74	32.36
1.205	28.56	33.09
1.210	28.99	33.82
1.215	29.61	34.55
1.220	30.24	35.28
1.225	30.88	36.03
1.230	31.53	36.78
1.235	32.17	37.53
1.240	32.82	
1.245	33.47	39.05
1.250	34.13	39.82
1.255	34.78	40.58
1.260	35.44	41.34
1.265	36.09	42.10
1.270	36.75	42.87
1.275	37.41	43.64
1.280	38.07	44.41
1.285	38.73	45.18
1.290	39.39	45.95
1.295	40.05	

NO₃, etc.—Continued.

	% HNO ₃	Kg. HNO ₃ in 1 l.
1	47.49	0.617
7	48.26	0.630
3	49.07	0.643
3	49.89	0.656
7	50.71	0.669
7	51.53	0.683
9	52.37	0.697
2	53.22	0.710
5	54.07	0.725
7	54.93	0.739
2	55.79	0.753
7	56.66	0.768
5	57.57	0.783
3	58.48	0.798
1	59.39	0.814
9	60.30	0.829
2	61.27	0.846
5	62.24	0.862
9	63.23	0.879
7	64.25	0.896
7	65.30	0.914
2	66.40	0.933
3	67.50	0.952
3	68.63	0.971
3	69.80	0.991
1	70.98	1.011
3	72.17	1.032
	73.39	1.053
	74.68	1.075
3	75.98	1.098
1	77.28	1.121
3	78.60	1.144
3	79.98	1.168
9	81.42	1.193
3	82.90	1.219
9	84.45	1.246
3	86.05	1.274
3	87.70	1.302
9	89.60	1.335
3	91.60	1.369
3	94.09	1.411
9	94.60	1.420
9	95.08	1.428
	95.55	1.436
9	96.00	1.444
3	96.39	1.451
1	96.76	1.457
3	97.13	1.464
3	97.50	1.470
7	97.84	1.476
9	98.10	1.481
3	98.32	1.486
3	98.53	1.490
3	98.73	1.494
7	98.90	1.497
2	99.07	1.501
4	99.21	1.504
5	99.34	1.507
6	99.46	1.510
5	99.57	1.512
4	99.67	1.515

Sp. gr. of N-HNO₃+Aq at 18°/4° = 1.0324.
(Loomis, W. Ann. 1896, 60. 550.)

Sp. gr. (reduced to a vacuum) of HNO₃ from
78–100% concentration at 4°/4°,
14.2°/4° and 24.2°/4°.

% HNO ₃	Sp. gr.		
	4°/4°	14.2°/4°	24.2°/4°
78.22	1.47129	1.45504	1.43964
79.14	1.46011	1.44372
79.59	1.47496
81.97	1.48391	1.46680	1.45092
84.90	1.49495
85.21	1.49581
85.80	1.47826	1.46224
87.55	1.50211
87.90	1.48491	1.46891
89.73	1.50898	1.49125
92.34	1.51804	1.49968	1.48264
94.04	1.51949	1.50149	1.48516
95.62	1.52192	1.50358	1.48677
96.64	1.52510	1.50632	1.48887
97.33	1.50911	1.49137
98.07	1.53212	1.51298	1.49543
99.97	1.54212	1.52236	1.50394

(Veley and Manley, Chem. Soc. 1903, 83.
1016.)

Sp. gr. at 20° of HNO₃+Aq containing M
g. mols. HNO₃ per liter.

M 0.025 0.05 0.075 0.10
Sp. gr. 1.000926 1.001798 1.002653 1.003498

M 0.25 0.5 0.75 1.0
Sp. gr. 1.008481 1.01686 1.02503 1.0336

M 2.0
Sp. gr. 1.0670

(Jones and Pearce, Am. Ch. J. 1907, 38. 732.)

For sp. gr. of HNO₃+H₂SO₄, see under
H₂SO₄.

Partition coefficient for HNO₃ between
ether and H₂O is increased by the addition
of nitrates. (Tanret, C. R. 1897, 124. 464.)

The hydrates described by Erdmann do
not exist. There are only two authentic hy-
drates, the mono- and the tri-hydrate.
(Küster, Ch. Z. 1904, 28. 132.)

The composition of the hydrates formed
by HNO₃ at different dilutions is calculated
from determinations of the lowering of the
fr.-pt. produced by HNO₃ and of the con-
ductivity and sp. gr. of HNO₃+Aq. (Jones,
Am. Ch. J. 1905, 34. 328.)

Dinitric acid, H₂N₄O₁₁=2N₂O₅, H₂O.

Fumes on air. Miscible with H₂O, with
evolution of much heat. (Weber, J. pr. (2)
6. 342.)

Nitrates.

All nitrates are sol. in H₂O except a few
basic compounds. Most nitrates are insol. in

conc. $\text{HNO}_3 + \text{Aq}$; many are sol. in alcohol; some are sol. in glycerine.

Aluminum nitrate, basic, $2\text{Al}_2\text{O}_3, 3\text{N}_2\text{O}_5 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Ordway, Sill. Am. J. (2) 26. 203.)

Basic aluminum nitrates containing 2 mols. or less of Al_2O_3 to one of N_2O_5 may be obtained sol. in H_2O , but the compounds containing more than 2 mols. Al_2O_3 are insol. in H_2O . (Ordway, l. c.)

$2\text{Al}_2\text{O}_3, \text{N}_2\text{O}_5 + 10\text{H}_2\text{O}$. (Ditte, C. R. 110. 782.)

$\text{Al}_2\text{O}_3 \cdot \text{H}_2\text{O}, \text{HNO}_3$. Sol. in H_2O . (Schlumberger, Bull. Soc. 1895, (3) 13. 59.)

Aluminum nitrate, $\text{Al}(\text{NO}_3)_3 + 9\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O , $\text{HNO}_3 + \text{Aq}$, or alcohol. (Berselius.)

Melts in its crystal H_2O at 72.7° . (Ordway.)

Sol. in 1 pt. strong alcohol. (Wenzel.)

Difficultly sol. in acetone. (Naumann, B. 1904, 87. 4328.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Ammonium nitrate, NH_4NO_3 .

Deliquescent.

Sol. in 0.502 pt. H_2O at 18° . (Karsten.)

Sol. in 0.54 pt. H_2O at 10° . (Harris, C. R. 24. 816.)

Much more sol. in hot than cold H_2O . (Harris.)

Sol. in 2 pts. H_2O at 15.5° and 0.5 pt. boiling H_2O . (Fourcroy.)

Sol. in 1 pt. cold, and 0.5 pt. boiling H_2O . (Fourcroy.)

Sol. in 0.5 pt. H_2O at 18° . (Berselius.)

Sol. in 2 pts. H_2O at 18° . (Abl.)

Decomp. by boiling H_2O .

Solubility in 100 pts. H_2O at t° .

t°	Pts. NH_4NO_3	t°	Pts. NH_4NO_3	t°	Pts. NH_4NO_3
0	97	24	205	48	351
1	101	25	210	49	358
2	105	26	216	50	365
3	109	27	221	51	372
4	113	28	226	52	379
5	117	29	232	53	387
6	121	30	238	54	395
7	125	31	244	55	402
8	130	32	250	56	410
9	134	33	256	57	418
10	139	34	262	58	425
11	143	35	268	59	433
12	148	36	274	60	441
13	152	37	280	61	449
14	157	38	286	62	457
15	161	39	292	63	465
16	166	40	298	64	473
17	170	41	304	65	481
18	175	42	311	66	490
19	180	43	317	67	499
20	185	44	324	68	508
21	190	45	331	69	517
22	195	46	337	70	526
23	200	47	344

(Mulder, Scheik. Verhandel. 1864. 95.)

100 pts. H_2O dissolve 183 pts. NH_4N 19.5° . (Mulder.)

Solubility in H_2O at t° .

t°	Specific gravity of the saturated solution	Mols. of N soluble in mols. of H_2O
12.2	1.2945	34.9
20.2	1.3116	43.7
23.0	1.3159	46.5
25.0	1.3197	48.1
27.7	1.3257	51.6
28.0	1.3260	51.8
30.0	1.3299	54.4
30.2	1.3306	54.6
31.9	1.3348	57.2
32.1	1.3344	57.1
32.7	1.3356	57.1
34.0	1.3375	58.1
35.0	1.3394	59
35.1	1.3397	59
35.6	1.3408	60
36.0	1.3412	61
36.6	1.3420	..
37.5	1.3432	62
38.0	1.3438	63
38.5	1.3440	64
39.0	1.3448	65
39.5	1.3460	65
40.0	1.3464	66

(Müller and Kaufmann, Z. phys. Cl 42. 499.)

Solubility in H_2O at t° .

t°	G. NH_4NO_3 per 100 g.		Solid phase
	solution	water	
0	54.19	118.3	NH_4NO_3 rhomb.
12.2	60.53	153.4	"
20.2	65.80	192.4	"
25.0	68.17	214.2	"
30.0	70.73	241.8	"
32.1	71.97	256.9	NH_4NO_3 rhomb. & rhomb. c.
35	72.64	265.8	NH_4NO_3 rhomb.
40	74.82	297.0	"
50	77.49	344.0	"
60	80.81	421.0	"
70	83.32	499.0	"
80	85.25	580.0	"
90	88.08	740.0	NH_4NO_3 rhomb.
100	89.71	871.0	"

(Seidell's Solubilities 1st ed. 28. Ca Müller & Kauffmann, see also Schwarz, Ostwald's Lehrb., p. 4)

100 g. $\text{NH}_4\text{NO}_3 + \text{Aq}$ contain:

54.19 g. NH_4NO_3 at 0° .

70.10 g. " " 30° .

84.03 g. " " 70° .

(de Waal, Dissert. Leiden, 1881)

g. NH_4NO_3 are contained in 100 g. H_2O + Aq sat. at 30° . (Schreinemakers Baat, Arch. neer. Sc. 1911, (2) 15.)

h. NH_4NO_3 mixed with 100 pts. H_2O as temperature from 13.6° to -13.6° , 37.2° , but if the initial temperature is ill fall only to -16.7° , the freezing-
the mixture. (Rüdorff, B. 2. 68.)

Sp. gr. of NH_4NO_3 + Aq at 18° .

H_2NO_3	Pts. H_2O	Sp. gr.
0	1800	1.0180
0	900	1.0331
0	360	1.0743

nsen and Gerlach, Z. anal. 28. 520.)

Sp. gr. of NH_4NO_3 + Aq at 15° .

NO_3	Sp. gr.	% NH_4NO_3	Sp. gr.
	1.0201	30	1.1304
	1.0419	40	1.1780
	1.0660	50	1.2279

(Kohlrausch, W. Ann. 1879. 1.)

p. gr. of NH_4NO_3 + Aq at 17.5° .

NO_3	Sp. gr.	% NH_4NO_3	Sp. gr.
	1.0042	33	1.1454
	1.0085	34	1.1502
	1.0127	35	1.1550
	1.0170	36	1.1598
	1.0212	37	1.1646
	1.0255	38	1.1694
	1.0297	39	1.1742
	1.0340	40	1.1790
	1.0382	41	1.1841
	1.0425	42	1.1892
	1.0468	43	1.1942
	1.0512	44	1.1994
	1.0555	45	1.2045
	1.0599	46	1.2096
	1.0642	47	1.2147
	1.0686	48	1.2198
	1.0729	49	1.2249
	1.0773	50	1.2300
	1.0816	51	1.2353
	1.0860	52	1.2407
	1.0905	53	1.2460
	1.0950	54	1.2514
	1.0995	55	1.2567
	1.1040	56	1.2621
	1.1085	57	1.2674
	1.1130	58	1.2728
	1.1175	59	1.2781
	1.1220	60	1.2835
	1.1265	61	1.2888
	1.1310	62	1.2942
	1.1358	63	1.3005
	1.1406	64	1.3059

(Gerlach, Z. anal. 27. 310.)

Sp. gr. of NH_4NO_3 + Aq.

% NH_4NO_3	Sp. gr. $16^\circ/16^\circ$
0	1.000000
0.6419	1.000271
1.4101	1.000593
2.7501	1.001153
5.4890	1.002300
11.7981	1.004916
23.4480	1.009758
47.9500	1.019952

(Dijken Z. phys. Ch. 1897, 24. 107.)

Sp. gr. $20^\circ/4^\circ$ of a normal solution of NH_4NO_3 = 1.030435; of a 0.5-normal solution = 1.014505. (Haigh, J. Am. Chem. Soc. 1912, 34. 1151.)

B.-pt. of NH_4NO_3 + Aq containing pts. NH_4NO_3 to 100 pts. H_2O . G = according to Gerlach (Z. anal. 28. 445); L = according to Legrand (A. ch. (2) 59. 426.)

$\frac{p}{m}$	G	L	$\frac{p}{m}$	G	L
101°	10	10	140°	682	770 5
102	20	20.5	141	719	...
103	30	31.3	142	737	840 6
104	41	42.4	143	765	...
105	52	53.8	144	793	915 5
106	63	65.4	145	823	...
107	74	77.3	146	853	995.5
108	85	89.4	147	883	...
109	96	101.9	148	914	1081 5
110	108	114.9	149	945	...
111	120	128.4	150	977	1173 5
112	132	142.4	151	1009	...
113	145	156.9	152	1043	1273
114	158	172	153	1079	...
115	172	188	154	1116	1383
116	187	204.4	155	1155	...
117	202	221.4	156	1196	1504
118	217	238.4	157	1238	...
119	232	256.8	158	1281	1637
120	248	275.3	159	1325	...
121	265	...	160	1370	1775
122	283	314	161	1417	...
123	301	...	162	1464	1923
124	319	354	163	1511	...
125	337	...	164	1558	2084
126	356	396	165	1606	...
127	376	...	166	1653	...
128	396	440 2	167	1700	...
129	417	...	168	1748	...
130	439	487 4	169	1796	...
131	461	...	170	1844	...
132	484	537 3	180	2400	∞
133	507	...	190	3112	...
134	530	590	200	4099	...
135	554	...	210	5618	...
136	578	645	220	8547	...
137	603	...	230	16950	...
138	629	705.5	240	∞	...
139	655

Very sol. in $\text{HNO}_3 + \text{Aq.}$ (Schulz, Zeit. Ch. 1869. 531.)

Solubility of NH_4NO_3 in HNO_3 .

Solution temp.	% by wt. NH_4NO_3	Solid phase
+8°	21.1	$\text{NH}_4\text{NO}_3, 2\text{HNO}_3$ (solution in HNO_3)
23.0	28.7	"
28.5	34.5	"
29.5 *	38.8	(solution in NH_4NO_3)
27.5	44.6	"
27.0	45.8	"
23.5	49.4	"
23.0	50.0	"
17.5	54.0	"
16.5	54.3	"
4.0	45.8	$\text{NH}_4\text{NO}_3, \text{HNO}_3$ labile (solution in HNO_3)
9.5	49.4	"
11.0	51.7	"
11.5	52.7	"
12.0	54.3	"
12.0	54.7	"
11.5	57.6	(solution in NH_4NO_3)
11.5	54.0	NH_4NO_3 (labile)
14.5	54.3	"
17.0	54.7	stable
26.0	55.9	"
27.0	56.2	"
33.5	57.5	"
49.0	60.4	"
79.0	68.1	"

* Mpt. of $\text{NH}_4\text{NO}_3, 2\text{HNO}_3$.
(Groschuff. Z. anorg. 1904, 40. 6.)

Solubility of NH_4NO_3 in $\text{NH}_4\text{OH} + \text{Aq.}$

Grams of NH_4NO_3	Grams of NH_3	Molecules of NH_4NO_3 in 100 molecules $\text{NH}_4\text{NO}_3 + \text{NH}_3$	Temperature at which the solutions are in equilibrium with the solid phase
.....	100.	about 168°
0.7578	0.0588	74.2	109.8
0.6439	0.0665	67.3	94.0
4.2615	0.7747	53.8	68.8
0.7746	0.1857	47.0	35.9
0.9358	0.2352	45.9	33.3
0.7600	0.2607	38.3	0
0.9675	0.3515	36.9	-10.5
0.8308	0.3700	32.3	-30.0
0.9526	1.2457	13.9	-44.5
1.3918	4.4327	6.25	-60
0	100	0	about -80

(Kuriloff, Z. phys. Ch. 1898, 25. 109.)

$\text{NH}_4\text{NO}_3 + \text{NH}_4\text{Cl}$.

100 pts. H_2O dissolve 29.1 pts. NH_4Cl and 173.8 pts. NH_4NO_3 . (Rüdorff, B. 6. 484.)

Sol. in sat. $\text{NH}_4\text{Cl} + \text{Aq}$ with pptn. of NH_4Cl until a state of equilibrium is reached. (Karsten.)

Addition of KClO_3 to $\text{NH}_4\text{Cl} + \text{Aq}$ pptn. of NH_4Cl , and dissolves any that may have been pptd. (Marguer R. 38. 306.)

See also under Ammonium chloride. $\text{NH}_4\text{NO}_3 + \text{KNO}_3$.

100 pts. H_2O dissolve:

	At 9°		At 11°		A
	(1)	(2)	(3)	(4)	(5)
KNO_3 ...	20.2	40.6		26.0	46.1
NH_4NO_3	88.8	143	130.4

2, Sat. at 11° with NH_4NO_3 and then with KNO_3 ; 5, sat. at 11° with NH_4NO_3 then at 15° with KNO_3 . (Mulder.)

Sol. in sat. $\text{HNO}_3 + \text{Aq}$ without c... (Karsten); with separation of KNO_3 (R...

Composition of solution is dependent relative excess of the salts present. (R...

100 pts. H_2O dissolve 77.1 pts. NaNO_3 162.9 pts. NH_4NO_3 at 16°. (Rüdorff 484.)

If a sat. solution of $\text{NH}_4\text{NO}_3 + \text{Aq}$ at sat. with $\text{Ba}(\text{NO}_3)_2$ at 9°, 100 p dissolve:

	At 11°	
NH_4NO_3	143	101.3
$\text{Ba}(\text{NO}_3)_2$	6.2

(Mulder.)

Solubility of $\text{NH}_4\text{NO}_3 + \text{AgNO}_3$ in H

t°	AgNO_3	NH_4NO_3	Solid phase
-7.3°	47.1	0	Ice + AgNO_3
-10.7	44.52	8.43	"
-14.9	42.0	16.80	Ice + AgNO_3
-14.8	39.51	18.79	+ AgNO_3
-18.7	15.99	37.30	Ice + AgNO_3
-17.4	0	41.2	+ NH_4NO_3
0	50.36	19.59	Ice + NH_4NO_3
18	55.36	22.06	$\text{AgNO}_3, \text{NH}_4\text{NO}_3$
30	58.89	23.42	"
55	63.32	26.12	"
109.6	67.9	32.1	"
0	22.13	44.87	$\text{AgNO}_3, \text{NH}_4\text{NO}_3$
18	27.07	49.22	"
30	29.76	52.50	"
40	32.68	52.22	$\text{AgNO}_3, \text{NH}_4\text{NO}_3$
55	36.60	52.38	"
101.5	47.5	52.5	$\text{AgNO}_3, \text{NH}_4\text{NO}_3$

(Schreinemakers and de Baat, Arch. 1911 (2) 15. 414.)

Solubility of NH_4NO_3 and AgNO_3 in H_2O at 30° .

Composition of the solution		Solid phase
H_2O	AgNO_3	
100	73.0	AgNO_3
99	69.08	"
98	63.27	"
95	58.84	$\text{AgNO}_3 + \text{AgNO}_3 \cdot \text{NH}_4\text{NO}_3$
90	58.93	"
83	57.93	$\text{AgNO}_3 \cdot \text{NH}_4\text{NO}_3$
72	55.32	"
60	52.45	"
47	45.85	"
30	41.09	"
14	35.62	"
0	29.77	$\text{NH}_4\text{NO}_3 + \text{AgNO}_3 \cdot \text{NH}_4\text{NO}_3$
1	29.86	"
9	29.66	"
2	27.75	NH_4NO_3
4	21.31	"
9	12.51	"
0	0	"

Schreinemakers and de Baat, Z. phys. Ch. 1909, 65, 572.)

Solubility of $\text{NH}_4\text{NO}_3 + \text{NaNO}_3$ in H_2O at t° .

g. per 100 g. H_2O		Sp. gr.
NH_4NO_3	NaNO_3	
0	73.33	1.354
105.5	66	1.407
118.4	0	1.264
0	83.9	1.375
24.03	81.21	1.386
42.81	79.34	1.392
64.6	77.00	1.401
110.9	75.81	1.417
152.	75.35	1.428
155.3	75.38	1.429
156.1	60.76	1.405
159	36.50	1.364
160	27.79	1.350
162.3	17.63	1.330
167.4	0	1.298
0	96.12	1.401
220.8	88.31	1.450
232.6	0	1.329

Jeff and Koltunoff, Z. anorg. 1914, 85, 251.)

Solubility of $\text{NH}_4\text{NO}_3 + (\text{NH}_4)_2\text{SO}_4$ in H_2O at 30° .

NH_4NO_3	$(\text{NH}_4)_2\text{SO}_4$	Solid phase
70.1	0	
67.63	2.38	NH_4NO_3
66.93	3.46	"
63.84	4.96	$\text{NH}_4\text{NO}_3 + (\text{NH}_4)_2\text{SO}_4$
58.06	8.22	$3\text{NH}_4\text{NO}_3$
52.76	11.42	$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{NH}_4\text{NO}_3$
49.80	13.27	$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{NH}_4\text{NO}_3 +$
37.20	19.48	$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3$
19.91	28.83	$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3 +$
12.05	34.7	$(\text{NH}_4)_2\text{SO}_4$
0	44.1	"

(Schreinemakers and Haenen, Chem. Weekbl. 1909, 6, 51.)

Solubility of $\text{NH}_4\text{NO}_3 + (\text{NH}_4)_2\text{SO}_4$ in H_2O .

Temp. $= 0^\circ$		
NH_4NO_3	$(\text{NH}_4)_2\text{SO}_4$	Solid phase
0	41.4	
5.61	37.89	$(\text{NH}_4)_2\text{SO}_4$
29.58	41.64	"
29.81	21.33	$(\text{NH}_4)_2\text{SO}_4 + (\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3$
31.04	20.40	$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3$
30.87	20.43	"
31.61	19.50	$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3 + (\text{NH}_4)_2\text{SO}_4$
45.99	9.53	$3\text{NH}_4\text{NO}_3$
49.12	6.00	$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{NH}_4\text{NO}_3$
54.19	0	$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{NH}_4\text{NO}_3 + \text{NH}_4\text{NO}_3$
		NH_4NO_3

Temp. $= 70^\circ$		
NH_4NO_3	$(\text{NH}_4)_2\text{SO}_4$	Solid phase
0	47.81	
11.10	40.81	$(\text{NH}_4)_2\text{SO}_4$
70.15	6.71	"
71.58	5.82	$(\text{NH}_4)_2\text{SO}_4 + (\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3$
73.48	5.14	$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3$
76.01	3.96	$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{NH}_4\text{NO}_3 + (\text{NH}_4)_2\text{SO}_4$
80.25	2.68	$3\text{NH}_4\text{NO}_3$
81.01	2.45	$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{NH}_4\text{NO}_3$
81.38	2.41	"
84.03	0	$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{NH}_4\text{NO}_3 + \text{NH}_4\text{NO}_3$
		NH_4NO_3

(de Waal, Dissert. Leiden, 1910.)

Very easily sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 826.)

1 pt. NH₄NO₃ dissolves in 2.29 pts. alcohol of 66.8% at 25°. (Pohl, W. A. B. 6. 599.)
1 pt. NH₄NO₃ dissolves in 1.1 pt. boiling alcohol. (Wenzel.)

100 pts. absolute methyl alcohol dissolve 17.1 pts. at 20.5°. (de Bruyn, Z. phys. Ch. 10. 783.)

100 g. absolute methyl alcohol dissolve 14.6 g. NH₄NO₃ at 14° and 16.3 g. at 18.5°. (Schiff and Monsacchi, Z. phys. Ch. 1896, 21. 277.)

100 pts. absolute ethyl alcohol dissolve 3.8 pts. at 20.5°. (de Bruyn, Z. phys. Ch. 10. 783.)

100 g. absolute ethyl alcohol dissolve 4.6 g. at 14°. (Schiff and Monsacchi, Z. phys. Ch. 1896, 21. 277.)

Solubility of NH₄NO₃ in H₂O is decreased by presence of ethyl alcohol but increased by presence of methyl alcohol. NH₄NO₃ is only very sl. sol. in abs. ethyl alcohol and the solubility increases slowly with rise in temp.; it is more sol. in abs. methyl alcohol and the solubility increases rapidly with rise in temp. (Fleckenstein, Phys. Zeit. 1905, 6. 419.)

Solubility in methyl alcohol + Aq. at 30°.

% by wt. H ₂ O	% by wt. alcohol	% by wt. NH ₄ NO ₃
29.9	0	70.1
21.6	24.5	53.9
20.6	31.3	48.1
16.5	46.0	37.5
11.5	59.4	29.1
0	83.3	16.7

(Schreinemakers, Z. phys. Ch. 1909, 65. 556.)

Solubility of NH₄NO₃ in ethyl alcohol + Aq. at 30°. Composition of sat. solution.

% by wt. H ₂ O	% by wt. alcohol	% by wt. NH ₄ NO ₃
29.9	0	70.1
26.9	18.6	54.5
23.2	39.3	37.5
18.3	58.5	23.2
11.6	76.5	11.9
5.8	86.2	8.0
0	96.4	3.6

(Schreinemakers, Z. phys. Ch. 1909. 65. 555.)

Solubility of NH₄NO₃ in alcohol

t°	% NH ₄ NO ₃	% Alcohol	% H ₂ O
0	54.19	0	45.
"	42.69	12.70	44.
"	1.96	97.93	0.
30	70.10	0	29.
"	59.83	10.60	29.
"	8.06	85.30	6.
"	3.60	96.51	0
70	84.03	0	15
"	72.37	11.12	16
"	61.11	22.87	16
"	41.25	44.64	14
"	24.71	67.23	8
"	7.51	92.49	(

(de Waal. Dissert. Leiden, 1910

Sp. gr. of alcoholic solution of ? at 15°.

Pts. NH ₄ NO ₃	Pts. alcohol	Sp. gr.
0	100	0.8
2	98	0.8
4	96	0.8
6	94	0.8

(Gerlach, Z. anal. 28. 521.)

Insol. in benzonitrile. (Naumann, 47. 1370.)

Insol. in methyl acetate. (Naum 1909, 42. 3790.)

Very sl. sol. in acetone. (Krug a roy, J. Anal. Ch. 6. 184.)

Sol. in acetone. (Eidmann, C. II. 1014.)

Ammonium hydrogen nitrate, NH₄H

Sol. in H₂O. (Ditte, C. R. 89. 576

Decomp. by H₂O. (Groschuff, B. 1487.)

Ammonium dihydrogen nitrate, NH₄H₂(NO₃)₂.

Sol. in H₂O. (Ditte.)

Solubility in H₂O.

Solution temp.	% by wt. NH ₄ NO ₃	%
—8.0	34.2	
—2.5	34.8	
+3.0	35.4	
3.5	36.0	
19.5	37.4	
25.0	38.1	
29.5 mpt.	38.8	

(Groschuff, Z. anorg. 1904, 40.

cerous nitrate, $3\text{NH}_4\text{NO}_3$,
 $3)_2 + 12\text{H}_2\text{O}$.

luescent. Very sol. in H_2O and
olxmann, J. pr. 84. 78.)

Hygroscopic. Sol. in H_2O .
B. 1900, 33. 3507.)

, $\text{Ce}(\text{NO}_3)_3 + 4\text{H}_2\text{O}$. As above.
A. ch. (4) 30. 64.)

in H_2O .

) dissolve at:

5° 45°

3.8 410.2 g. anhydrous salt,

5.06°

7.4 g. anhydrous salt.

f, Z. anorg. 1905, 45. 98.)

ceric nitrate, $2\text{NH}_4\text{NO}_3$,
 $4)_4$.

n H_2O without decomp. Sol. in
eyer, B. 1900, 33. 2137.)

cohol. (Meyer, Z. anorg. 1901,

in H_2O .

) dissolve at t° :

5.2° 45.3°

1.7 174.9 g. anhydrous salt,

5.60° 122°

3.8 735.4 g. anhydrous salt.

f, Z. anorg. 1905, 45. 94.)

Very deliquescent. (Holz-
84. 78.)

cobalt nitrate.

t. Sol. in H_2O . (Thenard.)

copper nitrate, $2\text{NH}_4\text{NO}_3$,
 $2)_2$.

n H_2O .

didymium nitrate, $2\text{NH}_4\text{NO}_3$,
 $3)_2 + 4\text{H}_2\text{O}$.

deliquescent.

gadolinium nitrate, $2\text{NH}_4\text{NO}_3$,
 $3)_2$.

s in the air. (Benedicks, Z.
22. 407.)

gold (auric) nitrate (Ammonium
ate), $\text{NH}_4\text{Au}(\text{NO}_3)_4$.

deliquescent.

u(NO_3)₆. (Schottländer, A. 217.

lanthanum nitrate, $2\text{NH}_4\text{NO}_3$,
 $3)_2 + 4\text{H}_2\text{O}$.

iescent. Sol. in H_2O . (Marig-

Ammonium magnesium nitrate, $2\text{NH}_4\text{NO}_3$,
 $\text{Mg}(\text{NO}_3)_2$.

Slowly deliquescent. Sol. in 10 pts. H_2O at
12.5°, and much less hot H_2O . (Fourcroy.)

Ammonium mercurous nitrate, $4\text{NH}_4\text{NO}_3$,
 $\text{Hg}_2(\text{NO}_3)_2 + 5\text{H}_2\text{O}$.

Sol. in H_2O . (Pagenstecher, Repert. 14.
188.)

Ammonium nickel nitrate.

Sol. in 3 pts. cold H_2O . (Thénard, Scher.
J. 10. 428.)

Ammonium praseodymium nitrate, $2\text{NH}_4\text{NO}_3$,
 $\text{Pr}(\text{NO}_3)_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (von Scheele, Z. anorg. 1898,
18. 356.)

Ammonium silver nitrate, NH_4NO_3 , AgNO_3 .

Very sol. in H_2O . (Russell and Maskelyne,
Roy. Soc. Proc. 26. 357.)

Sol. in H_2O without decomp. (Schreine-
makers and de Baat, Chem. Weekbl. 1910,
7. 6.)

See also solubility of $\text{NH}_4\text{NO}_3 + \text{AgNO}_3$
under NH_4NO_3 .

Ammonium thorium nitrate, $(\text{NH}_4)_2\text{Th}(\text{NO}_3)_6$.

Sol. in strong HNO_3 . (Meyer, Z. anorg.
1901, 27. 383.)

$\text{NH}_4\text{Th}(\text{NO}_3)_6 + 5\text{H}_2\text{O}$. Sol. in HNO_3 of
sp. gr. 1.25. (Meyer, Z. anorg. 1901, 27. 382.)

Ammonium uranyl nitrate, NH_4NO_3 ,
 $\text{UO}_2(\text{NO}_3)_2$.

Decomp. by H_2O . Sol. in conc. HNO_3 .
(Meyer, B. 1903, 36. 4057.)

Solubility in H_2O at t° .

t°	In 100 pts. by wt. of the solution			Solid phase
	Pts. by wt. UO_2	Pts. by wt. NH_4	Pts. by wt. total salt	
0.5	29.71	2.92	...	Double salt + $\text{UO}_2(\text{NO}_3)_2$
13.5	32.35	3.42	...	"
24.9 a	36.40	3.54	68.72	"
b	36.53	3.54	68.97	"
35.0	42.07	3.44	...	"
59.0	44.37	2.90	...	"
80.7 a	44.90	2.98	78.76	Double salt
b	45.01	2.98	78.79	"

Ammonium uranyl nitrate is decomp. by
 H_2O at temp. below 60°: above 60° it is sol.
in H_2O without decomp. (Rimbach, B. 1904,
37. 475.)

Ammonium nitrate ammonia, $2\text{NH}_4\text{NO}_3$,
 3NH_3 .

Known only as a solution of NH_3 in
 $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Troost, C. R. 94. 789.)

$\text{NH}_4\text{NO}_3, 3\text{NH}_3$. As above.

Ammonium nitrate mercuric chloride,
 $\text{NH}_4\text{NO}_3, 2\text{HgCl}_2$.
Insol. in H_2O . Ether dissolves out HgCl_2 .
(Kosmann, A. ch. (3) 27. 240.)
 $2\text{NH}_4\text{NO}_3, \text{HgCl}_2$. Sol. in H_2O . (Hofmann and Marburg, A. 1899, 305. 199.)

Ammonium nitrate sulphate, $2(\text{NH}_4)_2\text{O}, \text{N}_2\text{O}_5, 2\text{SO}_3, \text{H}_2\text{O}$.
Very hygroscopic and sol. in H_2O . (Friedheim, Z. anorg. 1894, 6. 297.)
 $2\text{NH}_4\text{NO}_3, (\text{NH}_4)_2\text{SO}_4$. (de Waal, Dissert. 1910.)
 $3\text{NH}_4\text{NO}_3, (\text{NH}_4)_2\text{SO}_4$. (de Waal.)
See also solubility of $\text{NH}_4\text{NO}_3 + (\text{NH}_4)_2\text{SO}_4$ under NH_4NO_3 .

Ammonium nitrate melitungstate, $\text{NH}_4\text{NO}_3, 2(\text{NH}_4)_2\text{W}_4\text{O}_{13} + 4\text{H}_2\text{O}$.
Decomposes by recrystallising out of H_2O .
(Marignac, A. ch. (3) 69. 61.)

Antimony nitrate, $\text{Sb}_2\text{O}_5, \text{N}_2\text{O}_5$.
Decomp. by cold H_2O . (Bucholz.)
Aqueous solution sat. at 10° contains 30.4% salt. (Eller.)
Sol. in strong, less sol. in dil. $\text{HNO}_3 + \text{Aq}$.
(Peligot, A. ch. (3) 20. 288.)
Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Barium nitrate, $\text{Ba}(\text{NO}_3)_2$.
Sol. in H_2O with absorption of heat.
100 pts. H_2O at 0° dissolve 5.0 parts $\text{Ba}(\text{NO}_3)_2$. (Gay-Lussac, A. ch. 11. 313.)
100 pts. H_2O at 0° dissolve 5.2 parts $\text{Ba}(\text{NO}_3)_2$. (Mulder.)
 $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ sat. at 20° contains 8.57 pts. $\text{Ba}(\text{NO}_3)_2$ to 100 pts. H_2O , and has 1.0679 sp. gr. (Karsten); sat. at 20° has 1.064 sp. gr., and contains 7.94 pts. $\text{Ba}(\text{NO}_3)_2$ to 100 pts. H_2O . (Michel and Krafft.)

100 pts. H_2O dissolve pts. $\text{Ba}(\text{NO}_3)_2$ at t° .

t°	Pts. $\text{Ba}(\text{NO}_3)_2$	t°	Pts. $\text{Ba}(\text{NO}_3)_2$
0	5.00	52.11	17.97
14.95	8.18	73.75	25.01
17.62	8.54	86.21	29.57
37.87	13.67	101.65	35.18
49.22	17.07

(Gay-Lussac, A. ch. (2) 11. 313.)

Solubility in 100 pts. H_2O at t° .			
t°	Pts. $\text{Ba}(\text{NO}_3)_2$	t°	Pts. $\text{Ba}(\text{NO}_3)_2$
0	5.0	52	17.97
1	5.1	53	18.11
2	5.3	54	18.25
3	5.5	55	18.39
4	5.7	56	18.53
5	6.0	57	18.67
6	6.2	58	18.81
7	6.4	59	18.95
8	6.6	60	19.09
9	6.8	61	19.23
10	7.0	62	19.37
11	7.3	63	19.51
12	7.5	64	19.65
13	7.7	65	19.79
14	7.9	66	19.93
15	8.1	67	20.07
16	8.3	68	20.21
17	8.5	69	20.35
18	8.8	70	20.49
19	9.0	71	20.63
20	9.2	72	20.77
21	9.5	73	20.91
22	9.7	74	21.05
23	9.9	75	21.19
24	10.1	76	21.33
25	10.4	77	21.47
26	10.6	78	21.61
27	10.8	79	21.75
28	11.1	80	21.89
29	11.3	81	22.03
30	11.6	82	22.17
31	11.8	83	22.31
32	12.1	84	22.45
33	12.3	85	22.59
34	12.6	86	22.73
35	12.8	87	22.87
36	13.1	88	23.01
37	13.4	89	23.15
38	13.7	90	23.29
39	14.0	91	23.43
40	14.2	92	23.57
41	14.5	93	23.71
42	14.8	94	23.85
43	15.1	95	23.99
44	15.4	96	24.13
45	15.6	97	24.27
46	15.9	98	24.41
47	16.2	99	24.55
48	16.5	100	24.69
49	16.8	101	24.83
50	17.1	101.9	24.97
51	17.4

(Mulder, calculated from his own experiments, Scheik. Verhandel. 1

Ba(NO₃)₂ + Aq contains % Ba(NO₃)₂ at t°.

	Ba(NO ₃) ₂ %	t°	Ba(NO ₃) ₂ %
	4.3	60.0	16.1
	4.9	73.0	19.4
	5.6	92.0	23.4
	5.6	110.0	27.4
	6.4	132.0	31.8
	7.1	134.0	32.5
	7.7	150.0	34.9
	9.7	152.0	35.4
	12.8	171.0	38.3
	14.9	215.0	45.8

(Étard, A. ch. 1894, (7) 2. 528.)

g. H₂O dissolve 8.54 g. Ba(NO₃)₂ at Gmelin-Kraut, Handbuch der anorg. ch.)

g. H₂O dissolve 7.87 g. Ba(NO₃)₂ at 12 g. at 17°. (Euler, Z. phys. Ch. 1904, .)

g. H₂O dissolve 0.72 gram-equivalents Ba(NO₃)₂ at 21.5°. (Euler, Z. phys. Ch. 1904, 9. 312.)

g. anhydrous Ba(NO₃)₂ are sol. in H₂O at 25°. (Parsons and Colson, J. chem. Soc. 1910, 32. 1385.)

g. Ba(NO₃)₂ are contained in 100 g. Ba(NO₃)₂ sat. at 0°. (Coppadoro, Gazz. ch. 1892, 42, I. 233.)

Solubility of Ba(NO₃)₂ in H₂O = 0.427 mol. at 0°. (Masson, Chem. Soc. 1911, 99.)

Solubility of Ba(NO₃)₂ in H₂O at 30° = 0.427 mol. (Coppadoro, Gazz. ch. it. 1913, 43.)

Solubility in H₂O.

g. of the sat. solution contain at:

t°	g. Ba(NO ₃) ₂
21.1°	8.46
35°	11.39 g. Ba(NO ₃) ₂ .

(Mendley, Chem. Soc. 1914, 105. 780.)

Sp. gr. of Ba(NO₃)₂ + Aq at 19.5°.

Sp. gr.	Ba(NO ₃) ₂ %	Sp. gr.
1.009	6	1.050
1.017	7	1.060
1.025	8	1.069
1.034	9	1.078
1.042	10	1.087

Prepared by Gerlach, Z. anal. 8. 286, from Kremers, Pogg. 95. 110.)

Sp. gr. of Ba(NO₃)₂ + Aq at 18°.

Ba(NO ₃) ₂ %	Sp. gr.
4.2	1.0340
8.4	1.0712

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of Ba(NO₃)₂ + Aq at 17.5°.

Ba(NO ₃) ₂ %	Sp. gr.	Ba(NO ₃) ₂ %	Sp. gr.
1	1.0085	6	1.0510
2	1.0170	7	1.0600
3	1.0255	8	1.0690
4	1.0340	Sat. sol.	1.0690
5	1.0425

(Gerlach, Z. anal. 27. 283.)

Sp. gr. of Ba(NO₃)₂ + Aq at room temp.

% Ba(NO ₃) ₂	Sp. gr.
5.25	1.0507
2.98	1.0274

(Wagner, W. Ann. 1883, 18. 264.)

Sp. gr. of Ba(NO₃)₂ + Aq at 25°.

Concentration of Ba(NO ₃) ₂ + Aq.	Sp. gr.
1/2 normal	1.0518
1/4 "	1.0259
1/8 "	1.0130

(Wagner, Z. phys. Ch. 1890, 5. 35.)

Ba(NO₃)₂ + Aq containing 6.08 % Ba(NO₃)₂ has sp. gr. 20°/20° = 1.0517.

Ba(NO₃)₂ + Aq containing 6.97 % Ba(NO₃)₂ has sp. gr. 20°/20° = 1.0597.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 279.)

Sp. gr. of Ba(NO₃)₂ + Aq at 20° containing M g. mols. salt per liter.

M	0.01	0.025	0.05
Sp. gr.	1.002031	1.005224	1.010591

M	0.075	0.10	0.15
Sp. gr.	1.015671	1.021143	1.031770

(Jones and Pearce, Am. Ch. J. 1907, 38. 708.)

Sp. gr. of sat. Ba(NO₃)₂ + Aq at t°.

t°	g. Ba(NO ₃) ₂ sol. in 100 g. H ₂ O	Sp. gr.
0	5.2	1.043
10	7.0	1.056
20	9.2	1.073
30	11.6	1.087
40	14.2	1.104
50	17.1	1.121
60	20.3	1.137
70	23.6	1.146

(Tschernaj, J. Russ. Phys. Chem. Soc. 1912, 44. 1565.)

Saturated Ba(NO₃)₂ + Aq contains:—
36.18 pts. Ba(NO₃)₂ to 100 pts. H₂O, and boils at 101.1°. (Griffiths.)

35.2 pts. $\text{Ba}(\text{NO}_3)_2$ to 100 pts. H_2O , and boils at 101.65° . (Gay-Lussac.)

34.8 pts. $\text{Ba}(\text{NO}_3)_2$ to 100 pts. H_2O , and boils at 101.9° . (Mulder.)

34.8 pts. $\text{Ba}(\text{NO}_3)_2$ to 100 pts. H_2O , and boils at 102.5° . (Kremers.)

Sat. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ forms a crust at 101.1° ; highest temp. observed was 101.5° . (Gerlach, Z. anal. 26. 427.)

B. pt. of $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ containing pts. $\text{Ba}(\text{NO}_3)_2$ to 100 pts. H_2O .

B.-pt.	Pts. $\text{Ba}(\text{NO}_3)_2$
100.5°	12.5
101.0	26.0
101.1	27.5

(Gerlach, Z. anal. 26. 440.)

Insol. in conc. $\text{HNO}_3 + \text{Aq}$, and much less sol. in dil. $\text{HNO}_3 + \text{Aq}$ or $\text{HCl} + \text{Aq}$ than in H_2O .

Solubility of $\text{Ba}(\text{NO}_3)_2$ in $\text{HNO}_3 + \text{Aq}$ at 30° .

Solid phase $\text{Ba}(\text{NO}_3)_2$		
Sp. gr. of sat. solution	G. mol. per l.	
	HNO_3	$\text{Ba}(\text{NO}_3)_2$
1.0891	0.0000	0.4270
1.0811	0.1318	0.3282
.....	0.2496	0.3268
1.0663	0.4995	0.2410
1.0619	0.7494	0.1785
1.0609	1.000	0.1353
1.0633	1.247	0.1056
1.0668	1.493	0.0847
1.0783	1.998	0.0598
1.1050	2.993	0.0334
1.1341	3.986	0.0218
1.1341	3.994	0.0223
1.1645	5.012	0.0147

(Masson, Chem. Soc. 1911, 99. 1136.)

Less sol. in dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ than in dil. $\text{HCl} + \text{Aq}$.

Solubility in $\text{NH}_4\text{Cl} + \text{Aq}$ is the same as in H_2O .

Less sol. in $\text{NH}_4\text{OH} + \text{Aq}$, $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$, or $\text{NH}_4\text{NO}_3 + \text{Aq}$ than in H_2O . (Pearson, Zeit. Ch. (2) 5. 662.)

$\text{Ba}(\text{NO}_3)_2$ is sol. in about:

13.33 pts. H_2O at ord. temp., and 4.67 pts. at 100° .

14.67 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (conc.) at ord. temp., and 5.67 pts. at 100° .

16.50 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (1 vol. conc. + 3 vols. H_2O) at ord. temp.

28.00 pts. $\text{HCl} + \text{Aq}$ (1 vol. conc. $\text{HCl} + 4$ vols. H_2O) at ord. temp.

29.00 pts. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ (1 vol. commercial $\text{HC}_2\text{H}_3\text{O}_2 + 1$ vol. H_2O) at ord. temp.

13.67 pts. $\text{NH}_4\text{Cl} + \text{Aq}$ (1 pt. $\text{NH}_4\text{Cl} + 10$ pts. H_2O) at ord. temp., and 4.67 pts. at 100° .

24.00 pts. $\text{NH}_4\text{NO}_3 + \text{Aq}$ (1 pt. $\text{NH}_4\text{NO}_3 + 10$ pts. H_2O) at ord. temp.

17.33 pts. $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$ (dil. $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$ neutralised by dil. $\text{HC}_2\text{H}_3\text{O}_2$) at ord. temp., and 4.33 pts. at 100° .

14.67 pts. $\text{NaC}_2\text{H}_3\text{O}_2 + \text{Aq}$ (dil. $\text{HC}_2\text{H}_3\text{O}_2$ neutralised by Na_2CO_3 and dil. with 4 vols. H_2O) at ord. temp., and 5.33 pts. at 100° .

17.33 pts. $\text{Cu}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$ (see Stolba, Z. anal. 2. 390) at ord. temp., and 6.00 pts. at 100° .

18.67 pts. grape sugar (1 pt. grape sugar + 10 pts. H_2O) at ord. temp. (Pearson, Zeit. Ch. 1869. 662.)

Sol. in sat. $\text{NH}_4\text{Cl} + \text{Aq}$ without pptn. at first, but finally NH_4Cl is pptd. until a certain state of equilibrium is reached. (Kersten.)

Solubility in $\text{BaO}_2\text{H}_2, 8\text{H}_2\text{O} + \text{Aq}$ at 25° .

Sp. gr. $25^\circ/25^\circ$	G. BaO as $\text{Ba}(\text{OH})_2$ in 100 g. H_2O	G. $\text{Ba}(\text{NO}_3)_2$ in 100 g. H_2O
1.0797	0	10.30
1.1002	1.55	10.66
1.1210	3.22	11.04
*1.1448	5.02	11.48

* This solution is sat. with respect to $\text{Ba}(\text{OH})_2, 8\text{H}_2\text{O}$ and $\text{Ba}(\text{NO}_3)_2$.

(Parsons. J. Am. Chem. Soc. 1910, 32. 1285)

See also under BaO_2H_2 .

Solubility in $\text{BaCl}_2 + \text{Aq}$ at t° .

t°	Sat. solution contains	
	% BaCl_2	% $\text{Ba}(\text{NO}_3)_2$
—7	21.4	4.0
—1	23.0	4.0
+1.5	22.6	4.4
2	5.0
10	24.7	6.1
21	24.5	5.6
32	26.6	7.7
35	26.4	7.7
38	26.7	7.8
48	28.1	8.0
53	28.5	9.0
53	28.3	9.2
66	28.0	10.0
73	30.0	10.5
79	30.3	11.2
90	32.1	12.5
155	32.5	23.1
162	33.1	23.4
210	32.5	31.9

(Étard, A. ch. 1894, (7) 3. 287.)

See also under BaCl_2 .

$\text{Ba}(\text{NO}_3)_2 + \text{Pb}(\text{NO}_3)_2$.

l. sol. in sat. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$. (Kar-

sten, *Ann. Chem. Phys.* 1897, 28, 365, 397.)

ity of $\text{Ba}(\text{NO}_3)_2 + \text{Pb}(\text{NO}_3)_2$ at 25°.

G. per l.		Sp. gr.
$\text{Ba}(\text{NO}_3)_2$	$\text{Pb}(\text{NO}_3)_2$	
.2	0	1.079
.9	17.63	1.088
.5	49.80	1.108
.7	68.10	1.119
0	97.20	1.140
.8	130.7	1.163
.0	177.3	1.198
.5	247.7	1.252
.9	334.3	1.294
.8	429.7	1.376
	553.8	1.459

Z. Kryst. Min. 1897, 28, 365, 397.)

em. $\text{Ba}(\text{NO}_3)_2 + \text{Pb}(\text{NO}_3)_2 + \text{Aq}$ sat. contain 3.22 g. $\text{Ba}(\text{NO}_3)_2$ and 38.59 g. $\text{Pb}(\text{NO}_3)_2$ and solution has sp. gr. = 1.350. Z. phys. Ch. 1904, 46, 313.)

pts. sat. $\text{Ba}(\text{NO}_3)_2 + \text{Pb}(\text{NO}_3)_2 + \text{Aq}$ contain 45.90 pts. of the three 19-20°. (v. Hauer, l. c.)

$\text{Ba}(\text{NO}_3)_2 + \text{Sr}(\text{NO}_3)_2$.

is. sat. $\text{Ba}(\text{NO}_3)_2 + \text{Sr}(\text{NO}_3)_2 + \text{Aq}$ contain 96 pts. of the two salts at 19-20°. (v. Hauer, l. c.)

$\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3$.

100 pts. H_2O dissolve:

	(Mulder)			
	(1)			
	29.7	28.8	5.4	8.9
$\text{Ba}(\text{NO}_3)_2$	5.4	8.9	
		34.2		
	(Karsten)		(Kopp)	
	(2)	(3)	(4)	(5)
	13.31	29.03	5.7	3.5
KNO_3	6.91	1.00	33.1	36.3
	20.22	30.03	38.8	39.8

l. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ sat. with KNO_3 at

sat. $\text{KNO}_3 + \text{Aq}$, $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ was

sat. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$, KNO_3 was

th salts in excess + Aq at 21.5°.

th salts in excess + Aq at 23°.

1 l. of the solution contains 59.1 g. $\text{Ba}(\text{NO}_3)_2$ + 124.2 g. KNO_3 = 183.35 g. mixed salts at 17°. Sp. gr. $\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3 + \text{Aq}$ = 1.120.

1 l. of the solution contains 88.7 g. $\text{Ba}(\text{NO}_3)_2$ + 213.6 g. KNO_3 = 302.3 g. mixed salts at 30°. Sp. gr. $\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3 + \text{Aq}$ = 1.191. (Euler, Z. phys. Ch. 1904, 49, 313.)

Solubility in $\text{KNO}_3 + \text{Aq}$ at 25°.

100 pts. of solution contain		Solid phase
pts. KNO_3	pts. $\text{Ba}(\text{NO}_3)_2$	
15.24	6.64	$\text{Ba}(\text{NO}_3)_2 + 2\text{KNO}_3$, $\text{Ba}(\text{NO}_3)_2$
14.69	6.60	"
14.79	6.62	"
16.30	5.49	2KNO_3 , $\text{Ba}(\text{NO}_3)_2$
21.99	3.04	"
27.66	2.01	$\text{KNO}_3 + 2\text{KNO}_3$, $\text{Ba}(\text{NO}_3)_2$
27.81	2.09	"
27.94	1.92	"
27.64	2.05	"

These results show that a double salt of potassium and barium nitrates is formed at 25°.

(Foote, Am. Ch. J. 1904, 32, 252.)

Solubility of $\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3$ at t°.

t°	$\text{Ba}(\text{NO}_3)_2$	KNO_3	Solid phase
9.1	6.25	0	$\text{Ba}(\text{NO}_3)_2$
	4.20	8.15	$\text{Ba}(\text{NO}_3)_2 + 2\text{KNO}_3$
	1.98	12.02	$\text{Ba}(\text{NO}_3)_2$
	0.98	16.80	2KNO_3 , $\text{Ba}(\text{NO}_3)_2$
	0	16.76	2KNO_3 , $\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3$
			KNO_3
21.1	8.46	0	$\text{Ba}(\text{NO}_3)_2$
	7.47	2.12	"
	6.35	5.98	"
	6.06	8.47	"
	5.98	13.24	$\text{Ba}(\text{NO}_3)_2 + 2\text{KNO}_3$
	3.35	18.24	$\text{Ba}(\text{NO}_3)_2$
	2.30	21.47	2KNO_3 , $\text{Ba}(\text{NO}_3)_2$
	1.76	24.86	"
	0	24.77	2KNO_3 , $\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3$
			KNO_3
35	11.39	■	$\text{Ba}(\text{NO}_3)_2$
	8.18	12.99	"
	8.08	17.48	"
	8.42	19.75	$\text{Ba}(\text{NO}_3)_2 + 2\text{KNO}_3$
	5.85	24	$\text{Ba}(\text{NO}_3)_2$
	5.02	26.05	2KNO_3 , $\text{Ba}(\text{NO}_3)_2$
	3.02	34.87	"
	1.77	34.98	2KNO_3 , $\text{Ba}(\text{NO}_3)_2 + \text{KNO}_3$
	0	35.01	KNO_3
			"

(Findlay, Chem. Soc. 1914, 106, 779.)

$\text{Ba}(\text{NO}_3)_2 + \text{NaNO}_3$.
 $\text{Ba}(\text{NO}_3)_2$ is sol. in sat. NaNO_3 + Aq without separation.

100 pts. H_2O dissolve:

	(Karsten) At 18.75°		
NaNO_3	86.6	88.14	...
$\text{Ba}(\text{NO}_3)_2$	3.77	8.9

	(Kopp) At 20.2°		
NaNO_3	87.7	88.6	...
$\text{Ba}(\text{NO}_3)_2$	3.6	9.2

Solubility of $\text{Ba}(\text{NO}_3)_2 + \text{NaNO}_3$ in H_2O at 0°.

% NaNO_3	% $\text{Ba}(\text{NO}_3)_2$	Solid phase
0	4.74	$\text{Ba}(\text{NO}_3)_2$
0.41	4.33	"
0.61	4.03	"
1.68	3.34	"
3.54	2.50	"
8.05	1.60	"
12.71	1.56	"
20.24	1.53	"
20.92	1.43	"
27.74	1.56	"
30.81	1.55	"
33.79	1.53	"
35.83	1.49	"
41.30	1.55	$\text{Ba}(\text{NO}_3)_2 + \text{NaNO}_3$
41.68	0.51	NaNO_3
42.47	0	"

(Coppadoro, Gazz. ch. it, 1912, 42 (1) 233.)

Solubility of $\text{Ba}(\text{NO}_3)_2 + \text{NaNO}_3$ in H_2O at 30°.

% NaNO_3	% $\text{Ba}(\text{NO}_3)_2$	Solid phase
0	10.33	$\text{Ba}(\text{NO}_3)_2$
2.33	8.58	"
7.09	5.28	"
12.07	3.89	"
14.41	3.54	"
17.87	3.20	"
19.06	3.07	"
23.55	2.81	"
41.22	2.27	"
48.22	2.11	$\text{Ba}(\text{NO}_3)_2 + \text{NaNO}_3$
48.50	1.00	NaNO_3
49.16	0	"

(Coppadoro, Gazz. ch. it. 1913, 43, I. 240.)

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 827.)
100 pts. hydrazine dissolves 81.1 pts. $\text{Ba}(\text{NO}_3)_2$ at 12.5–13°. (de Bruyn, R. t. c. 1899, 18. 297.)

100 pts. anhydrous hydroxylamine c 11.4 pts. $\text{Ba}(\text{NO}_3)_2$. (de Bruyn, R. 1892, 11. 18.)
Insol. in absolute alcohol.
Solubility in dilute alcohol increases with the temp. (Gerardin, A. ch. (4) 5. 14)

Solubility in ethyl alcohol + Aq at

% $\text{C}_2\text{H}_5\text{OH}$ in the solvent	% $\text{C}_2\text{H}_5\text{OH}$ in the solution	% $\text{Ba}(\text{NO}_3)_2$ in the solution
0	0	9.5
10.25	9.5	7.6
18.60	17.5	6.0
25.05	23.7	5.5
40.20	38.8	3.1
58.00	57.0	1.1
78.70	78.2	0.1
90.10	89.9	0.
99.40	99.39	0.

(D'Ans and Siegler, Z. phys. Ch. 1913

Completely insol. in boiling amyl alcohol. (Browning, Sill. Am. J. 143. 314.)

Solubility in organic solvents

Solvent	% $\text{Ba}(\text{NO}_3)_2$ in solution
Methyl alcohol	0.50
Ethyl alcohol	0.006
Acetone	0.005
Ether	very little
Paracetaldehyde	"

(D'Ans and Siegler, Z. phys. Ch. 1913

Solubility in phenol + Aq at 25°

Concentration of the phenol Mol./ Liter	Solubility of $\text{Ba}(\text{NO}_3)_2$ Mol./l
0.000	0.393
0.045	0.378
0.082	0.374
0.146	0.361
0.310	0.346
0.401	0.341
0.501	0.326
0.728 (sat.)	0.306

(Rothmund and Wilsmore, Z. phys. Ch. 40. 620.)

Insol. in benzonitrile. (Naumann, Z. 47. 1370.)
Insol. in methyl acetate (Naumann, 1909, 42. 3790); ethyl acetate. (B. 1904, 37. 3602.)
Insol. in acetone. (Krug and M. Anal. Ch. 6. 184.)
Difficultly sol. in acetone. (Naumann, 1904, 37. 4328.)
Sol. in acetone. (Eidmann, C. C. 1014.)

Barium mercurous nitrate, 2BaO , $2\text{Hg}_2\text{O}$, $3\text{N}_2\text{O}_5$.

Decomp. by H_2O . Sol. in hot dil. HNO_3 + aq and hot $\text{Hg}_2(\text{NO}_3)_2$ + Aq , from which it crystallises on cooling. (Städeler, A. 87. 129.)

Barium potassium nitrate, $\text{Ba}(\text{NO}_3)_2$, 2KNO_3 .

Ppt. (Wallbridge, Am. Ch. J. 1903, 30. 54.)

Solubility determinations show that the only double salt formed by barium and potassium nitrates at 25° is $\text{Ba}(\text{NO}_3)_2$, 2KNO_3 .

See $\text{Ba}(\text{NO}_3)_2$ + KNO_3 under $\text{Ba}(\text{NO}_3)_2$. Foote, Am. Ch. J. 1904, 32. 252.)

Barium nitrate metatungstate, $2\text{Ba}(\text{NO}_3)_2$, $\text{Ba}_2\text{W}_4\text{O}_{18}$ + $6\text{H}_2\text{O}$.

Efflorescent. Sol. in warm H_2O . (Péchar, A. ch. (6) 22. 198.)

Bismuth nitrate, basic, Bi_2O_3 , N_2O_5 + $2\text{H}_2\text{O}$.

Sol. in a large amount of H_2O . Sol. in HNO_3 + Aq . (Heintz.)

Sol. in 135 pts. H_2O at $90-93^\circ$. (Ruge, J. B. 163.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Sol. in much H_2O . (Yvon, C. R. 84. 1161.)

+ H_2O . (Ruge.)

$2\text{Bi}_2\text{O}_3$, N_2O_5 . Not acted upon by H_2O . (Ditte, C. R. 84. 1317.)

+ H_2O . (Yvon.)

Bi_2O_3 , $2\text{N}_2\text{O}_5$ + H_2O . (Ruge.)

$11\text{Bi}_2\text{O}_3$, $5\text{N}_2\text{O}_5$ + $16\text{H}_2\text{O}$. Not decomp. by H_2O . (Yvon.)

$5\text{Bi}_2\text{O}_3$, $4\text{N}_2\text{O}_5$ + $8\text{H}_2\text{O}$. Ppt. Not attacked by H_2O . (Schulten, Bull. Soc. 1903, (3) 29. 22.)

$5\text{Bi}_2\text{O}_3$, $5\text{N}_2\text{O}_5$ + $9\text{H}_2\text{O}$. Sol. in H_2O with decomp. (Schulten.)

$6\text{Bi}_2\text{O}_3$, $5\text{N}_2\text{O}_5$ + $8\text{H}_2\text{O}$, and + $9\text{H}_2\text{O}$. (Rutten, Z. anorg. 1902, 30. 368.)

At 25° the salt $\text{Bi}_{12}\text{O}_{18}(\text{NO}_3)_{10}$, $9\text{H}_2\text{O}$ is in equilibrium with HNO_3 + Aq from 0.03-0.32-N; the salt $\text{BiO}(\text{NO}_3)$, H_2O is in equilibrium with HNO_3 + Aq from 0.425-0.72-N.

At 50° the salt $\text{Bi}_4\text{O}_6(\text{NO}_3)_2$, H_2O is in equilibrium with HNO_3 + Aq from 0.057-0.285-N; the salt $\text{Bi}_{12}\text{O}_{18}(\text{NO}_3)_{10}$, $9\text{H}_2\text{O}$ is in equilibrium with HNO_3 + Aq from 0.285-0.446-N.

At 75° the salt $\text{Bi}_4\text{O}_6(\text{NO}_3)_2$, H_2O is in equilibrium with HNO_3 + Aq from 0.109-0.314-N. (Allan, Am. Ch. J. 1901, 25. 314.)

Bismuth nitrate, $\text{Bi}(\text{NO}_3)_3$.

Permanent. Decomp. by little H_2O with separation of a basic salt. This decomposition is prevented by slight excess of HNO_3 , and then the salt is completely sol. in a large amount of H_2O . (Rose.)

Sol. in dil. HNO_3 + Aq . Not decomp. by H_2O in presence of $\text{HC}_2\text{H}_3\text{O}_2$ or $\frac{1}{500}$ pt. NH_4NO_3 . (Löwe, J. pr. 74. 341.)

Completely sol. in HNO_3 + Aq containing 83 g. HNO_3 per liter. (Ditte.)

Solubility of $\text{Bi}(\text{NO}_3)_3$ in $2.3\text{N}-\text{HNO}_3$ + Aq . = 2.04 g. at Bi per l.; in $0.922\text{N}-\text{HNO}_3$ + Aq . = 2.23 g. at Bi per l. (Dubrisay, C. R. 1911, 153. 1077.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone. (Krug and M'Elroy.)

Solubility of $\text{Bi}(\text{NO}_3)_3$ in 6.67% acetone + $2.3\text{N}-\text{HNO}_3$ + Aq = 1.89 g. at Bi per l.; in 6.67% acetone + $0.922\text{N}-\text{HNO}_3$ + Aq . = 2.17 g. at Bi per l.; in 13.33% acetone + $0.922\text{N}-\text{HNO}_3$ + Aq = 2.08 g. at Bi per l. (Dubrisay, C. R. 1911, 153. 1077.)

When $\text{Bi}(\text{NO}_3)_3$ is mixed with mannite (dulcite, sorbite) in proportion to the mol. wts. and H_2O is added, a clear solution is obtained which is not pptd. by addition of much H_2O . These solutions are more stable the greater the proportion of mannitol. (Vanino and Hunser, Z. anorg. 1901, 28. 211.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. (Ditte.)

+ $5\text{H}_2\text{O}$. If treated with increasing amts. of H_2O , the amt. of Bi which dissolves decreases, and when 1 pt. is treated with 50,000 pts. H_2O , no Bi goes into solution. (Antony and Gigli, Gazz. ch. it. 1898, 28. 245.)

48.66 pts. are sol. in 100 pts. acetone at 0° .
41.70 " " " " 100 " " " 19° .

(Laszczynski, B. 1894, 27. 2287.)

+ $5\frac{1}{2}\text{H}_2\text{O}$. (Yvon, C. R. 84. 1161.)

+ $10\text{H}_2\text{O}$. Melts in crystal H_2O with decomp. at 74° . (Ordway.)

Bismuth caesium nitrate, $\text{Bi}(\text{NO}_3)_3$, 2CsNO_3 .

Ppt. (Wells, Am. Ch. J. 1901, 26. 277.)

Bismuth cobalt nitrate, $2\text{Bi}(\text{NO}_3)_3$, $3\text{Co}(\text{NO}_3)_2$ + $24\text{H}_2\text{O}$.

100 cc. sat. solution in HNO_3 + Aq (sp. gr. 1.325) contain 54.67 g. hydrated salt. (Jantsch Z. anorg. 1912, 76. 321.)

Bismuth magnesium nitrate, $2\text{Bi}(\text{NO}_3)_3$, $3\text{Mg}(\text{NO}_3)_2$ + $24\text{H}_2\text{O}$.

Deliquescent. Effloresces in dry air. Decomp. by H_2O . (Urbain and Lacombe, C. R. 1903, 137. 569.)

100 cc. sat. solution in HNO_3 + Aq (sp. gr. 1.325) contain 41.69 g. hydrated salt. (Jantsch Z. anorg. 1912, 76. 321.)

Bismuth manganous nitrate, $2\text{Bi}(\text{NO}_3)_3$, $3\text{Mn}(\text{NO}_3)_2$ + $24\text{H}_2\text{O}$.

Deliquescent. Effloresces in dry air. Decomp. by H_2O . (Urbain and Lacombe, C. R. 1903, 137. 569.)

100 cc. sat. solution in HNO_3 + Aq (sp. gr. 1.325) contain 65.77 g. hydrated salt. (Jantsch Z. anorg. 1912, 76. 321.)

Bismuth nickel nitrate, $2\text{Bi}(\text{NO}_3)_3, 3\text{Ni}(\text{NO}_3)_2 + 24\text{H}_2\text{O}$.

Deliquescent. Effloresces in dry air. Decomp. by H_2O . (Urbain and Lacombe, C. R. 1903, 37. 569.)

100 cc. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contain 46.20 g. hydrated salt at 16° . (Jantsch.)

Bismuth zinc nitrate, $2\text{Bi}(\text{NO}_3)_3, 3\text{Zn}(\text{NO}_3)_2 + 24\text{H}_2\text{O}$.

Deliquescent. Decomp. by H_2O . (Urbain and Lacombe, C. R. 1903, 137. 569.)

100 cc. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contain 57.51 g. hydrated salt at 16° . (Jantsch.)

Cadmium nitrate, basic, $\text{Cd}(\text{OH})\text{NO}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O , or ordinary alcohol. (Klinger, B. 16. 997.)

$12\text{CdO}, \text{N}_2\text{O}_5 + 11\text{H}_2\text{O}$. Sl. sol. in H_2O ; more sol. in H_2O than basic sulphate. (Habermann, 5. 432.)

$5\text{CdO}, 2\text{N}_2\text{O}_5 + 8\text{H}_2\text{O}$. Decomp. by cold H_2O . (Rousseau and Tite, C. R. 114. 1184.)

Cadmium nitrate, $\text{Cd}(\text{NO}_3)_2$.

Deliquescent, and very sol. in H_2O .
See +4, and $9\text{H}_2\text{O}$.

Sp. gr. of aqueous solution containing:

5 10 15 20 25% $\text{Cd}(\text{NO}_3)_2$,
1.0528 1.0978 1.1516 1.2134 1.2842

30 35 40 45 50% $\text{Cd}(\text{NO}_3)_2$.
1.3566 1.4372 1.5372 1.6474 1.7608

(Franz, J. pr. (2) 5. 274.)

Sp. gr. of $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ at 18° .

% $\text{Cd}(\text{NO}_3)_2$ 1 5 10 15
Sp. gr. 1.0069 1.0415 1.0869 1.136

% $\text{Cd}(\text{NO}_3)_2$ 20 25 30 35
Sp. gr. 1.1903 1.25 1.3125 1.3802

% $\text{Cd}(\text{NO}_3)_2$ 40 45 48
Sp. gr. 1.459 1.543 1.5978

(Grotrian, W. Ann. 1883, 18. 193.)

Sp. gr. of $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ at room temp. containing:

7.81 15.71 22.36% $\text{Cd}(\text{NO}_3)_2$.
1.0744 1.1593 1.2411

(Wagner, W. Ann. 1883, 18. 265.)

Sp. gr. of $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$.

$\text{Cd}(\text{NO}_3)_2$	t°	Sp. gr. at t°	Sp. gr. at 18°
0.0492	17.57	0.99912	0.99904
...	21.14	0.99839	...
0.100	18.00	...	0.99945
0.249	17.34	1.0008	1.0007
...	20.22	1.0002	...
0.464	18.00	...	1.0025
0.952	18.00	...	1.0065

(Wershofen, Z. phys. Ch. 1890, 5. 493.)

Sp. gr. of $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ at 25°

Concentration of $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$.	Sp. gr.
1-normal	1.095
$\frac{1}{2}$ - " "	1.047
$\frac{1}{5}$ - " "	1.024
$\frac{1}{16}$ - " "	1.011

(Wagner, Z. phys. Ch. 1890, 5.)

Sp. gr. of $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ at $18^\circ/4^\circ$

% $\text{Cd}(\text{NO}_3)_2$ 54.027 43.716
Sp. gr. 1.711 1.515

% $\text{Cd}(\text{NO}_3)_2$ 21.353 14.899
Sp. gr. 1.204 1.134

(de Muynck, W. Ann. 1894, 53.)

$\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ containing 7.89% has sp. gr. $20^\circ/20^\circ = 1.0673$.

$\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ containing $\text{Cd}(\text{NO}_3)_2$ has sp. gr. $20^\circ/20^\circ = 1.10$. (Le Blanc and Rohland, Z. phys. 19. 282.)

Sat. $\text{Cd}(\text{NO}_3)_2 + \text{Aq}$ boils at 132° . Almost entirely insol. in conc. H (Wurtz.)

Moderately sol. in liquid NH_3 . (Am. Ch. J. 1898, 20. 827.)

Sol. in alcohol.

Sol. in ethyl acetate. (Naumann 37. 3601.)

Sol. in acetone and in methylmann, C. C. 1899, II. 1014.)

+ $4\text{H}_2\text{O}$. M.-pt. of $\text{Cd}(\text{NO}_3)_2$ 59.5° . (Ordway; Tilden, Chem. Soc Solubility in H_2O .

Solubility in H_2O at t° .

t°	% $\text{Cd}(\text{NO}_3)_2$ in the solution	Mols 1 mol.
0	52.31	1
18	55.90	1
30	58.40	
40	61.42	
59.5 mpt.	76.54	

(Funk, B. 1899, 32. 105.)

Sat. solution of $\text{Cd}(\text{NO}_3)_2 + 4\text{H}_2\text{O}$ at 0° contains 52.3% $\text{Cd}(\text{NO}_3)_2$, 55.9% $\text{Cd}(\text{NO}_3)_2$. (Mylius, Z. an 74. 411.)

Sol. in liquid NH_3 . (Johnson more, Elektroch. Z. 1908, 14. 227.)

Sol. in acetone. (Naumann, B. 4328.)

Sol. in ethyl acetate. (Naumann 43. 314.)

+ $9\text{H}_2\text{O}$. Solubility in H_2O .

Sat. solution contains at:

-13° -1° $+1^\circ$
37.37 47.33 52.73% $\text{Cd}(\text{NO}_3)_2$

hydrate is formed at -16° . (Funk, *anorg.* 1899, **20**. 416.)

The composition of the hydrates formed by NO_3 , at different dilutions is calculated as determinations of the lowering of the σ . produced by $\text{Cd}(\text{NO}_3)_2$ and of the conductivity and sp. gr. of $\text{Cd}(\text{NO}_3)_2 + \text{Aq.}$ *anorg.* Am. Ch. J. 1905, **34**. 308.)

caesium uranyl nitrate, $\text{Cd}(\text{NO}_3)_2$,
 $(\text{UO}_2)(\text{NO}_3)_2 + 30\text{H}_2\text{O}$.

Sol. in H_2O and acids. Insol. in alcohol and alkalies + Aq. (Lancien, C. C. 1912, I. 1.)

caesium nitrate ammonia, $\text{Cd}(\text{NO}_3)_2$, 6NH_3 ,
 $+ \text{H}_2\text{O}$.
(André, C. R. 104. 987.)

caesium nitrate cupric oxide, $\text{Cd}(\text{NO}_3)_2$,
 $\text{CuO} + 5\text{H}_2\text{O}$.
Ppt. (Mailhe, C. R. 1902, **134**. 235.)

caesium nitrate cupric oxide, $\text{Cd}(\text{NO}_3)_2$,
 $3\text{CuO} + 5\text{H}_2\text{O}$.
(Mailhe, A. ch. 1902, (7) **27**. 383.)

caesium nitrate hydrazine, $\text{Cd}(\text{NO}_3)_2$,
 $3\text{N}_2\text{H}_4$.
Decomp. by hot H_2O . Sol. in warm H_2O . (Franzen, Z. anorg. 1908, **60**. 282.)

caesium nitrate, CsNO_3 .

100 pts. H_2O dissolve 10.58 pts. CsNO_3 at 0° . Sl. sol. in absolute alcohol. (Bunsen.)

Solubility of CsNO_3 in H_2O at t° .

t°	G. CsNO_3 per 100 g.		t°	G. CsNO_3 per 100 g.	
	Solu- tion	Water		Solu- tion	Water
0	8.54	9.33	60	45.6	83.8
10	12.97	14.9	70	51.7	107.0
20	18.7	23.0	80	57.3	134.0
30	25.3	33.9	90	62.0	163.0
40	32.1	47.2	100	66.3	197.0
50	39.2	64.4	106.2	68.8	220.3

(Berkeley, Trans. Roy. Soc. 1904, **203**.
A, 213.)

100 g. H_2O dissolve 26.945 g. CsNO_3 at 0° . (Haigh, J. Am. Chem. Soc. 1912, **34**. 18.)

Sp. gr. $20^{\circ}/4^{\circ}$ of a normal solution of CsNO_3 1.140905; of a 0.5 normal solution = 7001. (Haigh.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 14; Naumann, B. 1904, **37**. 4328.)

Solubility in glycol = 8% at ord. temp. (Coninck, Belg. Acad. Bull. 1905, 359.)

caesium hydrogen nitrate.

CsNO_3 , HNO_3 . Sol. in H_2O . (Wells, Am. Ch. J. 1901, **26**. 273.)

CsNO_3 , 2HNO_3 . (W.)

caesium cerium nitrate, $\text{Cs}_2\text{Ce}(\text{NO}_3)_6$.

Sol. in H_2O ; very sl. sol. in HNO_3 . (Meyer, Z. anorg. 1901, **27**. 371.)

Sol. in HNO_3 . (Meyer, B. 1900, **33**. 2137.)

caesium ferric nitrate, CsNO_3 , $\text{Fe}(\text{NO}_3)_3 + 7\text{H}_2\text{O}$.

Deliquescent. (Wells, Am. Ch. J. 1901, **26**. 276.)

caesium silver nitrate, CsNO_3 , AgNO_3 .

Sol. in H_2O . (Russell and Maskelyne, Roy. Soc. Proc. **26**. 357.)

caesium thorium nitrate, $\text{Cs}_2\text{Th}(\text{NO}_3)_6$.

Decomp. by H_2O ; sl. sol. in HNO_3 . (Meyer, Z. anorg. 1901, **27**. 384.)

caesium uranyl nitrate, $\text{Cs}(\text{UO}_2)(\text{NO}_3)_3$.

Decomp. by H_2O . Sol. in conc. HNO_3 . (Meyer, B. 1903, **36**. 4057.)

Decomp. by H_2O at low temp., so that the solid phase in contact with the solution consists of the double salt and CsNO_3 . At 16.1° 100 pts. by wt. of the solution in H_2O contain 31.39 pts. UO_2 and 6.59 pts. Cs . (Rimbach, B. 1904, **37**. 477.)

calcium nitrate, basic, $\text{Ca}(\text{NO}_3)_2$, $\text{CaO}_2\text{H}_2 + 2\frac{1}{2}\text{H}_2\text{O}$.

Decomp. by H_2O . (Werner, A. ch. (6) **27**. 570.)

+ H_2O . As above. (Rousseau and Tite, C. R. **114**. 1184.)

calcium nitrate, $\text{Ca}(\text{NO}_3)_2$.

Deliquescent. Very sol. in H_2O with evolution of much heat.

100 pts. H_2O at 0° dissolve 84.2 pts. $\text{Ca}(\text{NO}_3)_2$. (Poggiale.)

100 pts. H_2O at 0° dissolve 93.1 pts. $\text{Ca}(\text{NO}_3)_2$. (Mulder.)

Sol. in 0.25 pt. cold H_2O with reduction of temp. Sol. in all proportions in boiling H_2O . (Berzelius.)

Sol. in 2 pts. cold, and 0.6667 pt. boiling H_2O . (Fourcroy.)

Sat. $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 12.5° contains 33.8%. (Hassensfratz, A. ch. **28**. 29.)

Solubility in H_2O .

100 g. of the solution contain at:

55°	80°	90°	100°
78.16	78.20	78.37	78.43 g. $\text{Ca}(\text{NO}_3)_2$,

125°	147.5°	151° (bpt. of sat. solution at 760 mm.)
78.57	78.80	79.00 g. $\text{Ca}(\text{NO}_3)_2$.

The anhydrous salt is the stable solid phase

above 51.3° . (Bassett and Taylor, Chem. Soc. 1912, **101**. 580.)

100 g. sat. $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ contain 77.3 g. $\text{Ca}(\text{NO}_3)_2$ at 25°. (Taylor and Henderson, J. Am. Chem. Soc. 1915, 37. 1692.)

See also +2, 3, and 4H₂O.

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at room temp. containing:

17.55	30.10	40.13 % $\text{Ca}(\text{NO}_3)_2$.
1.1714	1.2739	1.3857

(Wagner, W. Ann. 1883, 18. 270.)

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 17.5°.

$\text{Ca}(\text{NO}_3)_2$	Sp. gr.	$\text{Ca}(\text{NO}_3)_2$	Sp. gr.
1	1.009	35	1.328
5	1.045	40	1.385
10	1.086	45	1.447
15	1.129	50	1.515
20	1.174	55	1.588
25	1.222	60	1.666
30	1.272

(Franz, J. pr. (2) 5. 274.)

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 17.5°.

$\text{Ca}(\text{NO}_3)_2$	Sp. gr.	$\text{Ca}(\text{NO}_3)_2$	Sp. gr.
10	1.076	40	1.368
20	1.163	50	1.483
30	1.261	60	1.605

(Gerlach, Z. anal. 27. 283.)

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 18°.

$\text{Ca}(\text{NO}_3)_2$	Sp. gr.	$\text{Ca}(\text{NO}_3)_2$	Sp. gr.
6.25	1.0487	37.5	1.3546
12.5	1.1016	50.0	1.5102
25.0	1.2198

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 24.65°. a=no. of g. $\times \frac{1}{2}$ mol. wt. dissolved in 1000 g. H₂O; b=sp. gr. if a is $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$, $\frac{1}{2}$ mol. wt.=118; c=sp. gr. if a is $\text{Ca}(\text{NO}_3)_2$, $\frac{1}{2}$ mol. wt.=82.

a	b	c	a	b	c
1	1.056	1.059	6	1.243	1.286
2	1.104	1.112	7	1.270	1.323
3	1.145	1.160	8	1.294	...
4	1.181	1.205	9	1.316	...
5	1.213	1.246	10	1.336	...

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 25°.

Concentration of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$.	Sp. gr.
1-normal	1.0596
$\frac{1}{2}$ "	1.0300
$\frac{1}{4}$ "	1.0151
$\frac{1}{8}$ "	1.0076

(Wagner, Z. phys. Ch. 1890, 5. 36.)

$\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ containing 7.15% Ca has sp. gr. 20°/20° = 1.0554.

$\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ containing 7.91% Ca has sp. gr. 20°/20° = 1.0613.

(Le Blanc and Rohland, Z. phys. Ch 19. 284.)

Sp. gr. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 20° cont M g. mols. of salt per liter.

M	0.0125	0.025	0.05	0
Sp. gr.	1.001846	1.003166	1.00604	1

M	0.25	0.5	0.75
Sp. gr.	1.03074	1.06011	1.08874

M	1.00	1.50
Sp. gr.	1.11751	1.17375

(Jones and Pearce, Am. Ch. J. 1907, 3)

Saturated $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ containing pts. $\text{Ca}(\text{NO}_3)_2$ to 100 pts. H₂O boils (Legrand); 152° (Kremers).

Forms a crust at 141°, and contain pts. $\text{Ca}(\text{NO}_3)_2$ to 100 pts. H₂O; highest observed, 151°. (Gerlach, Z. anal. 26.

B.-pt. of $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ containing $\text{Ca}(\text{NO}_3)_2$ to 100 pts. H₂O. G=ing to Gerlach (Z. anal. 26. 44' according to Legrand (A. ch. (2) 5

B.-pt.	G	L	B.-pt.	G
101°	10	15	127	215.5
102	20	25.3	128	222.5
103	30	34.4	129	230
104	40	42.6	130	237.5
105	50	50.4	131	245
106	60	57.8	132	253
107	70	64.9	133	261.5
108	80	71.8	134	270
109	89	78.6	135	278.5
110	98	85.3	136	287
111	106.5	91.9	137	296
112	114.5	98.4	138	305
113	122.5	104.8	139	314.5
114	130	111.2	140	324
115	137.5	117.5	141	333.5
116	144	123.8	142	343.5
117	150.5	130	143	354
118	157	136.1	144	364.5
119	163.5	142.1	145	375
120	170	148.1	146	386
121	176	...	147	397.5
122	182.5	160.1	148	409
123	189	...	149	420.5
124	195.5	172.2	150	432.5
125	202	...	151	444.5
126	208.5	184.5	151.97	455.6

Sat. $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ boils at 132°. way, Sill. Am. J. (2) 27. 14.)

Conc. HNO₃ precipitates $\text{Ca}(\text{NO}_3)_2$ its aqueous solution. (Mitscherlich 18. 159.)

Very sol. in conc. HNO₃. (Raw Soc. Chem. Ind. 1897, 16. 113.)

Solubility in HNO₃+Aq at 25°.

solution contain		Solid phase
g.	G. HNO ₃	
	0.00	Ca(NO ₃) ₂ ·4H ₂ O
	3.33	
	5.87	
	7.21	
	11.27	
	13.71	
	19.65	
	22.80	
	28.81	
	32.63	
	33.52	Ca(NO ₃) ₂ ·3H ₂ O
	35.63	
	41.66	
	45.70	
	40.56	
	45.70	Ca(NO ₃) ₂ ·2H ₂ O
	55.48	
	62.05	
	65.69	
	67.20	
	71.12	Ca(NO ₃) ₂
	74.77	
	78.56	
	80.83	
	85.83	
out)	90.90	
	96.86	

Results show that the hydrates of which are stable at 25° in contact with HNO₃+Aq are Ca(NO₃)₂·4H₂O, Ca(NO₃)₂·3H₂O, and Ca(NO₃)₂·2H₂O. (Taylor, Chem. Soc. 1912, 101. 582.)

Reaction with HC₂H₃O₂. (Persoz.) At 25° KNO₃+Aq with elevation of pptn. of a portion of KNO₃. (Vauquelin, A. ch. 11. 135.)

Solubility of Ca(NO₃)₂+NaNO₃ at t°.

g.	% NaNO ₃	Solid phase
	9.51	Ca(NO ₃) ₂ ·4H ₂ O " + NaNO ₃ NaNO ₃ "
	12.56	
	23.32	
	34.26	
	7.25	Ca(NO ₃) ₂ ·4H ₂ O " + NaNO ₃ NaNO ₃ " " "
	10.70	
	12.08	
	11.58	
	19.48	
	24.98	
	36.12	

and Rothmund, Z. anorg. 1914, 86. 373.)

Solubility of Ca(NO₃)₂+CaS₂O₃ at t°.

t°	% Ca(NO ₃) ₂	% CaS ₂ O ₃	Solid phase
9	46.02	5.46	Ca(NO ₃) ₂ ·4H ₂ O
	45.68	6.81	" CaS ₂ O ₃ ·6H ₂ O
	27.92	10.46	CaS ₂ O ₃ ·6H ₂ O
	10.49	22.81	"
	...	29.33	"
25	54.03	4.27	Ca(NO ₃) ₂ ·4H ₂ O
	50.25	9.10	"
	45.92	13.	" + CaS ₂ O ₃ ·6H ₂ O
	42.93	13.83	CaS ₂ O ₃ ·6H ₂ O
	32.01	17.09	"
	19.51	23.78	"
	8.15	29.85	"

(Kremann and Rothmund, Z. anorg. 1914, 86. 373.)

Very easily sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)
Sol. in 0.8 pt. alcohol (Macquer); 1 pt. boiling alcohol. (Bergmann.)
Dry Ca(NO₃)₂ is sol. in 7 pts. alcohol at 15° and 1 pt. boiling alcohol. (Bergmann.)

Sp. gr. of Ca(NO₃)₂+alcohol.

% Ca(NO ₃) ₂	Sp. gr. 20°/20°
0	0.7949
4.96	0.8278
6.47	0.8383

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 284.)

Solubility in ethyl alcohol+Aq at 25°.

% C ₂ H ₅ OH in the solvent	% C ₂ H ₅ OH in the solution	% Ca(NO ₃) ₂ in the solution
*0	0	82.5
*25.1	5.8	77.0
*50.1	15.2	69.52
*60.1	20.4	66.08
*63.9	22.4	64.94
70.4	26.5	62.3
72.0	27.39	61.96
73.4	28.5	61.15
75.3	29.9	60.3
*84.9	35.9	57.7
*99.1	48.1	51.4

* Metastable solutions.
(D'Ans and Siegler, Z. phys. Ch. 1913, 82. 43.)

Solubility of $\text{Ca}(\text{NO}_3)_2 \cdot 2\text{C}_2\text{H}_5\text{OH}$ in $\text{C}_2\text{H}_5\text{OH} + \text{Aq}$ at 25° .

% $\text{C}_2\text{H}_5\text{OH}$ in the solvent	% $\text{C}_2\text{H}_5\text{OH}$ in the solution	% $\text{Ca}(\text{NO}_3)_2$ in the solution
98.1	60.2	38.6
94.1.	54.6	41.9
85.8	42.5	50.97
80.5	35.8	55.3
75.3	29.9	60.28

(D'Ans and Siegler, *l. c.*)

See also under $+4\text{H}_2\text{O}$.

Solubility in organic solvents.

Solvent	% $\text{Ca}(\text{NO}_3)_2$ in the solution at 25°
Methyl alcohol	65.5
Ethyl alcohol	52.0
Propyl alcohol	36.5
Isobutyl alcohol	25.0
Amyl alcohol	13.3
Acetone	58.5

(D'Ans and Siegler, *l. c.*)

Solubility in H_2O at t° .
100 g. of the solution contain g. Ca at t° .

t°	G. $\text{Ca}(\text{NO}_3)_2$
-26.7	43.37
-10.0	47.31
0.0	50.50
$+5.0$	51.97
10.0	53.55
15.0	54.94
20.0	56.39
25.0	57.98
30.0	60.41
35.0	62.88
40.0	66.21
42.4	68.68
42.5	68.74
42.7	mpt. of $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
42.45	71.70

(Bassett and Taylor, *l. c.*)

Solubility of α and β modifications in
at t° .

α modification is the stable form.

t°	G. $\text{Ca}(\text{NO}_3)_2$ in 100 g. of solution	Solid phase
0	50.17	α $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
22.2	56.88	α "
25.0	57.90	α "
30.0	60.16	α "
30.0	61.57	β "
34.0	63.66	β "
35.0	62.88	α "
38.0	64.34	α "
38.0	66.65	β "
39.0	67.93	β "
¹ 39.6	69.50	β "
² 39.0	75.34	β "
40.0	66.21	α "
¹ 42.7	69.50	α "
² 42.4	71.70	α "

¹ mpt. of hydrate.
² reflex pt.

(Taylor and Henderson, *J. Am. Chem. Soc.* 1915, **37**, 1692.)

Sp. gr. of solution sat. at $18^\circ = 1.54$ containing 54.8% $\text{Ca}(\text{NO}_3)_2$. (Mylus, *B.* **30**, 1718.)

Sol. in 1.87 pts. ether-alcohol (1 : 1). (Fresenius, *Z. anal.* **32**, 191.)

Ether ppts. $\text{Ca}(\text{NO}_3)_2$ from its alcoholic solution. Easily sol. in boiling amyl alcohol. (Browning, *Sill. Am. J.* **143**, 53.)

Sol. in acetone. (Naumann, *B.* 1904, **37**, 4328.)

Insol. in benzonitrile. (Naumann, *B.* 1914, **47**, 1370.)

Insol. in methylal. (Eidmann, *C. C.* **1899**, II, 1014.)

1 g. $\text{Ca}(\text{NO}_3)_2$ is sol. in 1.44 g. methyl acetate at 18° . Sp. gr. $18^\circ/4^\circ$ of sat. solution = 1.313. (Naumann, *B.* 1909, **42**, 3795.)

Sol. in ethyl acetate. (Naumann, *B.* 1910, **43**, 314.)

$+2\text{H}_2\text{O}$. Solubility in H_2O .
100 g. of the solution contain at:
 49° 51°
77.49 78.05 g. $\text{Ca}(\text{NO}_3)_2$.

Solutions in stable equilibrium with the dihydrate can only exist between the limits of temp. 48.4° and 51.3° . (Bassett and Taylor, *Chem. Soc.* 1912, **101**, 580.)

$+3\text{H}_2\text{O}$. Solubility in H_2O .
100 g. of the solution contain at:
 40° 45° 50° 51°
70.37 71.45 73.79 74.73 g. $\text{Ca}(\text{NO}_3)_2$.
Mpt. of $\text{Ca}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O} = 51.1^\circ$.
(Bassett and Taylor, *l. c.*)

$+4\text{H}_2\text{O}$. $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ melts in its crystal H_2O at 44° . (Tilden, *Chem. Soc.* **45**, 409.)

bility in ethyl alcohol + Aq at 25°.

OH in solvent	% C ₂ H ₅ OH in the solution	% Ca(NO ₃) ₂ in the solution
	0	57.5
.3	3.5	56.1
.2	8.1	55.2
.2	14.1	52.9
.4	22.3	50.2
.4	29.4	49.0
.4	31.1	49.7
.4	31.2	52.0
.4	29.5	56.2
.1	28.3	58.9
.1	27.8	60.0
.1	27.3	60.7
.1	26.5	62.3

and Siegler, Z. phys. Ch. 1913, **82**. 42.)

nitrate hydrazine, Ca(NO₃)₂, 2N₂H₄,
I₂O.

(Fransen, Z. anorg. 1908, **60**. 288.)

nitrate tungstosilicate, Ca₂W₁₂SiO₄₀,
(NO₃)₂.

I₂O and +15H₂O. Decomp. by H₂O.
boff, Chem. Soc. 1897, **72**. (2) 176.)

nitrate, Ce(NO₃)₃ + 6H₂O.

very deliquescent. (Jolin.)

sol. in H₂O; sol. in 2 pts. alcohol.
lin.)

in acetone. (Eidmann, C. C. 1899. II.
aumann, B. 1904, **37**. 4328.)

trate, Ce(NO₃)₄.

uescent. Decomp. by hot H₂O.
us.)

in alcohol. (Dumas.)

compounds containing 12 mols. or
), to 1 mol. N₂O₅ may be obtained,
re sol. in H₂O. (Ordway.)

cobaltous nitrate, 2Ce(NO₃)₃,
(NO₃)₂ + 24H₂O.

uescent. Easily forms supersaturated
s. (Lange, J. pr. **82**. 129.)

sat. solution in HNO₃ + Aq (sp. gr.
contains 103.3 g. hydrous salt at
antsch, Z. anorg. 1912, **76**. 321.)

baltous nitrate, CeCo(NO₃)₆ + 8H₂O.

np. by H₂O when heated; sol. in cold
sol. in HNO₃. (Meyer, Z. anorg.
. 376.)

magnesium nitrate, 2Ce(NO₃)₃,
5(NO₃)₂ + 24H₂O.

ly deliquescent. Easily sol. in H₂O
ol, and easily forms supersaturated
s. (Holzmann, J. pr. **75**. 330.)

sat. solution in HNO₃ + Aq (sp. gr.
contains 63.8 g. hydrous salt at 16°.
, Z. anorg. 1912, **76**. 321.)

Ceric magnesium nitrate, CeMg(NO₃)₆.
+ 8H₂O.

Decomp. by H₂O; sol. in HNO₃ + Aq with-
out decomp. (Meyer, Z. anorg. 1901, **27**.
373.)

Cerous manganous nitrate, 2Ce(NO₃)₃,
3Mn(NO₃)₂ + 24H₂O.

Sol. in H₂O. (Lange, J. pr. **82**. 129.)

1 l. sat. solution in HNO₃ + Aq (sp. gr.
1.325) contains 193.1 g. hydrous salt at 16°.
(Jantsch.)

Ceric manganous nitrate, CeMn(NO₃)₆ +
8H₂O.

Decomp. by H₂O and dil. HNO₃; sol. in
conc. HNO₃ without decomp. (Meyer, Z.
anorg. 1901, **27**. 377.)

Cerous nickel nitrate, 2Ce(NO₃)₃, 3Ni(NO₃)₂,
+ 24H₂O.

Easily sol. in H₂O. (Holzmann, J. pr. **75**.
321.)

1 l. sat. solution in HNO₃ + Aq (sp. gr.
1.325) contains 80.3 g. hydrous salt at 16°.
(Jantsch.)

Ceric nickel nitrate, CeNi(NO₃)₆ + 8H₂O.

Decomp. by H₂O when heated; sol. in H₂O
in the cold; sl. sol. in HNO₃. (Meyer, Z.
anorg. 1901, **27**. 375.)

Cerous potassium nitrate, Ce(NO₃)₃, 2KNO₃,
+ 2H₂O.

Sol. in H₂O. (Lange, J. pr. **82**. 136.)

Ceric potassium nitrate, CeK₂(NO₃)₆.

Sol. in H₂O with decomp. (Meyer, Z.
anorg. 1901, **27**. 370.)

+ 1½H₂O. Efflorescent. (Holzmann, J.
pr. **75**. 324.)

Ceric rubidium nitrate, CeRu₂(NO₃)₆.

Very sol. in H₂O; sl. sol. in HNO₃. (Meyer.)

Ceric sodium nitrate.

Deliquescent. Decomp. by recrystalliza-
tion. (Holzmann.)

Cerous thallous nitrate, [Ce(NO₃)₃]Tl₂ +
4H₂O.

Very hygroscopic. Decomp. by H₂O.
(Jantsch, Z. anorg. 1911, **69**. 229.)

Cerous zinc nitrate, 2Ce(NO₃)₃, 3Zn(NO₃)₂ +
24H₂O.

Sol. in H₂O. Easily forms supersat. solu-
tions. (Lange, J. pr. **82**. 129.)

1 l. sat. solution in HNO₃ + Aq (sp. gr.
1.325) contains 124.1 g. hydrous salt at 16°.
(Jantsch, Z. anorg. 1912, **76**. 321.)

Ceric zinc nitrate, ZnCe(NO₃)₆ + 8H₂O.

Decomp. by H₂O; sol. in HNO₃ + Aq.
(Meyer, Z. anorg. 1901, **27**. 374.)

Ceroceric zinc nitrate (?), $\text{Ce}_2\text{O}_3 \cdot 2\text{ZnO}$, $6\text{N}_2\text{O}_5 + 18\text{H}_2\text{O}$ (?).

Easily sol. in H_2O . (Holzmann, J. pr. 75. 321.)

Chromic nitrate, basic, $\text{Cr}_2\text{O}(\text{NO}_3)_4$.

Sol. in H_2O . (Löwel.)

+12 H_2O . Sol. in H_2O . (Ordway.)

Chromic nitrate, $\text{Cr}(\text{NO}_3)_3 + 9\text{H}_2\text{O}$.

Very sol. in H_2O and alcohol. (Löwel.)

Melts in its crystal H_2O at 36.5° . Sat. $\text{Cr}(\text{NO}_3)_3 + \text{Aq}$ boils at 125.6° . (Ordway.)

Sp. gr. of $\text{Cr}(\text{NO}_3)_3 + \text{Aq}$.

M = concentration of solution in gram. mols.

W = wt. of 25 cc. of solution.

M 0.0934 0.1868 0.3736 0.5604 0.9340
W 25.4300 25.8828 26.7302 27.5524 29.3072

M 1.1208 1.3076 1.4944 1.8680

W 30.0668 30.8464 31.6327 33.3379

(Jones and Getman, Z. phys. Ch. 1904, 49. 426.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328; Eidmann, C. C. 1899, II. 1014.)

Chromic nitrate chloride, $\text{CrCl}_2(\text{NO}_3)_2$.

Sol. in H_2O and alcohol. (Schiff, A. 124. 177.)

$\text{Cr}(\text{NO}_3)_2\text{Cl}$. (Schiff.)

Chromic nitrate sulphate, $\text{Cr}_2(\text{SO}_4)(\text{NO}_3)_4$.

Hygroscopic. Completely sol. in H_2O .

$\text{Cr}_2(\text{SO}_4)_2(\text{NO}_3)_2$. Sol. in H_2O . (Schiff, A. 124. 174.)

Cobaltous nitrate, basic, 6CoO , $\text{N}_2\text{O}_5 + 5\text{H}_2\text{O}$.

Ppt. Gradually sol. in H_2O with deposition of CoO . (Winkelblech, A. 13. 155.)

Sol. in cold HCl , and $\text{HNO}_3 + \text{Aq}$. De-comp. by hot $\text{KOH} + \text{Aq}$.

4CoO , $\text{N}_2\text{O}_5 + 6\text{H}_2\text{O}$. Ppt. (Habermann, M. 5. 432.)

Cobaltous nitrate, $\text{Co}(\text{NO}_3)_2$.

Deliquescent in moist air. Very sol. in H_2O .

See +3, 6, and 9 H_2O .

Sp. gr. of aqueous solution at 17.5° containing:

5 10 15 20% $\text{Co}(\text{NO}_3)_2$,
1.0462 1.0906 1.1378 1.1936

25 30 35 40% $\text{Co}(\text{NO}_3)_2$,
1.2538 1.3190 1.3896 1.4662

Sp. gr. of sat. solution = 1.5382.

(Franz, J. pr. (2) 5. 274.)

Sp. gr. of $\text{Co}(\text{NO}_3)_2 + \text{Aq}$ at room temp. containing:

8.28 15.96 24.528% $\text{Co}(\text{NO}_3)_2$.

1.0732 1.1436 1.2288

(Wagner, W. Ann. 1883, 18. 268.)

Sp. gr. of $\text{Co}(\text{NO}_3)_2 + \text{Aq}$ at 20°

Concentration of $\text{Co}(\text{NO}_3)_2 + \text{Aq}$.	Sp. gr.
1-normal	1.075
$\frac{1}{2}$ " "	1.034
$\frac{1}{4}$ " "	1.015
$\frac{1}{8}$ " "	1.005

(Wagner, Z. phys. Ch. 1890, 5.

Sp. gr. at 20° of $\text{Co}(\text{NO}_3)_2 + \text{Aq}$ containing
M g. mols. $\text{Co}(\text{NO}_3)_2$ per liter.

M 0.01 0.025 0.05
Sp. gr. 1.001496 1.003863 1.007579

M 0.10 0.25 0.5
Sp. gr. 1.015084 1.03737 1.07415

M 1.0 1.5 2.0
Sp. gr. 1.14612 1.21720 1.28576

(Jones and Pearce, Am. Ch. J. 1907

Sol. in liquid NH_3 . (Guntz, 1909 (4) 5. 1009.)

100 g. sat. solution in glycol contains
 $\text{Co}(\text{NO}_3)_2$. (de Coninck, C. C. 1904

Sol. in ethyl acetate. (Naumann 37. 3601.)

+3 H_2O . Solubility in H_2O .

Sat. solution contains at:

55° 62° 70° 84° 91° mpt
61.74 62.88 64.89 68.84 77.21% (

(Funk, Z. anorg. 1899, 20. 40

+6 H_2O . Melts in its crystal H_2O (Ordway); 38° (Tilden).

Solubility in H_2O .

Sat. solution contains at:

—21° —10° —4° 0°
41.55 43.69 44.85 45.66% Co

+18° 41° 56° mpt.

49.73 55.96 62.88% $\text{Co}(\text{NO}_3)_2$.

(Funk, Z. anorg. 1899, 20. 40

Moderately sol. in liquid NH_3 .

Am. Ch. J. 1898, 20. 827.)

Easily sol. in alcohol. Sol. in 1 alcohol at 12.5° . (Wenzel.)

Easily sol. in acetone. (Krug and J. Anal. Ch. 6. 184.)

Sol. in methyl acetate. (Nau 1909, 42. 3790.)

Difficultly sol. in ethyl acetate (Naumann, B. 1910, 43. 314.)

+9 H_2O . Solubility in H_2O .

Sat. solution contains at:

—26° —23.5° —20.5°
39.45 40.40 42.77% Co

Cryohydrate is formed at -29° (Z. anorg. 1899, 20. 409.)

Sp. gr. at 20° of $\text{Cu}(\text{NO}_3)_2 + \text{Aq}$ containing
M g. mols. salt per liter.

M	0.01	0.025	0.05	0.075
Sp. gr.	1.001504	1.004076	1.007859	1.011715

M	0.25	0.50	0.75	0.935
Sp. gr.	1.040290	1.07723	1.11469	1.14262

Sp. gr.	1.5	2.0
M	1.22618	1.29262

(Jones and Pearce, Am. Ch. J. 1917, **38**, 719.)

Sat. $\text{Cu}(\text{NO}_3)_2 + \text{Aq}$ boils at about 173°.
(Griffiths.)

Insol. in fuming HNO_3 . (Ditte, A. ch. 1879 (5) **18**, 339.)

Solubility of $\text{Cu}(\text{NO}_3)_2 + \text{Pb}(\text{NO}_3)_2$ in H_2O
at 20°.

Sp. gr.	In 1 l. of solution				Solid phase
	Cu(NO ₃) ₂		Pb(NO ₃) ₂		
	g.	g. mol.	g.	g. mol.	
1.354	70.5	0.375	359.5	1.086	Pb(NO ₃) ₂
1.322	139.2	0.742	257.2	0.777	"
1.321	226.5	1.207	175.1	0.529	"
1.343	301.8	1.608	133.4	0.403	"
1.360	341.8	1.821	117.8	0.356	"
1.451	519.4	2.767	70.5	0.213	"
1.546	681.7	3.632	44.0	0.133	"
1.622	798.1	4.252	28.1	0.085	"
1.700	943.2	5.028	17.2	0.052	Pb(NO ₃) ₂ + Cu(NO ₃) ₂ ·6H ₂ O

(Fedotieff, Z. anorg. 1911, **73**, 178.)

Very sol. in liquid NH_3 . (Guntz, Bull. Soc. 1909, (4) **5**, 1007.)

Easily sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**, 827.)

Insol. in liquid HF . (Franklin, Z. anorg. 1905, **46**, 2.)

Insol. in ethyl acetate. (Naumann, B. 1910, **43**, 314.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, **47**, 1369.)

+ $3\text{H}_2\text{O}$. Melts in crystal H_2O at 114.5°.
(Ordway; Tilden, Chem. Soc. **45**, 409.)

Solubility in H_2O .

Sat. solution contains at:

25°	30°	40°	50°
60.01	60.44	61.51	62.62% $\text{Cu}(\text{NO}_3)_2$

60°	70°	80°	114.5° Mpt.
64.17	65.79	67.51	77.59% $\text{Cu}(\text{NO}_3)_2$

(Funk, Z. anorg. 1899, **20**, 413.)

100 pts. HNO_3 dissolve 2 pts. at 13°, considerably more on heating. (Ditte, A. ch. 1879, (5) **18**, 339.)

Sol. in 1 pt. strong alcohol at 12.5°. (Wenzel.)

Insol. in methyl acetate. (Naumann, B. 1909, **42**, 3790.)

+ $6\text{H}_2\text{O}$. Efflorescent. Melts in crystal H_2O at 35°. (Ordway.)

Solubility in H_2O .

Sat. solution contains at:

-21°	-10°	0°	+10°
39.52	42.08	45.00	48.79% $\text{Cu}(\text{NO}_3)_2$

18°	20°	26.4° mpt.
53.86	55.58	63.39% $\text{Cu}(\text{NO}_3)_2$

(Funk, Z. anorg. 1899, **20**, 413.)

Sat. solution of $\text{Cu}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$ in H_2O at 20° contains 5.04 g. mol. per l. Sp. gr. of sat. solution = 1.688. (Fedotieff, Z. anorg. 1911, **73**, 78.)

Sat. solution of $\text{Cu}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$ in H_2O contains 45.0 g. $\text{Cu}(\text{NO}_3)_2$ in 100 g. solution at 0°; 53.9 g. at 18°. (Mylus, Z. anorg. 1912, **74**, 411.)

+ $9\text{H}_2\text{O}$.

Solubility in H_2O .

Sat. solution contains at:

-23°	-21°	-20°
36.08	37.38	40.92% $\text{Cu}(\text{NO}_3)_2$

Cryohydrate is formed at -24°. (Funk, Z. anorg. 1899, **20**, 414.)

Cupric nitrate ammonia (Cuprammonium nitrate), $\text{Cu}(\text{NO}_3)_2 \cdot 4\text{NH}_3$.

Easily sol. in H_2O , from which it can be recrystallized. Sol. in alcohol. (Berselius.)

Sol. in 1 pt. liquid NH_3 . (Horn, Am. Ch. J. 1908, **39**, 216.)

$\text{Cu}(\text{NO}_3)_2 \cdot 5\text{NH}_3$. (Horn, Am. Ch. J. 1907, **37**, 620.)

$4\text{Cu}(\text{NO}_3)_2 \cdot 23\text{NH}_3$. (Horn.)

Cupric nitrate hydrazine, $\text{Cu}(\text{NO}_3)_2 \cdot \text{N}_2\text{H}_4$.

Decomp. by H_2O . (Hofmann and Marburg, A. 1899, **305**, 221.)

Cupric nitrate mercuric oxide, $\text{Cu}(\text{NO}_3)_2 \cdot \text{HgO} + 3\text{H}_2\text{O}$.

Sol. in HCl , HNO_3 and H_2SO_4 . (Fina, Gazz. ch. it. 1913, **43**, (2) 709.)

Didymium nitrate, basic, $4\text{Di}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5 + 15\text{H}_2\text{O}$.

Insol. in H_2O . (Marignac.)

$2\text{Di}_2\text{O}_3 \cdot 3\text{N}_2\text{O}_5$. (Becquerel, A. ch. (6) **14**, 257.)

Didymium nitrate, $\text{Di}(\text{NO}_3)_2$.

Anhydrous. Very sol. in H_2O . As sol. is 96% alcohol as in H_2O , and the solution is not precipitated by much ether. Insol. in pure ether. (Marignac, A. ch. (3) **36**, 161.)

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**, 827.)

Sol. in acetone. (Naumann, B. 1904, **37**, 4328; Eidmann, C. C. 1899, II, 1014.)

+ $6\text{H}_2\text{O}$. Very deliquescent. (Cleve, Bull. Soc. (2) **43**, 361.)

Neodymium nickel nitrate, $2\text{Di}(\text{NO}_3)_2, 3\text{Ni}(\text{NO}_3)_2 + 36\text{H}_2\text{O}$.

Very deliquescent. (Frerichs and Smith, 191. 355.)

See Neodymium and praseodymium.

Neodymium zinc nitrate, $2\text{Di}(\text{NO}_3)_2, 3\text{Zn}(\text{NO}_3)_2 + 69\text{H}_2\text{O}$.

Very deliquescent. (F. and S.)

See Neodymium and praseodymium.

Neodymium nitrate, $\text{Dy}(\text{NO}_3)_3 + 5\text{H}_2\text{O}$.

Very sol. in H_2O ; less sol. in $\text{H}_2\text{O} + \text{HNO}_3$. Insol. in alcohol. (Urbain, C. R. 1908, 146. 1909.)

Neodymium nitrate, basic, $2\text{Er}_2\text{O}_3, 3\text{N}_2\text{O}_5 + 9\text{H}_2\text{O}$.

Decomp. by H_2O . Sl. sol. in HNO_3 . (Bahr and Bunsen.)

$3\text{Er}_2\text{O}_3, 4\text{N}_2\text{O}_5 + 20\text{H}_2\text{O}$. (Cleve, Bull. Soc. (2) 21. 344.)

Neodymium nitrate, $\text{Er}(\text{NO}_3)_3 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O , alcohol, and ether. (Höglund.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Gadolinium nitrate, $\text{Gd}(\text{NO}_3)_3 + 6\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . (Benedicks, Z. anorg. 1900, 22. 406.)

+ $5\text{H}_2\text{O}$. Sol. in HNO_3 . (B.)

Gadolinium magnesium nitrate, $2\text{Gd}(\text{NO}_3)_3, 3\text{Mg}(\text{NO}_3)_2 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 352.3 g. hydrous salt at 16° . (Jantsch, Z. anorg. 1912, 76. 303.)

Gadolinium nickel nitrate, $2\text{Gd}(\text{NO}_3)_3, 3\text{Ni}(\text{NO}_3)_2 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 400.8 g. hydrous salt at 16° . (Jantsch.)

Gadolinium zinc nitrate, $2\text{Gd}(\text{NO}_3)_3, 3\text{Zn}(\text{NO}_3)_2 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 472.7 g. hydrous salt at 16° . (Jantsch.)

Gallium nitrate, $\text{Ga}(\text{NO}_3)_3$.

Very deliquescent, and sol. in H_2O . (Dupré.)

Glucinum nitrate, basic, $2\text{GfO}, \text{N}_2\text{O}_5 + 3\text{H}_2\text{O}$ (?).

Sol. in H_2O .

$3\text{GfO}, \text{N}_2\text{O}_5$. Sol. in H_2O . (Ordway, Sill. Am. J. (2) 26. 205.)

Compounds more basic than this are insol. in H_2O . (Ordway.)

Glucinum nitrate, $\text{Gl}(\text{NO}_3)_2 + 3\text{H}_2\text{O}$.

Very deliquescent. (Joy, Sill. Am. J. (2) 36. 90.)

Easily sol. in H_2O and alcohol. (Vauquelin.)

Melts in its crystal H_2O at 29.4° . (Ordway.)

Sat. $\text{Gl}(\text{NO}_3)_2 + \text{Aq}$ boils at 140.5° . (Ordway.)

Gold (auric) nitrate, basic, $\text{Au}_2\text{O}_3, \text{N}_2\text{O}_5 + \frac{2}{3}\text{H}_2\text{O}$, or **Auryl nitrate**, $(\text{AuO})\text{NO}_3 + \frac{1}{3}\text{H}_2\text{O}$.

(Schottländer, A. 217. 364.)

$2\text{Au}_2\text{O}_3, \text{N}_2\text{O}_5 + 2\text{H}_2\text{O} = \text{Au}_4\text{O}_6(\text{NO}_3)_2 + 2\text{H}_2\text{O}$. Slowly sol. in $\text{HNO}_3 + \text{Aq}$ at 100° . (Schottländer, A. 217. 356.)

Gold (auric) nitrate, $\text{Au}(\text{NO}_3)_3 + x\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in acetone. (Hanriot and Raoult, C. R. 1912, 155. 1086.)

Gold (auric) hydrogen nitrate, $\text{Au}(\text{NO}_3)_3, \text{HNO}_3 + 3\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$. (Schottländer, A. 217. 356.)

Gold (auric) potassium nitrate, $\text{KAu}(\text{NO}_3)_4$.

Easily sol. in H_2O .

$\text{HK}_2\text{Au}(\text{NO}_3)_6$. Decomp. immediately by H_2O .

$2\text{KAu}(\text{NO}_3)_4, \text{K}_2\text{HAu}(\text{NO}_3)_6$. (Schottländer, J. B. 1884. 453.)

Gold (auric) rubidium nitrate, $\text{RbAu}(\text{NO}_3)_4$.

Easily sol. in H_2O .

$\text{HRb}_2\text{Au}(\text{NO}_3)_6$. As above. (Schottländer.)

Gold (auric) thallium nitrate, $\text{TlAu}(\text{NO}_3)_4$.

Easily sol. in H_2O .

$6\text{Au}_2\text{O}_3, 2\text{Tl}_2\text{O}_3, 3\text{N}_2\text{O}_5 + 15\text{H}_2\text{O}$. Ppt. (Schottländer.)

Indium nitrate, $\text{In}(\text{NO}_3)_3 + 4\frac{1}{2}\text{H}_2\text{O}$.

Very deliquescent. Easily sol. in H_2O and absolute alcohol. (Winkler.)

+ $1\frac{1}{2}\text{H}_2\text{O}$.

Iron (ferrous) nitrate, $\text{Fe}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$.

100 pts. of crystals dissolve in 50 pts. H_2O at 0° , sp. gr. of solution = 1.44; 40.8 pts. H_2O at 15° , sp. gr. of solution = 1.48; 33.3 pts. H_2O at 25° , sp. gr. of solution = 1.50. (Ordway, Sill. Am. J. (2) 40. 325.)

Sat. solution contains at:

— 9° 0° $+18^\circ$ 24° 60.5° Mpt.
39.68 41.53 45.14 46.51 62.50% $\text{Fe}(\text{NO}_3)_2$.

(Funk, Z. anorg. 1899, 20. 406.)

Sat. solution of $\text{Fe}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$ in H_2O contains 41.5% $\text{Fe}(\text{NO}_3)_2$ at 0° ; 45.1% at 18° . (Mylius, Z. anorg. 1912, 74. 411.)

+9H₂O. Solubility in H₂O.
Sat. solution contains at:
—27° —21.5° —19° —15.5°
35.66 36.10 36.56 37.17% Fe(NO₃)₃.

Cryohydrate is formed at —28°. (Funk, Z. anorg. 1899, 20, 407.)

Fe(NO₃)₃+Aq decomposes on heating; less rapidly when dil., more readily in presence of excess of acid. (Ordway.)

Iron (ferric) nitrate, basic, 36Fe₂O₃, N₂O₅+48H₂O (?)

Easily sol. in H₂O. Sl. sol. in dil. HNO₃+Aq; very sl. sol. in alcohol. (Hausmann, A. 89, 111.)

8Fe₂O₃, N₂O₅+12H₂O. Sl. sol. in H₂O; very sl. sol. in cold or warm dil. HNO₃+Aq, more easily sol. in hot HCl+Aq. (Hausmann.)

+xH₂O. Sol. in H₂O; completely pptd. from aqueous solution by NaCl, NH₄Cl, KI, KClO₃, Na₂SO₄, CaSO₄, ZnSO₄, CuSO₄, KNO₃, NaNO₃, Ba(C₂H₃O₂)₂, or Zn(C₂H₃O₂)₂+Aq. More slowly pptd. by NH₄NO₃, Mg(NO₃)₂, Ba(NO₃)₂, or Pb(NO₃)₂+Aq. Not pptd. by alcohol, Pb(C₂H₃O₂)₂, Cu(C₂H₃O₂)₂, Hg(CN)₂, AgNO₃, or As₂O₃+Aq. (Ordway, Sill. Am. J. (2) 9, 30.)

4Fe₂O₃, N₂O₅+1½H₂O. Easily sol. in H₂O; sl. sol. in dil. HNO₃+Aq, and in alcohol. (Hausmann.)

+3H₂O. Insol. in H₂O or HNO₃+Aq; sol. in HCl+Aq. (Scheurer-Kestner, C. R. 87, 927.)

+9H₂O. Not deliquescent; easily sol. in H₂O. (Ordway.)

3Fe₂O₃, N₂O₅+2H₂O. Insol. in H₂O. (Scheurer-Kestner.)

2Fe₂O₃, N₂O₅+H₂O. Decomp. by H₂O. (Scheurer-Kestner.)

+8H₂O. (S.-K.)

Fe₂O₃, N₂O₅. Decomp. by H₂O. (S.-K.)

Fe₂O₃, 2N₂O₅. Sol. in H₂O or alcohol in all proportions. Insol. in HNO₃+Aq.

N₂O₅ with 1, 2, 3, 4, 5, 6, and 8Fe₂O₃ forms compounds, sol. in H₂O. (Ordway.)

Solubility determinations show that there are no definite basic nitrates of iron formed from solutions at 25°, and that the solid phase under these conditions is a solid solution of Fe₂O₃, HNO₃ and H₂O. The normal salt, Fe₂O₃, 3N₂O₅, 18H₂O is stable in solutions containing about 30–45% N₂O₅. In higher concentrations of nitric acid it appears to be metastable and a new salt, Fe₂O₃, 4N₂O₅, 18(?)H₂O is the stable form. (Cameron, J. phys. Chem. 1909, 13, 252.)

Iron (ferric) nitrate, Fe(NO₃)₃.

+H₂O. (Scheurer-Kestner, A. ch. (3) 65, 113.)

+6H₂O. Deliquescent, and sol. in any amount of H₂O. (Schönbein, Pogg. 39, 141.)

Sol. in acetone. (Naumann, B. 1904, 37, 4328.)

+9H₂O. Deliquescent. Sol. in alcohol. Sl. sol. in HNO₃+Aq. 2 with 1 pt. H₂O lower the temperat (Scheurer-Kestner.)

Sp. gr. of solution at 17.5° conta
5 10 15 20 25%
1.0398 1.0770 1.1182 1.1612 1.2110

30 35 40 45 50%
1.2622 1.3164 1.3746 1.4338 1.4972
55 60 65% Fe(NO₃)₃
1.5722 1.6572 1.7532

(Frans, J. pr. (2) 5, 274.)

Nearly insol. in conc. HNO₃+A below 15.5°.

Easily sol. in alcohol.
Melts in crystal H₂O at 47.2°.

Sat. Fe(NO₃)₃+Aq boils at 12 way.)

Lanthanum nitrate, La(NO₃)₃+6H₂O

Very deliquescent; easily sol. in alcohol. (Mosander.) Melts in H₂O at 40°; boils at 124.5°.

La(NO₃)₃+Aq sat. at 25° conta La(NO₃)₃, or 100 g. H₂O dissolv La(NO₃)₃ at 25°. (James and W J. Am. Chem. Soc. 1912, 34, 1169.)

Sol. in acetone. (Naumann, B 4328; Eidmann, C. C. 1899, II, 10)

Lanthanum magnesium nitrate, 2 3Mg(NO₃)₂+24H₂O.

Deliquescent in moist air. (Ho pr. 76, 350.)

1 l sat. solution in HNO₃+A 1.325) contains 63.8 g. hydrous (Jantsch, Z. anorg. 1912, 76, 321.)

Lanthanum manganous nitrate, 2 3Mn(NO₃)₂+24H₂O.

Sol. in H₂O. (Damour and Dev 1 l. sat. solution in HNO₃+A 1.325) contains 193.1 g. hydrous (Jantsch)

Lanthanum nickel nitrate, 2 3Ni(NO₃)₂+36H₂O.

Very sol. in H₂O. (Frerichs and 191, 355.)

+24H₂O. 1 l sat. solution in Aq (sp. gr. 1.325) contains 80.3 salt at 16°. (Jantsch.)

Lanthanum rubidium hydrogen [La(NO₃)₃]Rb, HNO₃+6H₂O.

Sol. in H₂O and HNO₃. (Jantsch 1911, 69, 225.)

Lanthanum thallous nitrate, [La(N 4H₂O).

Hydroscopic. (Jantsch, Z. anorg 228.)

n zinc nitrate, $2\text{La}(\text{NO}_3)_3$, $\text{NO}_3)_2+24\text{H}_2\text{O}$.
l. in H_2O . (Damour and Deville, J. 35.)
t. solution in HNO_3+Aq (sp. gr. contains 124.1 g. hydrous salt at 16° . Z. anorg. 1912, 76. 321.)
O. (Frerichs and Smith, A. 191.

rate, basic, 2PbO , $\text{N}_2\text{O}_5+\text{H}_2\text{O}=(\text{H})\text{NO}_3$.
5.15 pts. H_2O at 19.2° . (Pohl, W. A.) Very sl. sol. in cold, much more). (Berzelius.) Sol. in $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2$ Guignet, C. R. 56. 358.)
n H_2O ; sol. in acids. (Athanesco, 1895, (3) 13. 178.)
) . (André, C. R. 100. 639.)
 $\text{N}_2\text{O}_5+1\frac{1}{2}\text{H}_2\text{O}$. Sl. sol. in pure H_2O . H_2O containing HCl . (Berzelius.)
) . Sol. in 119.2 pts. cold, and 10.5 ag H_2O . Sol. in $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2+\text{Aq}$, l. in KNO_3+Aq . (Vogel, jr. A. 94.

O , $3\text{N}_2\text{O}_5+5\text{H}_2\text{O}$. (Wakemann and n. Ch. J. 9. 299.)
) . (André, C. R. 100. 639.)
 $\text{N}_2\text{O}_5+\text{H}_2\text{O}$. Nearly insol. in H_2O . pr. 98. 385.)
, $3\text{N}_2\text{O}_5+4\text{H}_2\text{O}$. Less sol. in H_2O $\text{NO}_3)\text{OH}$, and not decomp. by boiling Wakemann and Wells, Am. Ch. J. 9.

ate, $\text{Pb}(\text{NO}_3)_2$.
 H_2O with absorption of much heat.

$\text{b}(\text{NO}_3)_2$ dissolves in $7\frac{1}{2}$ pts. cold H_2O . ich.)
 $(\text{NO}_3)_2$ dissolves in 1.989 pts. H_2O at 17.5° a liquid of 1.3978 sp. gr. (Karsten.)
 $(\text{NO}_3)_2$ dissolves in 1.707 pts. H_2O at 22.3° ; a. H_2O at 24.7° . (Kopp.)
.87 pts. H_2O at 17.5° . (Schiff, A. 109. 326.)
 $\text{Pb}(\text{NO}_3)_2+\text{Aq}$ sat. at 102.2° contain 52.5 $\text{O}_3)_2$, or 100 pt. H_2O dissolve 110.526 pts. at 102.2° . (Griffiths.)
7.5 pts. cold H_2O and much less hot H_2O .)
. boiling H_2O dissolve 13 pts. $\text{Pb}(\text{NO}_3)_2$. t.)

s. $\text{Pb}(\text{NO}_3)_2+\text{Aq}$ sat. at $19-20^\circ$ con-0 pts. salt. (v. Hauer, W. A. B. 53, 2.

lissolves:
 10° 25° 45° 65° 85° 100°
.07 1.65 1.25 0.99 0.83 0.72 pts. H_2O .
(Kremers, Pogg. 92. 497.)

$\text{Pb}(\text{NO}_3)_2+\text{Aq}$ sat. at 15° contains . $\text{Pb}(\text{NO}_3)_2$ and 928.58 g. H_2O , and gr. 1.39. (Michel and Krafft, A. ch. 171.)

Solubility in 100 pts. H_2O at t° .					
t°	Pts. $\text{Pb}(\text{NO}_3)_2$	t°	Pts. $\text{Pb}(\text{NO}_3)_2$	t°	Pts. $\text{Pb}(\text{NO}_3)_2$
0	36.5	36	65.9	72	99.7
1	37.4	37	66.7	73	100.7
2	38.3	38	67.6	74	101.7
3	39.1	39	68.5	75	102.6
4	39.8	40	69.4	76	103.6
5	40.5	41	70.3	77	104.6
6	41.2	42	71.2	78	105.6
7	42.0	43	72.1	79	106.6
8	42.8	44	73.0	80	107.6
9	43.6	45	74.0	81	108.6
10	44.4	46	74.9	82	109.6
11	45.2	47	75.9	83	110.6
12	46.0	48	76.8	84	111.5
13	46.8	49	77.7	85	112.5
14	47.5	50	78.7	86	113.5
15	48.3	51	79.6	87	114.5
16	49.1	52	80.5	88	115.4
17	49.9	53	81.5	89	116.4
18	50.7	54	82.4	90	117.4
19	51.5	55	83.3	91	118.4
20	52.3	56	84.3	92	119.4
21	53.1	57	85.2	93	120.3
22	53.9	58	86.1	94	121.3
23	54.7	59	87.1	95	122.3
24	55.6	60	88.0	96	123.2
25	56.4	61	89.0	97	124.2
26	57.3	62	90.0	98	125.2
27	58.1	63	90.9	99	126.1
28	59.0	64	91.9	100	127.0
29	59.8	65	92.8	101	128.0
30	60.7	66	93.8	102	128.9
31	61.6	67	94.8	103	129.9
32	62.4	68	95.7	104	130.9
33	63.3	69	96.7	104.7	131.5
34	64.1	70	97.7
35	65.0	71	98.7

(Mulder, Scheik. Verhandel. 1864. 66.)

100 g. H_2O dissolve 52.76 g. $\text{Pb}(\text{NO}_3)_2$ at 17° . (Euler, Z. phys. Ch. 1904, 49. 315.)
Solubility of $\text{Pb}(\text{NO}_3)_2$ in H_2O at $20^\circ=1.52$ g. mol. per l. Sp. gr. of sat. solution = 1.419. (Fedotieff, Z. anorg. 1911, 73. 178.)
Sat. $\text{Pb}(\text{NO}_3)_2+\text{Aq}$ at 0° contains 26.7% $\text{Pb}(\text{NO}_3)_2$; at 18° , 29.1% $\text{Pb}(\text{NO}_3)_2$. (Mylius, Z. anorg. 1912, 74. 411.)

Sp. gr. of $\text{Pb}(\text{NO}_3)_2+\text{Aq}$ at 19.5° .

$\text{Pb}(\text{NO}_3)_2$	Sp.gr.	$\text{Pb}(\text{NO}_3)_2$	Sp. gr.
5	1.045	25	1.266
10	1.093	30	1.334
15	1.144	35	1.414
20	1.203

(Kremers, calculated by Gerlach, Z. anal. 8. 286.)

Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at 17.5° .

$\text{Pb}(\text{NO}_3)_2$	Sp. gr.	$\text{Pb}(\text{NO}_3)_2$	Sp. gr.
5	1.044	25	1.263
10	1.092	30	1.333
15	1.144	35	1.409
20	1.200	sat. sol.	1.433

(Gerlach, Z. anal. 27. 283.)

Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ sat. at $8^\circ = 1.372$. (Anthon.)Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at 17.5° .

$\text{Pb}(\text{NO}_3)_2$	Sp. gr.	$\text{Pb}(\text{NO}_3)_2$	Sp. gr.
1	1.0080	20	1.1902
2	1.0163	21	1.2016
3	1.0247	22	1.2132
4	1.0331	23	1.2251
5	1.0416	24	1.2372
6	1.0502	25	1.2495
7	1.0591	26	1.2620
8	1.0682	27	1.2747
9	1.0775	28	1.2876
10	1.0869	29	1.3907
11	1.0963	30	1.3140
12	1.1059	31	1.3276
13	1.1157	32	1.3416
14	1.1257	33	1.3558
15	1.1359	34	1.3702
16	1.1463	35	1.3848
17	1.1569	36	1.3996
18	1.1677	37	1.4146
19	1.1788

(Schiff, calculated by Gerlach, Z. anal. 8. 286.)

Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at t° .

t°	% $\text{Pb}(\text{NO}_3)_2$	Sp. gr.
14	5	1.0451
14	10	1.0939
14.5	15	1.1468
14.3	20	1.2045
15	25	1.2678
15	32.28	1.3716

(Long, W. Ann. 1880, 11. 40.)

Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at room temp. containing:

17.93 32.22% $\text{Pb}(\text{NO}_3)_2$.
1.1786 1.3619

(Wagner, W. Ann. 1883, 18. 267.)

Sp. gr. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ at 25° .

Concentration of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$	Sp. gr.
1-normal	1.1380
$\frac{1}{2}$ -	1.0699
$\frac{1}{4}$ -	1.0351
$\frac{1}{8}$ -	1.0175

(Wagner, Z. phys. Ch. 1890, 5. 36.)

 $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ containing 15.93% $\text{Pb}(\text{NO}_3)_2$ has sp. gr. $20^\circ/20^\circ = 1.1558$. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ containing 30.57% $\text{Pb}(\text{NO}_3)_2$ has sp. gr. $20^\circ/20^\circ = 1.3436$. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ containing 30.69% $\text{Pb}(\text{NO}_3)_2$ has sp. gr. $20^\circ/20^\circ = 1.3465$.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 279.)

Sat. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ boils at 103.5° . (Kreners.)Sat. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ boils at 102.2° , and contains 140 pts. $\text{Pb}(\text{NO}_3)_2$ to 100 pts. H_2O (Griffiths.)Sat. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ boils at 103.5° . (Gerlach, Z. anal. 26. 427.)B.-pt. of $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ containing p $\text{Pb}(\text{NO}_3)_2$ to 100 pts. H_2O , according Gerlach (Z. anal. 26. 449).

B.-pt.	Pts. $\text{Pb}(\text{NO}_3)_2$	B.-pt.	Pts. $\text{Pb}(\text{NO}_3)_2$
100.5°	11	102.5°	87
101	26	103	111
101.5	44	103.5	137
102	65

Insol. in conc. $\text{HNO}_3 + \text{Aq}$.Solubility of $\text{Pb}(\text{NO}_3)_2 + \text{Ba}(\text{NO}_3)_2$.See under $\text{Ba}(\text{NO}_3)_2$.Solubility of $\text{Pb}(\text{NO}_3)_2 + \text{Cu}(\text{NO}_3)_2$.See under $\text{Cu}(\text{NO}_3)_2$.Sol. in sat. $\text{KNO}_3 + \text{Aq}$ without pptn., 11 pts. H_2O at 18.75° dissolving 114 pts. mix salt, viz. 84.1 pts. $\text{Pb}(\text{NO}_3)_2$ and 29.9 pts. KNO_3 . (Karsten.)100 pts. H_2O dissolve 119.6 pts. $\text{Pb}(\text{NO}_3)_2$ and 67.1 pts. KNO_3 at 21.2° . (Rüdorff, B. 484.)100 g. H_2O dissolve 95.39 g. $\text{Pb}(\text{NO}_3)_2$ and 61.05 g. KNO_3 at 20° . (Le Blanc and Noy, Z. phys. Ch. 1890, 6. 386.)Sol. in sat. $\text{NaNO}_3 + \text{Aq}$ without pptn., 11 pts. H_2O at 18.75° dissolving 121.9 pts. mix salt, viz. 87.8 pts. $\text{Pb}(\text{NO}_3)_2$ and 34.1 pts. NaNO_3 . (Karsten.)Solubility of $\text{Pb}(\text{NO}_3)_2 + \text{NaNO}_3$.Solid phase = $\text{Pb}(\text{NO}_3)_2$.

t° of saturation	% NaNO_3	% $\text{Pb}(\text{NO}_3)_2$
32	34.42	19.00
35.5	34.15	20.33
39.5	33.71	21.35
44.	33.35	22.19
49.1	32.94	23.15
55	32.60	23.93
58	32.47	24.24
62	32.33	24.57
65	32.14	24.60

ty of $\text{Pb}(\text{NO}_3)_2 + \text{NaNO}_3$ —Continued.

Solid phase = NaNO_3		
uration	% NaNO_3	% $\text{Pb}(\text{NO}_3)_2$
5	40.97	13.62
	42.04	13.38
	43.18	12.88
8	44.63.	12.78
	45.11	12.94
25	46.03	12.45
	47.28	12.50
	49.03	11.76
	49.92	11.56

saac, Chem. Soc. 1908, 93. 398.)

sol. in $\text{KNO}_3 + \text{NaNO}_3 + \text{Aq.}$
pts. sat. $\text{Pb}(\text{NO}_3)_2 + \text{Sr}(\text{NO}_3)_2 + \text{Aq}$
45.98 pts. of the two salts at 19.20°.
ier, J. pr. 98. 137.)

ility of $\text{Pb}(\text{NO}_3)_2 + \text{Sr}(\text{NO}_3)_2$ at 25°.

per 100 cc.		Mol. per cent in solid phase	
$\text{Pb}(\text{NO}_3)_2$	$\text{Sr}(\text{NO}_3)_2$	$\text{Pb}(\text{NO}_3)_2$	$\text{Sr}(\text{NO}_3)_2$
1	0	100.	0
7	4.56	99.05	0.95
2	8.14	98.11	1.89
4	17.81	97.02	2.98
8	18.74	96.06	3.94
3	35.03	83.84	16.16
3	37.54	32.88	67.12
	71.04	0	100.

ock, Z. Kryst. Min. 1897, 28. 365.)

easily sol. in liquid NH_3 . (Franklin,
h. J. 1898, 20. 828.)
pts. alcohol of 0.9282 sp. gr. dissolve:
8° 22° 40° 50°
5.82 8.77 12.8 14.9 pts. $\text{Pb}(\text{NO}_3)_2$.
(Gerardin, A. ch. (4), 5. 129.)

pts. absolute methyl alcohol dissolve
ts. at 20.5°.
pts. absolute ethyl alcohol dissolve 0.04
20.5°. (de Bruyn. Z. phys. Ch. 10.

y sl. sol. in acetone. (Krug and M'El-
. Anal. Ch. 6. 184.)
sl. in cold, sl. sol. in hot CS_2 . (Arctow-
. anorg. 1894, 6. 257.)
sl. in benzonitrile. (Naumann, B. 1914,
70.)
sl. in methyl acetate. (Naumann, B.
42. 3790); ethyl acetate. (Naumann,
10, 43. 314.)
l. weight determined in pyridine. (Wer-
. anorg. 1897, 15. 21.)

Solubility of $\text{Pb}(\text{NO}_3)_2$ in pyridine at t°.

t°	G. $\text{Pb}(\text{NO}_3)_2$ per 100 g. $\text{C}_5\text{H}_5\text{N}$	Solid phase
—19.4	2.93	$\text{Pb}(\text{NO}_3)_2, 4\text{C}_5\text{H}_5\text{N}$
—14.5	2.14	"
—10	1.90	"
0	3.54	"
5.4	3.93	"
8.7	5.39	"
14.72	6.13	"
19.97	6.78	"
24.75	8.56	"
30.03	10.98	"
34.97	13.20	"
40.03	16.94	"
45.	22.03	"
49.97	29.37	"
51 tr. pt.	...	" + $\text{Pb}(\text{NO}_3)_2, 3\text{C}_5\text{H}_5\text{N}$
59.52	36.70	$\text{Pb}(\text{NO}_3)_2, 3\text{C}_5\text{H}_5\text{N}$
70	47.29	"
80	61.60	"
89.93	90.21	"
94.94	128.06	"
96 tr. pt.	...	" + $\text{Pb}(\text{NO}_3)_2, 2\text{C}_5\text{H}_5\text{N}$
99.89	143.36	$\text{Pb}(\text{NO}_3)_2, 2\text{C}_5\text{H}_5\text{N}$
104.90	152	"
109.90	163.80	"

(Walton and Judd, J. Am. Chem. Soc. 1911,
33. 1036.)

Lead mercurous nitrate, $2\text{PbO}, 2\text{Hg}_2\text{O}, 3\text{N}_2\text{O}_5$.
Decomp. by H_2O . Sol. in warm dil. HNO_3 ,
or $\text{Hg}_2(\text{NO}_3)_2 + \text{Aq}$ without decomp. (Städ-
eler, A. 87. 129.)

Lead silver nitrate, $\text{Pb}(\text{NO}_3)_2, 2\text{AgNO}_3$.
Sol. in H_2O . (Stürenberg, Pogg. 74. 115.)

Lead silver nitrate iodide, $\text{Pb}(\text{NO}_3)_2, 8\text{AgNO}_3$,
 4AgI .
Decomp. by H_2O . (Stürenberg.)
 $\text{Pb}(\text{NO}_3)_2, 2\text{AgNO}_3, 2\text{AgI}$. Decomp. by
 H_2O . (Stürenberg.)

Lead nitrate nitrite, basic, $4\text{PbO}, \text{N}_2\text{O}_5, \text{N}_2\text{O}_3$,
 $+2\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3, \text{Pb}(\text{OH})\text{NO}_2$.
Sl. sol. in cold, easily in hot H_2O . Sol. in
80 pts. H_2O at 23° (Chevreuil); 85 pts. at ord.
temp. (Bromeis, A. 72. 38); 10.6 pts. at 100°
(Chevreuil).

+2H₂O.

Solubility in acetic acid.

Normality of acid	g. PbO per 100 cc. sat. solution	Normality of acid	g. PbO per 100 cc. sat. solution
0	0.601	0.25	5.450
0.05	1.323	0.50	9.690
0.10	2.185	0.75	15.874

(Chilesotti, Att. Acad. Linc. 1908, (5) 17, 18.
475.)

Formula is $3\text{Pb}(\text{OH})\text{NO}_3$, $5\text{Pb}(\text{OH})\text{NO}_3 + \text{H}_2\text{O}$. (v. Lorenz, W. A. B. 84, 2. 1133.)

+ $3\text{H}_2\text{O}$. (v. Lorenz.)

4PbO , N_2O_5 , $3\text{N}_2\text{O}_5 + 4\text{H}_2\text{O}$. Sol. in H_2O . (Bromeis.)

6PbO , N_2O_5 , $2\text{N}_2\text{O}_5 + 3\frac{1}{2}\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3$, $2\text{Pb}(\text{OH})\text{NO}_3 + \frac{1}{2}\text{H}_2\text{O}$. (v. Lorenz.)

6PbO , $2\text{N}_2\text{O}_5$, $\text{N}_2\text{O}_5 + 3\frac{1}{2}\text{H}_2\text{O} = 2\text{Pb}(\text{OH})\text{NO}_3$, $\text{Pb}(\text{OH})\text{NO}_3 + \frac{1}{2}\text{H}_2\text{O}$. (v. Lorenz.)

7PbO , N_2O_5 , $\text{N}_2\text{O}_5 + 3\text{H}_2\text{O}$. Less sol. in H_2O than 4PbO , N_2O_5 , $\text{N}_2\text{O}_5 + 2\text{H}_2\text{O}$; sol. in cold conc. $\text{HNO}_3 + \text{Aq}$. (Peligot, A. 39. 338.)

8PbO , N_2O_5 , $3\text{N}_2\text{O}_5 + 4\frac{1}{2}\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3$, $3\text{Pb}(\text{OH})\text{NO}_3 + \frac{1}{2}\text{H}_2\text{O}$. (v. Lorenz.)

10PbO , N_2O_5 , $4\text{N}_2\text{O}_5 + 5\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3$, $4\text{Pb}(\text{OH})\text{NO}_3$. (v. Lorenz.)

12PbO , N_2O_5 , $5\text{N}_2\text{O}_5 + 6\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3$, $5\text{Pb}(\text{OH})\text{NO}_3$. (v. Lorenz.)

10PbO , N_2O_5 , $2\text{N}_2\text{O}_5 + 4\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3$, $2\text{Pb}(\text{OH})\text{NO}_3$, $2\text{PbO} + \frac{1}{2}\text{H}_2\text{O}$. (v. Lorenz.)

14PbO , N_2O_5 , $3\text{N}_2\text{O}_5 + 6\text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_3$, $3\text{Pb}(\text{OH})\text{NO}_3$, $3\text{PbO} + \text{H}_2\text{O}$. (Bromeis.)

14PbO , $3\text{N}_2\text{O}_5$, $\text{N}_2\text{O}_5 + 6\text{H}_2\text{O} = 3\text{Pb}(\text{OH})\text{NO}_3$, $\text{Pb}(\text{OH})\text{NO}_3$, $3\text{PbO} + \text{H}_2\text{O}$. (Bromeis.)

16PbO , $2\text{N}_2\text{O}_5$, $3\text{N}_2\text{O}_5 + 6\text{H}_2\text{O} = 4\text{Pb}(\text{OH})\text{NO}_3$, $6\text{Pb}(\text{OH})\text{NO}_3$, 5PbO , $\text{Pb}(\text{OH})_2$. (v. Lorenz.)

16PbO , $3\text{N}_2\text{O}_5$, $5\text{N}_2\text{O}_5 + 10\text{H}_2\text{O} = 3\text{Pb}(\text{OH})\text{NO}_3$, $5\text{Pb}(\text{OH})\text{NO}_3 + \text{H}_2\text{O}$. (v. Lorenz.)

26PbO , $6\text{N}_2\text{O}_5$, $7\text{N}_2\text{O}_5 + 21\text{H}_2\text{O} = 6\text{Pb}(\text{OH})\text{NO}_3$, $7\text{Pb}(\text{OH})\text{NO}_3 + 4\text{H}_2\text{O}$. (v. Lorenz.)

Lead nitrate phosphate, $\text{Pb}(\text{NO}_3)_2$, $\text{Pb}_3(\text{PO}_4)_2 + 2\text{H}_2\text{O}$.

Completely insol. in cold H_2O . Decomp. by boiling H_2O into its constituents. Sol. in a little conc. $\text{HNO}_3 + \text{Aq}$ without decomp. (Gerhardt, A. 72. 83.)

Lead nitrate phosphite, $\text{Pb}(\text{NO}_3)_2$, PbHPO_3 .

Decomp. by H_2O . Sol. in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$ (33.3 g. per litre) dissolves 1 g. salt at 15° . If less than 31 g. per litre of $\text{Pb}(\text{NO}_3)_2$ are present the salt is decomp. (Amat, A. ch. (6) 24. 317.)

Lead nitrate potassium nitrite, $\text{Pb}(\text{NO}_3)_2$, $2\text{KNO}_2 + \text{H}_2\text{O}$.

Difficultly sol. in H_2O . (Lang, J. B. 1862. 102.)

3PbO , $3\text{K}_2\text{O}$, $4\text{N}_2\text{O}_5$, $2\text{N}_2\text{O}_5 + 3\text{H}_2\text{O}$. Sol. in H_2O . (Hayes, Sill. Am. J. (2) 31. 226.)

Lithium nitrate, LiNO_3 .

Very deliquescent, and sol. in H_2O .

100 pts. H_2O dissolve:

at 0° 20° 40° 70° 100° 110°

48.3 75.7 169.4 196.1 227.3 256.4 pts. LiNO_3 . (Kremers, Pogg. 99. 47.)

Forms supersaturated solutions which crystallise when temp. is $10^\circ + 1^\circ$. (Kremers, Pogg. 99. 520.)

Sat. solution boils at over 200° . (Pogg. 99. 43.)

1 pt. LiNO_3 dissolves in 200 pts (Schultz, Zeit. Ch. (2) 5. 531.)

100 pts. of the sat. solution contain

64.2° 70.9°

64.9 66.1 pts. anhydrous

(Donnan and Burt, Chem. Soc. 339.)

See $+\frac{1}{2}\text{H}_2\text{O}$, and $3\text{H}_2\text{O}$.

Sp. gr. of $\text{LiNO}_3 + \text{Aq}$ at 19.5° pts. LiNO_3 in 100 pts. H_2O :

12.7	14.2	26.4	41.8 p
1.069	1.077	1.134	1.197
54.8	57.5	77.4	79.4 p
1.245	1.255	1.315	1.319

(Kremers, Pogg. 114. 45.)

Sp. gr. of $\text{LiNO}_3 + \text{Aq}$.

g. LiNO_3 in 1000 g. of solution	Sp. gr.
0	1.000
4 8526	1.002
10 9128	1.005
17 9016	1.009

(Dijken, Z. phys. Ch. 1897, 24.)

Sp. gr. $20^\circ/4^\circ$ of a normal solution = 1.03803; of a 0.5-normal solution (Haigh, J. Am. Chem. Soc. 1912, 34)

Very easily sol. in liquid NH_3 . (Am. Ch. J. 1898, 20. 828.)

Sol. in strong alcohol.

Sol. in acetone. (Eidmann, C. II. 1014; Naumann, B. 1904, 37. 4)

Solubility in acetone = 0.343 g. at 18° (Roshdestwensky and Chem. Soc. 1911, 99. 2140.)

Insol. in benzonitrile. (Naur 1914, 47. 1370.)

Difficultly sol. in ethyl acetate. (mann, B. 1910, 43. 314.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Solubility in H_2O .

100 pts. of the sat. solution contain

43.6° 50.5° 55.0° 60.0°

60.8 61.3 63.0 63.6 pts. anhy

61.1° is the temp. at which LiNO_3 goes over into Li_2NO_3 . (Donnan: Chem. Soc. 1903, 82. 339.)

3H₂O.

Solubility in H₂O.

0 pts. of the sat. solution contain pts. anhydrous salt at t°.

t°	Pts. anhydrous salt
0.10	34.8
10.50	37.9
12.10	38.2
13.75	39.3
19.05	40.4
22.10	42.9
27.55	47.3
29.47	53.67
29.78	55.09
29.87	56.42
29.86	56.68
29.64	57.48
29.55	58.03

pt. of LiNO₃+3H₂O is 29.88°.

Donnan and Burt, Chem. Soc. 1903, 83. 337.)

magnesium nitrate, basic, Mg₃N₂O₈.

sol. in H₂O and alcohol. Sol. in acids. (Dewar, A. 71. 241.)

5H₂O. Decomp. by H₂O. (Didier, C. R. 122. 936.)

magnesium nitrate, Mg(NO₃)₂.

anhydrous. Deliquescent.

in 1 pt. H₂O at 15.6°. Sol. in 4 pts. abs. alcohol at 15°, and 2 pts. at boiling temp. More sol. in alcohol than in that of 0.900. (Kirwan.)
in 0.3458 pt. strong alcohol at 82.5°. (Wenzel.)
in 10 pts. strong alcohol at 15°. (Bergmann.)
in 9 pts. strong alcohol on heating. (Bergmann.)

solubility in H₂O in presence of the anhydrous salt.

the solution contains at:

89° 77.5° 67°
3.14 65.67 67.55% Mg(NO₃)₂.
(Funk, Z. anorg. 1899, 20. 396.)
+6, and 9H₂O.

Sp. gr. of Mg(NO₃)₂+Aq at 14°.

(NO ₃) ₂ · H ₂ O	Sp. gr.	% Mg(NO ₃) ₂ · 6H ₂ O	Sp. gr.
1	1.0034	30	1.1347
5	1.0202	35	1.1649
9	1.0418	40	1.1909
5	1.0639	45	1.2176
9	1.0869	49	1.2397
5	1.1103

(Oudemans, Z. anal. 7. 419.)

Sp. gr. of Mg(NO₃)₂+Aq at 21°.

% Mg(NO ₃) ₂ · +6H ₂ O	Sp. gr.	% Mg(NO ₃) ₂ · +6H ₂ O	Sp. gr.
2	1.0078	28	1.1216
4	1.0158	30	1.1312
6	1.0239	32	1.1410
8	1.0321	34	1.1508
10	1.0405	36	1.1608
12	1.0490	38	1.1709
14	1.0577	40	1.1811
16	1.0663	42	1.1914
18	1.0752	44	1.2019
20	1.0843	46	1.2124
22	1.0934	48	1.2231
24	1.1026	50	1.2340
26	1.1120

(Schiff, calculated by Gerlach, Z. anal. 8. 286.)

Sp. gr. of Mg(NO₃)₂+Aq at 18°.

% Mg(NO ₃) ₂	Sp. gr.	% Mg(NO ₃) ₂	Sp. gr.
5	1.0378	15	1.1181
10	1.0763	17	1.1372

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of Mg(NO₃)₂+Aq at room temp. containing:

18.62 34.19 39.77% Mg(NO₃)₂.
1.1025 1.2000 1.4298

(Wagner, W. Ann. 1883, 18. 273.)

Sp. gr. of Mg(NO₃)₂+Aq at 25°.

Concentration of Mg(NO ₃) ₂ +Aq	Sp. gr.
1-normal	1.0512
1/2- " "	1.0259
1/4- " "	1.0130
1/8- " "	1.0066

(Wagner, Z. phys. Ch. 1890, 5. 38.)

Sp. gr. of solution sat. at 18° = 1.384, containing 43.1% Mg(NO₃)₂. (Mylius, B. 1897, 30. 1718.)

Sp. gr. of Mg(NO₃)₂+Aq.

1/2 Mg(NO ₃) ₂ g. in 1000 g. of solution	Sp. gr. 16°/16°
0	1.000000
0.8099	1.000660
1.5621	1.001253
3.3398	1.002539
7.4410	1.005523
15.161	1.011151
29.356	1.021580
58.353	1.043329
81.025	1.060773

(Dijken, Z. phys. Ch. 1897, 24. 107.)

Sp. gr. of $\text{Mg}(\text{NO}_3)_2 + \text{Aq}$ at 20.1° .
p = per cent strength of solution; d = observed density; w = volume concentration in grams per cc. ($\frac{pd}{100} = w$.)

p	d	w
35.02	1.3110	0.46695
31.15	1.2655	0.39420
25.03	1.2057	0.30172
19.55	1.1551	0.22585
13.43	1.1028	0.14815
10.09	1.0753	0.10850
6.650	1.0480	0.06968
4.672	1.0330	0.04826
4.001	1.0276	0.04112
1.372	1.0085	0.01383

(Barnes, J. phys. Chem. 1898, 2. 545.)

Sp. gr. of $\text{Mg}(\text{NO}_3)_2 + \text{Aq}$ at 20° containing M g. mols. of salt per liter.

M	0.02	0.05	0.10	0.15
Sp. gr.	1.00224	1.005626	1.011118	1.016557

M	0.20	0.50	1.00	1.274
Sp. gr.	1.022026	1.054804	1.107865	1.136615

(Jones and Pearce, Am. Ch. J. 1907, 38. 707.)

Less sol. in $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ than in H_2O . (Dijonval.)
Very easily sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)
+ $2\text{H}_2\text{O}$. Mpt. 127° . (Wasiljew, C. C. 1909, II. 1966.)
+ $4\text{H}_2\text{O}$. Mpt. 45.5° . (W.)
+ $6\text{H}_2\text{O}$. Deliquescent. Sol. in H_2O and alcohol. Sol. in 0.5 pt. cold H_2O , and 9 pts. cold alcohol of 0.84 sp. gr.; very sl. sol. in abs. alcohol. (Graham.)
Melts in its crystal H_2O at 90° , and the resulting liquid boils at 143.4° . (Ordway, Sill. Am. J. (2) 27. 14.)

Solubility in H_2O .
Sat. solution contains at:

-18°	-4.5°	0°	18°
38.03	39.50	39.96	42.33% $\text{Mg}(\text{NO}_3)_2$
40°	80°	90° (mpt.)	
45.87	53.69	57.81% $\text{Mg}(\text{NO}_3)_2$	

(Funk, Z. anorg. 1899, 20. 305.)

+ $9\text{H}_2\text{O}$. Solubility in H_2O .
Sat. solution contains at:

-23°	-20.5°	-18°
35.44	36.19	38.03% $\text{Mg}(\text{NO}_3)_2$

Cryohydrate is formed at -29° . (Funk, Z. anorg. 1899, 20. 398.)

Magnesium neodymium nitrate, $3\text{Mg}(\text{NO}_3)_2$, $2\text{Nd}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.
1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 97.7 g. hydrous salt at 16° . (Jantsch, Z. anorg. 1912, 76. 303.)

Magnesium praseodymium nitrate, $3\text{Mg}(\text{NO}_3)_2$, $2\text{Pr}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.
1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 7.70 g. hydrous salt (Jantsch.)

Magnesium samarium nitrate, $3\text{Mg}(\text{NO}_3)_2$, $\text{Sm}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.
(Demarçay, C. R. 1900, 130. 1187.)
1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 24.55 g. hydrous salt (Jantsch.)

Magnesium thorium nitrate, $\text{MgTh}(\text{NO}_3)_6 + 6\text{H}_2\text{O}$.
Hydroscopic; sol. in HNO_3 . (Mc anorg. 1901, 27. 385.)

Magnesium nitrate ammonia, $\text{Mg}(\text{NO}_3)_2 \cdot 6\text{NH}_3$.
Sl. sol. in liquid NH_3 . (Franklin, Chem. Soc. 1913, 35. 1459.)

Manganous nitrate, basic, 2MnO , $3\text{H}_2\text{O}$.
Sol. in H_2O . (Gorgeu.)

Manganous nitrate, $\text{Mn}(\text{NO}_3)_2$.
Deliquescent. Easily sol. in H_2O and alcohol.
See +3, and $6\text{H}_2\text{O}$.

Sp. gr. of $\text{Mn}(\text{NO}_3)_2 + \text{Aq}$ at 8°

% $\text{Mn}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$	Sp. gr.	% $\text{Mn}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$
5	1.0253	45
10	1.0517	50
15	1.0792	55
20	1.1078	60
25	1.1137	65
30	1.1688	70
35	1.2012	71
40	1.2352	..

(Oudemans, Z. anal. 7. 421.)

Sp. gr. of aqueous solutions contain

10	20	30	% $\text{Mn}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$
6.237	12.474	18.711	% $\text{Mn}(\text{NO}_3)_2$
1.052	1.107	1.165	
40	50	60	% $\text{Mn}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$
24.948	31.185	37.422	% $\text{Mn}(\text{NO}_3)_2$
1.230	1.302	1.381	
70	80	% $\text{Mn}(\text{NO}_3)_2 + 6\text{H}_2\text{O}$	
43.659	49.896	% $\text{Mn}(\text{NO}_3)_2$	
1.466	1.558		

(Gerlach, Z. anal. 28. 477.)

Sp. gr. of $\text{Mn}(\text{NO}_3)_2 + \text{Aq}$ at room containing:

18.309	29.602	49.309% $\text{Mn}(\text{NO}_3)_2$
1.1482	1.3227	1.5056

(Wagner, W. Ann. 1892, 12. 271)

Sp. gr. of $\text{Mn}(\text{NO}_3)_2 + \text{Aq}$ at 25°.	
Concentration of $\text{Mn}(\text{NO}_3)_2 + \text{Aq}$	Sp. gr.
1-normal	1.0690
$\frac{1}{2}$ " "	1.0349
$\frac{1}{4}$ " "	1.0174
$\frac{1}{8}$ " "	1.0093

(Wagner, Z. phys. Ch. 1890, 5. 39.)

Sol. in liquid NH_3 . (Guntz, Bull. Soc. 1909, (4) 5. 1006.)

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

+ H_2O . Deliquescent. (Guntz, Bull. Soc. 1909 (4) 5. 1005.)

+3 H_2O . From solution in HNO_3 . (Schultz-Sellack, Zeit. Ch. 1870. 646.)

Solubility in H_2O .

Sat. solution contains at:

27° 29° 30° 34° 35.5° mpt.
65.66 66.99 67.38 71.31 76.82% $\text{Mn}(\text{NO}_3)_2$.
(Funk, Z. anorg. 1899, 20. 403.)

+6 H_2O . Melts in its crystal H_2O at 25.8° and boils at 129.4°. (Ordway.)

Solubility in H_2O .

Sat. solution contains at:

-29° -26° -21° -16° -5°
12.29 43.15 44.30 45.52 48.88% $\text{Mn}(\text{NO}_3)_2$,

0° +11° 18° 25.8° mpt.
10.49 54.50 57.33 62.37% $\text{Mn}(\text{NO}_3)_2$.

Cryohydrate is formed at -36°. (Funk, Z. anorg. 1899, 20. 403.)

Manganous neodymium nitrate, $3\text{Mn}(\text{NO}_3)_2$, $2\text{Nd}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 296 g. hydrous salt at 16°. (Jantsch, Z. anorg. 1912, 76. 303.)

Manganous praseodymium nitrate, $3\text{Mn}(\text{NO}_3)_2$, $2\text{Pr}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 23.4 g. hydrous salt at 16°. (Jantsch.)

Manganous samarium nitrate, $3\text{Mn}(\text{NO}_3)_2$, $2\text{Sm}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 50.04 g. hydrous salt at 16°. (Jantsch.)

Manganous thorium nitrate, $\text{MnTh}(\text{NO}_3)_6 + 8\text{H}_2\text{O}$.

Ppt. (Meyer, Z. anorg. 1901, 27. 388.)

Manganous nitrate cupric oxide, $\text{Mn}(\text{NO}_3)_2$, $3\text{CuO} + 3\text{H}_2\text{O}$.

Ppt. (Mailhe, C. R. 1902, 134. 234.)

Manganous nitrate hydrazine, $\text{Mn}(\text{NO}_3)_2$, $2\text{N}_2\text{H}_4$.

Not decomp. by H_2O . (Franzen, Z. anorg. 1908, 60. 286.)

Mercurous nitrate, basic, $2\text{Hg}_2\text{O}$, $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$.

Ppt. Decomp. by boiling with H_2O . (Marignac, A. ch. (3) 27. 332.)

Slowly sol. in cold, rapidly in hot $\text{HCl} + \text{Aq}$; insol. in NH_4Cl , and $\text{NH}_4\text{NO}_3 + \text{Aq}$.

+10 H_2O . Slowly sol. in normal HNO_3 . (Reuss, Dissert. 1886.)

4 Hg_2O , 3 $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. Sol. in a small quantity of H_2O ; decomp. by a large amt. of H_2O or by warm H_2O . (Rose, Pogg. 83. 154.)

Is 3 HgO , 2 $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$ according to Gerhardt.

+5 H_2O . (Reuss, Dissert. 1886.)

5 Hg_2O , 3 $\text{N}_2\text{O}_5 + 2\text{H}_2\text{O}$. (Marignac.) Is 2 Hg_2O , $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. (Lefort, A. 56. 247.)

Sol. in boiling, less sol. in cold H_2O . (Marignac, l. c.)

+4 H_2O , and +6 H_2O . (Reuss, Dissert. 1886.)

8 Hg_2O , 5 $\text{N}_2\text{O}_5 + 5\text{H}_2\text{O}$, and +11 H_2O . (Reuss.)

11 Hg_2O , 6 $\text{N}_2\text{O}_5 + 25\text{H}_2\text{O}$. (Reuss.)

16 Hg_2O , 9 $\text{N}_2\text{O}_5 + 19\text{H}_2\text{O}$, +23 H_2O , and +31 H_2O . (Reuss.)

3 Hg_2O , $\text{N}_2\text{O}_5 + 2\text{H}_2\text{O}$. (Cox, Z. anorg. 1904, 40. 177.)

Mercurous nitrate, HgNO_3 .

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Fairly sol. in boiling CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in methylamine. (Franklin, J. Am. Chem. Soc. 1906, 28. 1419.)

+ H_2O . Completely sol. in a little warm H_2O , but decomp. by more H_2O . Completely sol. as acid salt in H_2O containing HNO_3 . (Marignac, A. ch. (3) 27. 332.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

+1 $\frac{1}{4}$ H_2O , +1 $\frac{1}{2}$ H_2O , +1 $\frac{1}{2}$ H_2O , etc. (Reuss, Dissert. 1896.)

Mercuric nitrate, basic, 6HgO , N_2O_5 (?).

Insol. in hot H_2O . (Kane.)

3 HgO , $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. Decomp. to oxide by washing with cold H_2O . Sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Millon, A. ch. (3) 18. 361.)

2 HgO , $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. Sl. deliquescent. Decomp. by H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Millon.)

+2 H_2O . Decomp. by cold H_2O . Deliquescent. Sol. in H_2O containing HNO_3 . (Marignac.)

+3 H_2O . (Ditte, J. B. 1854. 366.)

Mercuric nitrate, $\text{Hg}(\text{NO}_3)_2$.

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Neither dissolved nor attacked by liquid NO_2 . (Frankland, Chem. Soc. 1901, 79. 1361.)

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Difficultly sol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Sol. in methylal. (Eidmann, C. C. 1899. II, 1014.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Deliquescent. Very sol. in a little H_2O . H_2O precipitates basic salt from conc. $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$. Insol. in alcohol. Decomp. by ether. (Millon.)

+ H_2O . Extraordinarily sol. in H_2O . (Cox, Z. anorg. 1904, 40. 159.)

+ $8\text{H}_2\text{O}$. Melts at 6° in crystal H_2O . (Ditte.)

Mercuromercuric nitrate, Hg_2O , 2HgO , N_2O_5 .

Boiling H_2O gradually dissolves out $\text{Hg}_2(\text{NO}_3)_2$, and leaves residue of HgO and Hg . (Brooks, Pogg. 66. 63.)

$2\text{Hg}_2\text{O}$, HgO , $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. (Ray, Chem. Soc. 1905, 87. 175.)

Hg_2O , 2HgO , $\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. (Ray.)

Mercurous hydrogen nitrate, 4HgNO_3 , HNO_3 , $+8\text{H}_2\text{O}$.

(Reuss, Dissert. 1886.)

5HgNO_3 , $3\text{HNO}_3 + 26\text{H}_2\text{O}$. (Reuss.)

Mercuric silver nitrate, $\text{Hg}(\text{NO}_3)_2$, 2AgNO_3 .

Easily sol. in H_2O without decomp. (Berzelius.)

Mercurous strontium nitrate, 2SrO , $2\text{Hg}_2\text{O}$, $3\text{N}_2\text{O}_5$.

Decomp. by H_2O . Much more sol. in H_2O than the corresponding Ba compound.

Readily sol. in warm dil. $\text{HNO}_3 + \text{Aq}$ or $\text{Hg}_2(\text{NO}_3)_2 + \text{Aq}$ without decomposition. (Städeler, A. 87. 131.)

Mercurous thallos nitrate, HgNO_3 , TiNO_3 .

Miscible with H_2O . (Retgers, N. Jahrb. Miner, 1896. II, 183.)

Mercuric nitrate bromide, $\text{Hg}(\text{NO}_3)_2$, HgBr_2 . (Morse, Z. phys. Ch. 1902, 41. 733.)

Mercuric nitrate cadmium oxide, $\text{Hg}(\text{NO}_3)_2$, $\text{CdO} + 2\text{H}_2\text{O}$.

Ppt. (Mailhe, Bull. Soc. 1901, (3) 25. 788.)

+ $3\text{H}_2\text{O}$. Decomp. by H_2O . (Mailhe.)

Mercuric nitrate cobaltous oxide, $\text{Hg}(\text{NO}_3)_2$, $\text{CoO} + 3\text{H}_2\text{O}$.

Ppt. Mailhe, C. R. 1901, 132. 1275.)

+ $4\text{H}_2\text{O}$. Decomp. by H_2O . (Mailhe, A. ch. 1902, (7) 27. 369.)

Mercuric nitrate cupric oxide, $\text{Hg}(\text{NO}_3)_2$, $\text{CuO} + 2\text{H}_2\text{O}$, and $+4\text{H}_2\text{O}$.

(Mailhe, Bull. Soc. 1901, (3) 25. 791.)

+ $5\text{H}_2\text{O}$. Decomp. by H_2O . (Mailhe, A. ch. 1902, (7) 27. 365.)

Mercuric nitrate cyanide, $\text{Hg}(\text{NO}_3)_2$, $\text{Hg}(\text{CN})_2$.

Very sol. in H_2O . Very sol. in methyl alcohol and solution is not decomp. at bpt. Ethyl alcohol apparently decomp. it. (Prassia, Gazz. ch. it. 1898, 28. (2) 115.)

Mercurous nitrate hydrazine, 2HgNO_3 , N_2H_4 .

Decomp. by H_2O . Stable in dil. $\text{HNO}_3 + \text{Aq}$ solution. (Hofmann and Marburg, A. 1899, 305. 215.)

Ppt.; very unstable. (Hofmann, B. 1897, 30. 2021.)

$\text{Hg}(\text{NO}_3)_2$, N_2H_4 . Sol. in dil. HCl and HNO_3 . (Hofmann and Marburg, A. 1899, 305. 215.)

Ppt.; sol. in acids; decomp. by alkali. (Hofmann, B. 1897, 30. 2021.)

Mercuric nitrate iodide, $\text{Hg}(\text{NO}_3)_2$, 2HgI_2 .

Decomp. by long boiling with H_2O . (Riegel, Jahrb. Pharm. 11. 396.)

$2\text{Hg}(\text{NO}_3)_2$, 3HgI_2 . Easily decomp. by H_2O ; less easily by alcohol or ether. (Riegel.)

$\text{Hg}(\text{NO}_3)_2$, HgI_2 . Decomp. very quickly by $\text{HNO}_3 + \text{Aq}$ or alcohol of 0.814 sp. gr. (Souville, J. Pharm. 26. 474.)

Mercuric nitrate manganous oxide, $\text{Hg}(\text{NO}_3)_2$, $\text{MnO} + 2\text{H}_2\text{O}$.

Decomp. by H_2O . (Mailhe, Bull. Soc. 1901, (3) 25. 790.)

+ $3\text{H}_2\text{O}$. (Mailhe.)

+ $4\text{H}_2\text{O}$. (Mailhe, A. ch. 1902, (7) 27. 370.)

Mercuric nitrate nickel oxide, $\text{Hg}(\text{NO}_3)_2$, $\text{NiO} + 2\text{H}_2\text{O}$.

(Mailhe, Bull. Soc. 1901, (3) 25. 788.)

+ $4\text{H}_2\text{O}$. Decomp. by H_2O . (Mailhe, A. ch. 1902, (7) 27. 369.)

Mercurous nitrate phosphate, HgNO_3 , $\text{Hg}_3\text{PO}_4 + \text{H}_2\text{O}$.

Insol. in H_2O , but decomp. by boiling therewith. Insol. in $\text{H}_3\text{PO}_4 + \text{Aq}$ or alcohol. Completely sol. in hot $\text{NH}_4\text{Cl} + \text{Aq}$. Decomp. by cold $\text{KOH} + \text{Aq}$, and warm $\text{K}_2\text{CO}_3 + \text{Aq}$. (Wittstein.)

2HgNO_3 , Hg_2O , $5\text{Hg}_3\text{PO}_4 + \text{H}_2\text{O}$. (Haack, A. 262. 192.)

Mercuric nitrate silver bromide, $\text{Hg}(\text{NO}_3)_2$, AgBr .

(Morse, Z. phys. Ch. 1902, 41. 733.)

nitrate silver cyanide, basic,
 $(\text{H})\text{NO}_3, \text{AgCN} + 2\text{H}_2\text{O}$.
dt, Z. anorg. 1895, 9. 431.)
 $(\text{H})\text{NO}_3, 5\text{Ag}_2\text{O}, 20\text{AgCN} + 7\text{H}_2\text{O}$.
)

nitrate silver iodide, $\text{Hg}(\text{NO}_3)_2$,
 $+ \frac{1}{2}\text{H}_2\text{O}$.
p. by H_2O . (Preuss, A. 29. 328.)

nitrate sulphide, $\text{Hg}(\text{NO}_3)_2, 2\text{HgS}$.
. sol. in hot H_2O . Insol. in HNO_3 +
comp. by hot H_2SO_4 or aqua regia,
not $\text{HCl} + \text{Aq}$. (Barfoed, J. pr. 93.

aqua regia. (Denigès, Bull. Soc.
17. 355.)
 $\text{O}_3)_2, \text{HgO}, 6\text{HgS} + 12\text{H}_2\text{O}$. Insol.
and $\text{HNO}_3 + \text{Aq}$ of 1.2 sp. gr. (Gramp,
14. 299.)

nitrate zinc oxide, $\text{Hg}(\text{NO}_3)_2, \text{ZnO} +$

Decomp. by H_2O . (Mailhe, C. R.
. 1274.)

ium nitrate, $\text{Mo}_2\text{O}_3, \text{N}_2\text{O}_5$ (?).
dil. $\text{HNO}_3 + \text{Aq}$. (Berzelius.)
 $2\text{N}_2\text{O}_5$ (?). Sol. in dil. $\text{HNO}_3 + \text{Aq}$.
i.)

um nickel nitrate, $2\text{Nd}(\text{NO}_3)_3$,
 $\text{NO}_3)_2 + 24\text{H}_2\text{O}$.
t. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr.
contains 116.6 g. hydrous salt at 16° .
Z. anorg. 1912, 76. 303.)

um rubidium nitrate, $[\text{Nd}(\text{NO}_3)_3]\text{Rb}_2$,
 O_3O .
copic. (Jantsch, Z. anorg. 1911,

um zinc nitrate, $2\text{Nd}(\text{NO}_3)_3$,
 $\text{NO}_3)_2 + 24\text{H}_2\text{O}$.
t. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr.
contains 177 g. hydrous salt at 16° .

rate, basic.
 $1 \text{H}_2\text{O}$. (Proust.)
 $\text{N}_2\text{O}_5 + 5\text{H}_2\text{O}$. Insol. in cold or hot
abermann, M. 5. 432.)
 $\text{N}_2\text{O}_5 + 4\text{H}_2\text{O}$. Not decomp. by boil-
(Rousseau and Tite, C. R. 114.

rate, $\text{Ni}(\text{NO}_3)_2$.
ty in H_2O . See +3, 6, and $9\text{H}_2\text{O}$.
of aqueous solution at 17.5° contain-

10 15 20 % $\text{Ni}(\text{NO}_3)_2$,
.0903 1.1375 1.1935

30 35 40 % $\text{Ni}(\text{NO}_3)_2$.
.3193 1.3896 1.4667
(Franz, J. pr. (2) 5. 295.)

Sp. gr. of $\text{Ni}(\text{NO}_3)_2 + \text{Aq}$ containing g.
 $\text{Ni}(\text{NO}_3)_2$ (anhydrous) in 1000 g. H_2O at
 24.4° :
91.5 g. (= $\frac{1}{2}$ mol.) 183 274.5 369 460.5 549
1.073 1.141 1.205 1.266 1.324 1.378
(Gerlach, Z. anal. 28. 468.)

Sp. gr. of $\text{Ni}(\text{NO}_3)_2 + \text{Aq}$ at room temp.
containing:
16.493 30.006 40.953% $\text{Ni}(\text{NO}_3)_2$.
1.1363 1.2776 1.3879
(Wagner, W. Ann. 1883, 18. 269.)

Concentration of $\text{Ni}(\text{NO}_3)_2 + \text{Aq}$		Sp. gr.
1-normal		1.0755
$\frac{1}{2}$ " "		1.0381
$\frac{1}{3}$ " "		1.0192
$\frac{1}{4}$ " "		1.0096

(Wagner, Z. phys. Ch. 1890, 5. 39.)

Sp. gr. at 20° of $\text{Ni}(\text{NO}_3)_2 + \text{Aq}$ containing
M g. mols. of salt per liter.
M 0.01 0.025 0.05 0.075
Sp. gr. 1.001521 1.003882 1.007792 1.011541
M 0.1 0.25 0.5 0.75
Sp. gr. 1.015307 1.03837 1.07611 1.11310
M 1.0 1.5 2.0
Sp. gr. 1.14562 1.22134 1.29459
(Jones and Pearce, Am. Ch. J. 1907, 38. 720.)

Sol. in liquid NH_3 . (Guntz, Bull. Soc.
1909, (4) 5. 1008.)
Moderately sol. in liquid NH_3 . (Franklin,
Am. Ch. J. 1898, 20. 828.)
Solubility in glycol = 7.5%. (de Coninck,
C. C. 1905, II. 1234.)
Insol. in benzonitrile. (Naumann, B. 1914,
47. 1370.)
 $+ 3\text{H}_2\text{O}$. Solubility in H_2O .
Sat. solution contains at:
58° 60° 64° 70°
61.61 61.99 62.76 63.95% $\text{Ni}(\text{NO}_3)_2$,
90° 95° mpt.
70.16 77.12% $\text{Ni}(\text{NO}_3)_2$.
(Funk, Z. anorg. 1899, 20. 411.)

$+ 6\text{H}_2\text{O}$. Not deliquescent in dry air. Sol.
in 2 pts. cold H_2O and in alcohol. (Tupputi.)
Mpt. of $\text{Ni}(\text{NO}_3)_2 + 6\text{H}_2\text{O} = 56.7^\circ$. (Ord-
way; Tilden, Chem. Soc. 45. 409.)
Sat. solution boils at 136.7° . (Ordway.)
Solubility in H_2O .
Sat. solution contains at:
-21° -12.5° -10° -6°
39.94 41.59 42.11 43.00% $\text{Ni}(\text{NO}_3)_2$,
0° +20° 41° 56.7° mpt.
44.32 49.06 55.22 62.76% $\text{Ni}(\text{NO}_3)_2$.
(Funk, Z. anorg. 1899, 20. 410.)

Sat. solution of $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ contains 44.3% $\text{Ni}(\text{NO}_3)_2$ at 0° , and 48.7% $\text{Ni}(\text{NO}_3)_2$ at 18° . (Mylius, Z. anorg. 1912, 74. 411.)

Sp. gr. of $\text{Ni}(\text{NO}_3)_2 \cdot \text{Aq}$ containing in 1000 g. H_2O at 24.4° , g. $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$.

145.5 g. ($= \frac{1}{2}$ mol.) 291 436.5 582
1.069 1.128 1.179 1.224

727.5 873 1018.5 1164
1.264 1.299 1.329 1.357

(Gerlach, Z. anal. 28. 468.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Insol. in absolute alcohol.

Sl. sol. in acetone. (Krug and M'Elroy.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

+9 H_2O . Solubility in H_2O .

Sat. solution contains at:

-23° -21° -10.5°
39.02 39.48 44.13% $\text{Ni}(\text{NO}_3)_2$.

Cryohydrate is formed at -27° . (Funk, Z. anorg. 1899, 20. 411.)

Nickel praseodymium nitrate, $3\text{Ni}(\text{NO}_3)_2$, $2\text{Pr}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 9.28 g. hydrous salt at 16° . (Jantsch, Z. anorg. 1912, 76. 303.)

Nickel samarium nitrate, $3\text{Ni}(\text{NO}_3)_2$, $2\text{Sm}(\text{NO}_3)_3 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 29.11 g. hydrous salt at 16° . (Jantsch.)

Nickel thorium nitrate, $\text{NiTh}(\text{NO}_3)_6 + 8\text{H}_2\text{O}$.

Sol. in $\text{HNO}_3 + \text{Aq}$. (Meyer, Z. anorg. 1901, 27. 387.)

Nickel uranyl nitrate, $10\text{Ni}(\text{NO}_3)_2$, $3(\text{UO}_2)(\text{NO}_3)_2$.

Sol. in H_2O and acids; insol. in aq. alkali. (Lancien, C. C. 1912, I. 208.)

Nickel nitrate ammonia, $\text{Ni}(\text{NO}_3)_2$, $4\text{NH}_3 + 2\text{H}_2\text{O}$.

Efflorescent. Easily sol. in cold H_2O ; decomp. by boiling. Insol. in alcohol. (Erdmann, J. pr. 97. 395; Ephraim, B. 1913, 46. 3106.)

+1 $\frac{1}{2}$ H_2O . (André, C. R. 106. 936.)

Nickel nitrate chloride ammonia, $6\text{Ni}(\text{NO}_3)_2$, NiCl_2 , $30\text{NH}_3 + 16\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Schwarz, W. A. B. 1850. 272.)

Nickel nitrate cupric oxide, $\text{Ni}(\text{NO}_3)_2$, $3\text{CuO} + 3\text{H}_2\text{O}$.

Ppt. (Maible, C. R. 1902, 134. 234.)

Nickel nitrate hydrazine, $\text{Ni}(\text{NO}_3)_2$.

Insol. in H_2O . Decomp. by heat. Easily sol. in dil. acids. (Fransen, Z. 1908, 60. 267.)

Palladium nitrate, basic, $\text{Pd}(\text{NO}_3)_2 + 4\text{H}_2\text{O}$.

Ppt. Insol. in H_2O . (Kane.)

Palladium nitrate, $\text{Pd}(\text{NO}_3)_2 + x\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O comp. by much H_2O or alcohol. (R.)

Decomp. by cold or hot H_2O . (R. 143.)

Platinic nitrate, $\text{Pt}(\text{NO}_3)_4$ (?).

Known only in solution, which is on evaporating. (Berzelius.)

$\text{Pt}(\text{NO}_3)_2$, $3\text{PtO}_2 + 5\text{H}_2\text{O}$. Insol. (Prost, Bull. Soc. (2) 46. 156.)

Platinum nitrate sulphocarbamide, $4\text{CS}(\text{NH}_2)_2$.

Very sol. in H_2O . Unstable. (K. J. pr. 1894, (2) 50. 490.)

Potassium nitrate, KNO_3 .

Not deliquescent, but, according to 100 pts. KNO_3 under a bell jar with up 339 pts. H_2O in 22 days, and finally deliquesce completely.

Sol. in H_2O with absorption of heat. 16 pts. $\text{KNO}_3 + 100$ pts. H_2O at 1° the temperature 10.2° . If the initial 23° it falls to 12.8° , if 0° it does not -2.7° , which is the freezing-point mixture. (Rüdorff, Pogg. 136. 276.)

$\text{KNO}_3 + \text{Aq}$ sat. at 18.1° has 1.1601 sp. gr. contains 22.72% KNO_3 , or 100 pts. H_2O at 18° 29.45 pts. KNO_3 . (Karsten, 1840.)

Sol. in 3.745 pts. H_2O at 15° . (Gerlach.)

Sol. in 3 pts. H_2O at 21° (Schiff, A. 284 solution has 1.1683 sp. gr.

Sol. in 3 pts. cold, and 0.5 pt. boiling H_2O cryo.)

$\text{KNO}_3 + \text{Aq}$ sat. at 18° has sp. gr. 1.151, or 21.63% KNO_3 , or 100 pts. H_2O dissolve 29.45 pts. KNO_3 at 18° . (Longchamp.)

Sol. in 4 pts. H_2O at 16° , and 0.25 pt. at 1° fault.)

100 pts. H_2O at 114.5° dissolve 284.61 pts. KNO_3 .

Sol. in 7 pts. cold, and 1 pt. boiling H_2O mann.)

Sol. in 6.15 pts. cold H_2O at 18.75° . (Ab. 100 pts. H_2O at 15.5° dissolve 26.6 pts.; 100 pts. (Ure's Dictionary.)

$\text{KNO}_3 + \text{Aq}$ sat. at 10° contains 33.3% KNO_3 .

$\text{KNO}_3 + \text{Aq}$ sat. in the cold contains 28% KNO_3 cryo.)

$\text{KNO}_3 + \text{Aq}$ sat. at 12.5° contains 34.8% KNO_3 frats.)

Solubility of KNO_3 in 100 pts. H_2O

t°	Pts. KNO_3	t°	Pts.
0	13.2	45.10	
5	16.7	54.72	
11.67	22.2	65.45	
17.91	29.3	79.72	
24.94	38.4	97.06	

(Gay-Lussac, A. ch. 11. 314)

Solubility of KNO₃ in 100 pts. H₂O at t°.

t°	Pts. KNO ₃
16.0	26.7
29	43.5
44.2	71.4

(Nordenskjöld, Pogg. 136. 312.)

100 pts. H₂O dissolve at:

10°	18°	27°	41°	53°
21.2	27.9	40.1	66.3	93.3

pts. KNO₃.
(Gerardin, A. ch. (4) 5. 150.)

100 pts. KNO₃+Aq sat. at 14° contain 43.4 pts. KNO₃; at 15°, 18.81 pts. KNO₃.
(v. Hauer, J. pr. 98. 177.)

100 pts. H₂O dissolve at:

4°	16.3°	68.3°
16	27.2	132.1

pts. KNO₃.
(Andreae, J. pr. (2) 29. 456.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. KNO ₃	t°	Pts. KNO ₃	t°	Pts. KNO ₃
0	13.3	39	62	78	165
1	13.8	40	64	79	168
2	14.6	41	66	80	172
3	15.5	42	68	81	175
4	16.4	43	70	82	179
5	17.1	44	72	83	182
6	17.8	45	74	84	185
7	18.5	46	76	85	189
8	19.3	47	78	86	192
9	20.2	48	81	87	196
10	21.1	49	83	88	199
11	22.0	50	86	89	203
12	23.0	51	88	90	206
13	24.0	52	91	91	210
14	25.0	53	93	92	214
15	26.0	54	96	93	218
16	27.0	55	98	94	222
17	28.1	56	101	95	226
18	29.1	57	103	96	230
19	30.2	58	106	97	234
20	31.2	59	108	98	238
21	32.3	60	111	99	243
22	33.5	61	113	100	247
23	34.7	62	116	101	252
24	36.0	63	119	102	256
25	37.3	64	121	103	261
26	38.6	65	124	104	266
27	40.0	66	127	105	272
28	41.4	67	130	106	278
29	42.9	68	133	107	284
30	44.5	69	136	108	289
31	46.0	70	139	109	295
32	48	71	142	110	301
33	50	72	146	111	307
34	52	73	149	112	313
35	54	74	152	113	319
36	56	75	155	114	326
37	58	76	159	114.1	327.4
38	60	77	162

(Mulder, Scheik. Verhandel. 1864. 89.)

100 pts. H₂O dissolve 493 pts. KNO₃ at 125°. (Tilden and Shenstone, Phil. Trans. 1884. 23.)

Rhombohedral KNO₃ is more easily soluble than the prismatic, and easily forms super-saturated solutions. (Frankenheim.)

Sat. KNO₃+Aq contains at:

139°	158°	160°	175°	180°
79.8	83.7	83.9	84.0	84.2%
190°	215°	225°	258°	283°
86.0	89.0	90.4	91.6	96.5%

KNO₃.
(Étard, A. ch. 1894, (7) 2. 526.)

Solubility in 100 pts. H₂O at t°.

t°	G. KNO ₃	Sp. gr. t°/4°
0.40	13.43	1.0817
14.90	25.78	1.1389
30.80	47.52	1.2218
44.75	74.50	1.3043
60.05	111.18	1.3903
76	156.61	1.4700
91.65	210.20	1.5394
114*	311.64	1.6269

* B.-pt. of sat. solution.

(Berkeley, Phil. Trans. 1904, 203, A. 189.)

100 g. H₂O dissolve 37.79 g. KNO₃ at 25°.

100 g. H₂O dissolve 3.08 g. equiv. KNO₃ at 20°; 3.27 at 21.5°. (Euler, Z. phys. Ch. 1904, 49. 312.)

1 l. H₂O dissolves 384.48 g. KNO₃ at 25°. (Armstrong and Eyre, Proc. Roy. Soc. 1910, A, 84. 123.)

1 l. sat. KNO₃+Aq contains 2.8 g. mols. KNO₃. (Rosenheim and Weinheber, Z. anorg. 1911, 69. 263.)

100 g. H₂O dissolve 38.485 g. KNO₃ at 25°. (Haigh, J. Am. Chem. Soc. 1912, 34. 1148.)

Sat. KNO₃+Aq contains at:

50°	58°	62°	68°
46.39	51.55	53.64	57.04%

KNO₃.
(Tschugaeff, Z. anorg. 1914, 86. 160.)

Solubility in H₂O.

100 g. of the sat. solution contain at:

9.1°	21.1°	35°
16.76	24.77	35.01

g. KNO₃.

(Findlay, Chem. Soc. 1914, 105. 780.)

Sp. gr. of solution sat. at 15°=1.134. (Michel and Krafft.)

Sp. gr. of solution sat. at 16°=1.138. (Stolba, J. pr. 97. 503.)

Sp. gr. of solution sat. at 18°=1.1601, and contains 29.45% KNO₃. (Karsten.)

Sp. gr. of KNO₃+Aq at 19.5°.

% KNO ₃	Sp. gr.	% KNO ₃	Sp. gr.
4.871	1.0307	17.965	1.1198
9.618	1.0618	21.488	1.1457
14.044	1.0920

(Kremers, Pogg. 95. 120.)

Sp. gr. of KNO ₃ +Aq at 21°.			
% KNO ₃	Sp. gr.	% KNO ₃	Sp. gr.
1	1.0058	13	1.0819
2	1.0118	14	1.0887
3	1.0178	15	1.0956
4	1.0239	16	1.1026
5	1.0300	17	1.1097
6	1.0363	18	1.1169
7	1.0425	19	1.1242
8	1.0490	20	1.1316
9	1.0555	21	1.1390
10	1.0621	22	1.1464
11	1.0686	23	1.1538
12	1.0752	24	1.1613

(Schiff, A. 110. 75.)

Sp. gr. of KNO ₃ +Aq at 15°.			
% KNO ₃	Sp. gr.	% KNO ₃	Sp. gr.
1	1.00641	12	1.07905
2	1.01283	13	1.08596
3	1.01924	14	1.09286
4	1.02566	15	1.09977
5	1.03207	16	1.10701
6	1.03870	17	1.11426
7	1.04534	18	1.12150
8	1.05197	19	1.12875
9	1.05861	20	1.13599
10	1.06524	21	1.14361
11	1.07215

(Gerlach, Z. anal. 8. 286.)

Sp. gr. of KNO ₃ +Aq at 17.5°.					
% KNO ₃	Sp. gr.	% KNO ₃	Sp. gr.	% KNO ₃	Sp. gr.
1	1.006	8	1.051	15	1.099
2	1.012	9	1.058	16	1.106
3	1.019	10	1.065	17	1.113
4	1.025	11	1.072	18	1.120
5	1.032	12	1.078	19	1.127
6	1.038	13	1.085	20	1.134
7	1.045	14	1.092

(Hager, Comm. 1883.)

Sp. gr. of KNO ₃ +Aq at 18°.			
% KNO ₃	Sp. gr.	% KNO ₃	Sp. gr.
5	1.0305	20	1.133
10	1.0632	22	1.148
15	1.097

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of KNO ₃ +Aq at 20°, containing mols. KNO ₃ in 100 mols. H ₂ O.			
Mols. KNO ₃	Sp. gr.	Mols. KNO ₃	Sp. gr.
0.5	1.01730	4	1.12264
1	1.03373	5	1.14888
2	1.06524

(Nicol, Phil. Mag. (5) 16. 122.)

Sp. gr. of KNO ₃ +Aq at 25°.	
Concentration of KNO ₃ +Aq.	Sp. gr.
1-normal	1.0605
1/2 " "	1.0305
1/3 " "	1.0161
1/4 " "	1.0075

(Wagner, Z. phys. Ch. 1890, 5. 37)

Sp. gr. of KNO₃+Aq at 20.1°.
p=per cent strength of solution;
served density; w=volume conc. in
per cc. ($\frac{pd}{100}=w$.)

p	d	w
25.54	1.1783	0.30
21.95	1.1510	0.25
17.88	1.1200	0.20
13.93	1.0913	0.15
8.706	1.0553	0.09
5.393	1.0331	0.05
4.389	1.0264	0.04
2.848	1.0165	0.02
2.030	1.0113	0.02
0.741	1.0030	0.00

(Barnes, J. Phys. Chem. 1898, 2. 5)

Sp. gr. 20°/4° of a normal solution of
= 1.05954; of a 0.5 normal solution = 1.0
(Haigh, J. Am. Chem. Soc. 1912, 34. 1)

Sp. gr. of sat. KNO ₃ +Aq at t°.		
t°	G. KNO ₃ sol. in 100 g. H ₂ O	Sp.
0	13.27	1.0
10	20.89	1.1
20	31.59	1.2
30	45.85	1.3
40	63.90	1.4
50	85.51	1.5
60	109.00	1.6
70	138.00	1.7

(Tschernaj, J. Russ. Phys. Chem. Soc. 44. 1565.)

The saturated solution boils at 114.1 der); 114.5° (Griffiths); 115.9° (Legrardin); 117° (Magnus); 118° (Kremer (Le Page).
The saturated solution forms a e 111°, and boils at 115°; highest ten served, 115.3°. (Gerlach, Z. anal. 28.

pt of KNO₃+Aq containing pts. KNO₃ to 100 pts. H₂O. G=according to Gerlach (Z. anal. 26. 444); L=according to Legrand (A. ch. (2) 52. 426).

n.	G	L	B.-pt.	G	L
5°	7.5	...	107°	120.5	119.0
	15.2	12.2	108	141.5	140.6
5	23	...	109	164	163.0
	31	26.4	110	188.5	185.9
5	39	...	111	215	209.2
	47.5	42.2	112	243	233.0
5	56	...	113	274	257.6
	64.5	59.6	114	306	283.3
5	73	...	115	338.5	310.2
	82	78.3	115.9	...	335.1
	101	98.2

pt. KNO₃ dissolves in 1.4 pts. HNO₃; at n 3.8 pts., and at 123° in 1 pt. HNO₃+Aq 423 sp. gr. (Composition 2HNO₃, 3H₂O.) ultz, Zeit. Ch. (2) 5. 531.)

Solubility of KNO₃ in HNO₃ at 0°.

G. per 100 cc. solution		Sp. gr.
KNO ₃	HNO ₃	
12.65	0.00	1.079
10.02	3.71	...
8.38	8.38	1.093
7.49	13.58	1.117
7.49	19.47	1.144
7.68	30.04	1.202
10.42	42.86	1.289
28.64	75.95	1.498

(Engel, C. R., 1887, 104. 913.)

Solubility in HNO₃+Aq.

ation np.	% by wt. KNO ₃	Solid phase
	24.4	KNO ₃ , 2HNO ₃ (solution in HNO ₃) (stable)
0	32.6	"
0	34.8	"
5	37.2	"
0	44.5	(mpt.)
5	47.8	(Solution in KNO ₃) (labile)
5	48.6	"
0	50.9	"
0	37.2	KNO ₃ , HNO ₃ (labile)
5	44.5	"
5	47.2	(stable)
5	47.8	"
5	48.6	"
0	49.4	"
0	50.1	(labile)
5	50.9	"
0	49.4	KNO ₃ (labile)
0	50.9	(stable)
0	51.7	"

(Groschuff, Z. anorg. 1904, 40. 10.)

Sol. in sat. NH₄Cl+Aq. Solution thus obtained contains 43.07 pts. mixed salts, or 100 pts. H₂O dissolve 75.66 pts. mixed salts, viz. 38.62 pts. KNO₃ and 39.84 pts. NH₄Cl. (Karsten.) See also under NH₄Cl.

Solubility of KNO₃+K₂CO₃ and KHCO₃ in H₂O, see under K₂CO₃ and KHCO₃.

Sol. in sat. BaCl₂+Aq with pptn. of Ba(NO₃)₂.

Sol. in sat. NH₄NO₃+Aq, at first without pptn., but afterwards NH₄NO₃ is pptd. (Karsten.)

Sol. in NH₄NO₃+Aq with pptn. of NH₄NO₃. (Rüdorff, B. 6. 485.)

See also under NH₄NO₃.

Sol. in sat. Ba(NO₃)₂+Aq, but soon a double salt separates. (Karsten.)

See also under Ba(NO₃)₂.

Sol. in Ca(NO₃)₂+Aq. (Longchamp.)

See also under Ca(NO₃)₂.

Sol. in sat. Pb(NO₃)₂+Aq without pptn.

100 pts. H₂O dissolve 119.6 pts. Pb(NO₃)₂ and 67.1 pts. KNO₃ at 21.2°. (Rüdorff, B. 6. 484.) See also under Pb(NO₃)₂.

Solubility in AgNO₃+Aq at t°.

t°	Sat. solution contains		
	% KNO ₃	% AgNO ₃	% total salt
-7	10.5	39.4	49.9
-7.5	10.5	40.5	51.0
-4	11.3	42.5	53.8
+20	23.6	46.4	70.0
33	26.3	50.9	77.2
34	27.3	51.1	78.4
36	29.4	52.0	81.4
38	81.7
46	82.3
54	33.1	55.0	88.1
54	...	55.8	...
61	89.5
68	34.3	54.6	88.9
96	37.8	55.4	93.2
105	38.5	55.6	94.1
142	41.5	55.8	97.3

(Étard, A. ch. 1894, (7) 3. 286.)

Solubility of mixed crystals of KNO₃ and AgNO₃ in H₂O at 25°.

G. per l.		Mg. mols. per l.		Mol. % AgNO ₃ in solu- tion	Mol. % AgNO ₃ in solid phase
AgNO ₃	KNO ₃	AgNO ₃	KNO ₃		
45.9	321.8	270	3180	7.83	0.2896
110.7	322.6	6513	3184	16.96	0.6006
176.8	333.7	1040	3298	23.97	0.9040
259.6	364.0	1258	3597	29.81	1.054
365.6	456.4	2151	4511	32.28	1.604
507.9	387.2	2988	3816	43.85	2.439
745.9	398.6	4388	3960	57.70	8.294

(Herz, Z. Kryst. Min. 1897, 28. 405.)

Solubility of $\text{KNO}_3 + \text{AgNO}_3$ at 30° .

Composition of the solution		Solid phase
% KNO_3	% AgNO_3	
0	73.0	AgNO_3
5.53	71.65	"
11.22	69.01	$\text{AgNO}_3 + \text{AgNO}_3, \text{KNO}_3$
13.44	65.08	$\text{AgNO}_3, \text{KNO}_3$
17.38	57.85	"
25.05	46.32	$\text{KNO}_3 + \text{AgNO}_3, \text{KNO}_3$
25.00	46.46	"
26.58	39.09	KNO_3
29.22	23.59	"
30.45	11.51	"
31.30	0	"

(Schreinemakers, Z. phys. Ch. 1909, 65. 576.)

 $\text{KNO}_3 + \text{NaNO}_3$.

100 pts. H_2O dissolve 34.53 pts. KNO_3 and 91.16 pts. NaNO_3 at 15.6° , and solution has sp. gr. = 1.478. (Page and Keightley.)

100 pts. $\text{KNO}_3 + \text{NaNO}_3 + \text{Aq}$ sat. at 14° contain 52.17 pts. of the two salts; sat. at 13° contain 53.15 pts. of the two salts. (v. Hauer.)

100 pts. H_2O dissolve at 18.75° 29.45 pts. KNO_3 and 89.53 pts. NaNO_3 , if sat. $\text{KNO}_3 + \text{Aq}$ is treated with NaNO_3 , and 35.79 pts. KNO_3 and 88.00 pts. NaNO_3 by the opposite process. 134.38 pts. of the two salts are dissolved if a mixture of the salts is treated with H_2O at 18.75° . (Karsten.)

100 pts. H_2O dissolve 39.34 pts. KNO_3 and 94.60 pts. NaNO_3 , or 133.94 pts. of the two salts at 20° . (Nicol, Phil. Mag. (5) 13. 385.)

Solubility of mixtures of KNO_3 and NaNO_3 .

% NaNO_3 in mixture before solution	Total amt mixed salts dissolved in 100 pts. H_2O at 20°	Pts. NaNO_3 dissolved	Pts. KNO_3 dissolved	% NaNO_3 in mixture after solution and evap. to dryness
100	86.8	86.8	0	100
90	109.6	96.4	13.2	88
80	136.5	98.0	38.5	71.8
70	136.3	.	.	.
60	137.6	90.0	47.6	65.4
50	106.1	66.0	40.1	62.2
45.7*	88.0	53.3	34.7	60.6
40	81.1	45.6	35.6	56.2
30	73.5	.	.	.
20	54.1	20.8	33.3	38.5
10	40.9	9.4	31.5	22.9
0	33.6	0	33.6	0

* $\text{NaNO}_3 + \text{KNO}_3$.

(Carnelley and Thomson, Chem. Soc. 53. 782.)

Solubility of $\text{KNO}_3 + \text{NaNO}_3$ in H_2O at

In 1000 ccm. H_2O		Solid phase
NaNO_3	KNO_3	
805.0	.	NaNO_3
848.3	301.9	$\text{NaNO}_3, \text{KNO}_3$
.	208.9	KNO_3

(Kremann and Zitek, M. 1909, 30. 3)

Solubility of $\text{KNO}_3 + \text{NaNO}_3$ in H_2O at

In 1000 ccm. H_2O		Solid phase
NaNO_3	KNO_3	
913.58	.	NaNO_3
910.60	123.60	"
1019.65	435.85	$\text{NaNO}_3, \text{KNO}_3$
1018.40	437.70	"
931.30	422.00	KNO_3
346.70	390.00	"
.	377.35	"

(Kremann and Zitek.)

Solubility of $\text{KNO}_3 + \text{NaNO}_3$ in H_2O

% KNO_3	% NaNO_3	Solid phase
38.70	39.62	KNO_3
41.60	66.31	"
46.35	100.10	$\text{KNO}_3 + \text{NaNO}_3$
39.08	98.99	NaNO_3
20.98	94.44	"

(Uyeda, Mem. Col. Sc. Kyoto, 1910, 1)

Solubility in $\text{NaNO}_3 + \text{Aq}$ at 20° , 15.6° , and 91° . Data, given in the original, that each salt increases the solubility of the other. (Leather, Mem. Dept. Agric. 1914, 3. 177; Chem. Soc. 1915, 100. (2))

100 pts. H_2O dissolve 28.92 pts. KNO_3 , 53.68 pts. NaNO_3 , and 26.44 pts. NaNO_3 at 15.6° , and solution has sp. gr. = 1.44. (Page and Keightley, Chem. Soc. (2) 10. 56)

Solubility of $\text{KNO}_3 + \text{NaNO}_3 + \text{NaCl}$ at 25° .

% KNO_3	% NaNO_3	% NaCl	Solid phase
38.44	22.87	32.58	$\text{NaCl} + \text{KNO}_3$
38.57	44.40	27.67	"
42.55	63.26	23.59	$\text{NaNO}_3, \text{KNO}_3$
17.77	61.12	23.94	"
28.08	62.92	23.70	"
44.72	82.82	9.56	"

(Uyeda, Mem. Col. Sc. Kyoto, 1910, 1)

 $\text{KNO}_3 + \text{Sr}(\text{NO}_3)_2$.

1 l. H_2O sat. with both salts at 25° contains 552 g. KNO_3 + 1074 g. $\text{Sr}(\text{NO}_3)_2$. (Leather and Noyes, Z. phys. Ch. 1890, 8. 300.)

Solubility of $\text{KNO}_3 + \text{Sr}(\text{NO}_3)_2$ at t° .

$\%$ $\text{Sr}(\text{NO}_3)_2$	Solid phase
5.49	KNO_3
9.17	"
17.10	"
31.24	"
34.91	"
39.56	$\text{KNO}_3 + \text{Sr}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
40.37	$\text{Sr}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
41.12	"
40.70	"
23.70	KNO_3
38.52	$\text{KNO}_3 + \text{Sr}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
40.22	$\text{Sr}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$
44.19	"
47.7	"

Morgan and Morris, Chem. Soc. 1914, 105. 779.)

TiNO_3 .
 H_2O dissolve 43.5 g. $\text{TiNO}_3 + 104.2$ g. H_2O at 58° . (Rabe, Z. anorg. 1902, 31.

Solubility of mixed crystals of $\text{KNO}_3 + \text{TiNO}_3$ in H_2O at 25° .

G. per l.	Sp. gr.
KNO_3	
351.0	1.2632
329.0	1.1903
332.4	1.1956
333.7	1.2050
333.3	1.2196
321.0	1.2436
330.5	1.2617
428.3	1.2950
245.1	1.2050
0.0	1.0964

Z. Kryst. Min. 1897, 28. 405.)

KBr .

Solubility in $\text{KBr} + \text{Aq}$.

litre of the solution contains

4.5°	at 25.2°	
Mol. KNO_3	Mol. KBr	Mol. KNO_3
2.228	0.0	3.217
2.026	0.38	3.026
1.835	0.93	2.689
1.730	1.37	2.492
1.589	2.08	2.216
1.406	2.87	1.958
1.308	3.55	1.807

ren, C. R., 1900, 130. 910.)

$\text{KNO}_3 + \text{KCl}$.

100 pts. H_2O dissolve pts. of the two salts:

	At 12.9°	At 15.3°
KNO_3	18.8	18.9
KCl	28.5	29.8

(Kopp.)

100 pts. H_2O dissolve 315.2 pts. KCl and 19.1 pts. KNO_3 at 20.0° . (Rüdorff, B. 6. 484.)
100 pts. H_2O dissolve 18.95 pts. $\text{KNO}_3 + 32.84$ pts. KCl , or 51.79 pts. of the mixed salts at 20° . (Nicol, Phil. Mag. (5) 31. 385.)

Solubility of KCl with addition of KNO_3 at 17.5° .

Sp. gr.	100 ccm. of solution contain g.		
	KCl	H_2O	KNO_3
1.1730	29.39	87.85	0
1.1980	27.50	85.68	6.58
1.2100	27.34	84.76	8.83
1.2250	26.53	83.58	12.48
1.2360	25.98	82.84	14.83
1.2390	25.96	82.65	15.22
1.2388	25.95	82.43	15.49
1.2410	26.24	82.63	15.33

KNO_3 separated out in last four solutions.

Solubility of KNO_3 with addition of KCl at 20.5° .

Sp. gr.	100 ccm. of solution contain g.		
	KNO_3	H_2O	KCl
1.1625	27.68	88.51	0
1.1700	24.39	87.89	4.72
1.1765	22.44	87.47	7.74
1.1895	20.23	86.48	12.23
1.1983	18.96	85.69	15.15
1.2150	17.67	84.23	19.61
1.2265	17.11	83.40	22.17
1.2400	16.79	82.24	24.96

(Bodländer, Z. phys. Ch. 7. 359.)

Solubility in KCl+Aq at t°.				Solubility of KCl in KNO ₃ +Aq			
t°	100 ml. solution contains			t°	Concentration of KNO ₃ % mol per l.	G. salt dissolved in 1 l. H ₂ O	M. sol.
	% KNO ₃	% KCl	% total salt				
-11.4	4.0	18.4	22.4	0°	0	283.55	...
-11	3.9	18.1	22.0	"	1/4	284.25	...
-10	3.8	18.6	22.5	"	1/2	283.60	...
-4	...	19.7	...	"	1	287.60	...
+2.5	6.3	19.9	26.2	25°	0	364.15	...
4.5	7.3	20.7	28.0	"	1/4	365.00	...
4.5	7.8	19.8	27.6	"	1/2	361.65	...
8.5	7.9	20.9	28.8	"	1	358.80	...
10.5	8.4	21.0	29.4	"	1 1/2	355.20	...
13.5	8.9	21.8	30.7				
14	10.2	21.3	31.5				
17	9.7	22.6	32.3				
23	12.5	21.8	34.3				
27	14.9	21.3	36.2				
29	16.1	21.0	37.1				
34	18.9	21.2	39.3				
36	18.9	21.6	40.5				
37.5	19.2	21.6	40.8				
42	21.0	21.0	42.0				
42.5	21.8	20.9	42.7				
48	25.3	20.3	45.6				
50	28.3	20.8	49.1				
53	...	20.2	...				
53	48.7				
56	29.5	19.5	49.0				
61	34.5	18.3	52.8				
62	35.4				
71	40.5	17.3	57.8				
81	47.1	15.4	62.5				
85	48.2	15.3	63.5				
90	52.8	13.3	66.1				
96	54.1	12.6	66.7				
97	56.6	12.4	69.0				
104	59.9	10.8	70.7				
105	...	10.9	...				
120	69.6	7.7	77.3				
121	69.1	7.6	76.7				

(Étard, A. ch. 1894, (7) 3. 285.)

Solubility in KCl+Aq.

1 litre of the solution contains at			
14.5°		at 25.2°	
Mol. KCl	Mol. KNO ₃	Mol. KCl	Mol. KNO ₃
0.0	2.228	0.0	3.217
0.182	2.172	0.26	3.066
0.424	2.057	0.66	2.853
0.880	1.830	1.35	2.510
1.778	1.576	2.06	2.218
2.204	1.515	2.78	2.015
2.635	1.423	3.04	1.946
3.172	1.355	.	.

(Touren, C. R. 1900, 130. 909.)

(Armstrong and Eyre, Proc. R. Soc. 12. 84. 127.)

Solubility in KCl+Aq at 20°, 30°, 91°. Data, given in the original, show each salt diminishes the solubility in the other. (Leather, Mem. Dept. Agri. 1914, 3. 177; Chem. Soc. 1915, 106. (KNO₃+NaCl.

NaCl is sol. in sat. KNO₃+Aq, mixed solution is capable of dissolving KNO₃. An amount of H₂O, which pure, could only dissolve 100 pts. KCl in this way be made to take up 15% (Longchamp, A. ch. (2) 9. 8.)

Sol. in sat. NaCl+Aq.

100 pts. H₂O dissolve:

	Longchamp 4° (1)	Rüdorff	
		14° (2)	18° (3)
NaCl	35.96	38.5	38.9
KNO ₃	26.01	28.7	36.1
	61.97	67.2	75.0

	Karsten 18.75°		
	(5)	(6)	(7)
NaCl	36.53	38.25	39.19
KNO ₃	33.12	29.45	28.12
	69.65	67.70	77.72

1, 2, 3, 4, and 8. Both salts in eq. 5. Sat. NaCl+Aq treated with K. 6. Sat. KNO₃+Aq treated with N. 7. The two salts simultaneously with H₂O.

100 pts. H₂O dissolve 31.44 pts. KCl, and 38.58 pts. NaCl at 1 solution has sp. gr. = 1.33. (P. Keightley.)

solubility in NaCl+Aq at t°.

	Sat. solution contains		
	% KNO ₃	% NaCl	% total salt
	7.3	22.5	29.8
5	7.9	22.6	30.5
5	8.7	22.1	31.8
	10.1	22.5	32.6
	10.9	23.0	33.9
	12.7	23.3	36.0
	12.9	23.8	36.7
	16.6	22.8	39.4
	19.0	22.8	41.8
	19.8	22.0	41.8
	18.9	22.5	41.5
5	20.4	22.3	42.7
	...	21.2	...
	21.8	20.7	42.5
	20.7.	22.3	43.0
	45.0
5	24.7	20.7	45.4
5	25.0	20.2	45.2
5	25.9	20.2	46.1
	26.7	20.7	47.4
	27.9	20.2	48.1
	29.8	19.5	49.3
2	31.1	20.0	51.1
	38.5	16.5	55.0
	39.4	17.1	56.5
5	40.9	15.3	56.2
	49.7	14.0	63.7
	53.9	13.6	67.5
	54.8	12.9	67.7
	57.4	12.6	70.0
	61.4	10.4	71.8
	64.7	9.5	74.2
	70.0	9.0	79.0
	69.9	9.3	79.2
	71.3	8.4	79.7
	72.2	9.0	81.2
	73.8	8.0	81.8
	73.6	7.9	81.5
	72.9	8.8	81.7
	73.0	7.6	80.6
	74.2	7.9	82.1
	75.7	7.6	83.3
	77.7	7.6	85.3
	80.7	5.8	86.5
	79.1	5.9	85.0

Stard, A. ch. 1894, (7) 3. 283.)

H₂O dissolve 41.14 g. KNO₃ and NaCl at 25°; 168.8 g. KNO₃ and NaCl at 80°. (Soch, J. phys. Ch. 46.)
insol under NaCl.
insol in sat. CuSO₄+Aq, forming a double salt which soon separates out.
slowly and slightly sol. in MgSO₄+Aq with pptn. of MgSO₄. (Karsten.)
insol in sat. K₂SO₄+Aq.
insol in NO₂+Aq dissolves some K₂SO₄, and insol in O₂+Aq slowly dissolves some KNO₃.

without pptn., but K₂SO₄ is afterwards pptd. (Karsten.)

100 pts. H₂O dissolve:

	Mulder 18.75° (1)	Karsten 18.75° (2)	Kopp		Mulder 18.75° (5)
			20° (3)	40° (4)	
KNO ₃	29.90	29.42	26.9	59.35	...
K ₂ SO ₄	...	4.0	6.6	5.75	10.8

2. H₂O sat. with KNO₃ and K₂SO₄ simultaneously, or to a sat. solution of one salt the other was added.

3 and 4. H₂O sat. with both salts simultaneously.

Mulder doubts the results of 3 and 4.

Solubility in K₂SO₄+Aq at t°.

t°	In 100 ccm. of the solution		Sp. gr. of solution
	G. KNO ₃	G. K ₂ SO ₄	
15	216.5	50.7	1.165
25	308.5	47.66	1.210

(Euler, Z. phys. Ch. 1914, 40. 313.)

Slowly sol. in sat. Na₂SO₄ at first without pptn., but afterwards K₂SO₄ or NaSO₄ separates out.

Sol. in sat. ZnSO₄+Aq with pptn. of double salt. (Karsten.)

Sol. in sat. KClO₃+Aq, from which solution it is not pptd. by salts which would ppt. it from aqueous solution. (Karsten.)

Hydrazine dissolves 21.7 pts. KNO₃ at 12.5-13°. (de Bruyn, R. t. c. 1899, 18. 297.)

Neither dissolved nor attacked by liquid NO₂. (Frankland, Chem. Soc. 1901, 79. 1361.)

Very sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in absolute alcohol; in dilute alcohol it dissolves proportional to the amount of H₂O present, but always less is dissolved than the H₂O would dissolve by itself. (Gerardin.)

100 pts. alcohol containing % by weight of alcohol dissolve pts. KNO₃ at 15°.

10	20	30	40	50	60	80%
13.2	8.5	5.6	4.3	2.8	1.7	0.4 pts. KNO ₃ .

(Schiff, A. 118. 365.)

Solubility in 100 pts. alcohol at t° . D = sp. gr. of alcohol; S = solubility.

D = 0.9904		D = 0.9848		D = 0.9793		D = 0.9726	
t°	S	t°	S	t°	S	t°	S
12	18.1	12	14.6	10	10.20	14	8.8
21	25.0	21	21.7	10	10.19	25	13.6
33	40.4	36	37.8	13	11.74	34	20.3
43	58.6	41	45.0	18	14.52	44	31.3
53	79.1	56	72.9	20	16.35	47	34.2
61	94.5			31	25.81	60	52.3
62	95.7			34	28.63		
				40	36.66		
				41	37.20		
				50	50.14		
				53	56.01		
				61	72.24		
				62	73.36		

D = 0.9573		D = 0.9390		D = 0.8967		D = 0.8429	
t°	S	t°	S	t°	S	t°	S
14	5.4	16	4.13	12	1.61	15	0.29
25	9.0	24	6.00	33	3.62	22	0.39
33	13.2	40	10.94	47	5.77	40	0.62
	19.1	51	16.51	57	6.97	54	0.78
57	29.1	60	21.54			60	1.10
65	36.2	64	24.22				

(Gerardin, A. ch. (4) 5. 151.)

Solubility of KNO_3 in alcohol at 18° .

Sp. gr.	100 cem contain g		
	Alcohol	Water	KNO_3
1.1475		89.63	25.12
1.1085	3.30	87.44	20.11
1.1010	5.24	86.26	18.60
1.0805	8.69	83.18	16.18
1.0655	14.08	77.93	14.54
1.0490	16.27	76.36	12.27
1.0375	19.97	72.93	10.85
0.9935	28.11	64.74	6.50
0.9585	37.53	54.21	4.11
0.9456	42.98	48.15	3.37
0.9050	51.23	27.32	1.95
0.8722	61.65	24.74	0.83
0.8375	69.60	13.95	0.20

(Bodländer, Z. phys. Ch. 7. 316.)

Solubility in alcohol.

Wt. % alcohol	G. KNO_3 per 100 g. alcohol	
	at 30°	at 40°
0	45.6	64.5
8.25	32.3	47.1
17.0	22.4	33.3
25.7	15.1	24.1
35.0	11.4 (34.5°)	16.7
44.9	7.0	11.6 (44)
54.3	4.5	7.2 (35)
65.0	2.7	4.4
75.6	1.3	2.0 (76)
88.0	0.4	0.6 (96)

(Bathrick, J. phys. Ch. 1896, 1. 100)

Solubility of KNO_3 in ethyl alcohol + 30° .

% by wt. H_2O	% by wt. alcohol	% by wt. l
68.7	0	31.3
69.2	10.1	20.7
67.3	17.0	16.7
64.1	23.8	12.1
58.8	32.2	9.0
50.8	43.1	6.1
39.8	56.9	3.3
33.9	63.8	2.3
22.3	76.8	0.8
7.5	92.3	0.1

(Schreinemakers, Z. phys. Ch. 1909, 66)

Solubility in ethyl alcohol at 25° .

Concentration of alcohol in g. mol per l. H_2O	Solubility in 1 l. H_2O	M. solub
0	384.48	3.
$\frac{1}{4}$	368.30	3.
$\frac{1}{2}$	354.40	3.
1	327.00	3.

(Armstrong and Eyre, Proc. R. Soc. (A) 84. 127.)

Solubility of KNO_3 in methyl alcohol + 30° .

% by wt. H_2O	% by wt. alcohol	% by wt
68.7	0	31.
68.9	7.8	28.
66.4	17.3	16.
61.0	27.8	11.
53.9	38.4	7.
39.2	57.0	3.
0.99	98.58	0.

(Schreinemakers, Z. phys. Ch. 1909, 66)

Solubility of $\text{KNO}_3 + \text{AgNO}_3$ in 51.6% $\text{C}_2\text{H}_5\text{OH} + \text{Aq}$ at 30° .

KNO_3	% AgNO_3	Solid phase
3	0	KNO_3
55	5.15	"
11	16.47	"
26	21.28	$\text{KNO}_3 + \text{AgNO}_3, \text{KNO}_3$
32	36.94	$\text{AgNO}_3, \text{KNO}_3 + \text{AgNO}_3$
	37	AgNO_3

Barinmakers, Z. phys. Ch. 1909, 65. 556.)

100 g. 40% ethyl alcohol sat. with $\text{KNO}_3 + \text{Cl}$ at 25° contain 13.74 g. $\text{KNO}_3 + 15.78$ g. Cl . (Soch, J. phys. Ch. 1898, 2. 43.)
Insol. in propyl alcohol. (Schlamp, Z. phys. Ch. 1894, 14. 277.)
Almost insol. in ether. (Braconnot.)
Very sl. sol. in acetone. (Krug and Elroy.)
Sol. in acetone. (Eidmann, C. C. 1899. II, 4.)

Solubility in acetone + Aq at 40° .

Wt. % acetone	G. KNO_3 per 100 g. solvent
0	64.5
8.5	51.3
16.8	38.9
25.2	22.8
34.3	24.7
44.1	17.0
53.9	11.9
64.8	7.2
76.0	3.0
87.6	0.7

(Bathrick, J. phys. Ch. 1896, 1. 160.)

100 pts. glycerine (sp. gr. 1.225) dissolve 10 g. KNO_3 . (Vogel, N. Rep. Ph. 16. 557.)
100 g. trichlorethylene dissolve 0.01 g. NO_3 at 15° . (Wester and Bruins, Pharm. weekbl. 1914, 51. 1443.)
Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 257.)
Insol. in benzonitrile. (Naumann, B. 1914, 1370.)
Insol. in methyl acetate (Naumann, B. 19, 42. 3790); ethyl acetate. (Naumann, 1910, 43. 314.)
100 g. H_2O sat. with sugar and KNO_3 dissolve 224.7 g. sugar + 41.9 g. KNO_3 , or saturation contains 61.36 g. sugar + 11.45 g. NO_3 at 31.25° . (Köhler, Z. Ver. Zuckerind. 7, 47. 447.)

Potassium hydrogen nitrate, $\text{KNO}_3, \text{HNO}_3$.
Very hygroscopic. Decomp. by H_2O . (Groschuff, B. 1904, 37. 1489.)

Potassium dihydrogen nitrate, $\text{KNO}_3, 2\text{HNO}_3$.
Decomp. by H_2O . (Ditte, A. ch. (5) 18. 1.)

Solubility in H_2O .

Solution temp.	% by wt. KNO_3	% by wt. HNO_3	% by wt. H_2O
mpt. $+22^\circ$	44.5	55.5	0
20.5	44.1	55.0	0.9
18.0	43.8	54.5	1.7
12.0	43.0	53.6	3.4
6.0	42.3	52.7	5.0
0	41.6	51.8	6.6

(Groschuff, Z. anorg. 1904, 40. 11.)

Potassium silver nitrate, $\text{KNO}_3, \text{AgNO}_3$.
Sol. in H_2O . (Russell and Maskelyne, Roy. Soc. Proc. 26. 357.)
 $3\text{KNO}_3, \text{AgNO}_3$. Sol. in H_2O . (Rose, Pogg. 106. 320.)

Potassium thallic nitrate, $2\text{KNO}_3, \text{Ti}(\text{NO}_3)_3 + \text{H}_2\text{O}$.
Decomp. by H_2O . (Meyer, Z. anorg. 1900, 24. 361.)

Potassium thorium nitrate, $4\text{KNO}_3, \text{Th}(\text{NO}_3)_4$.
Very sol. in H_2O and alcohol. (Berzelius.)
Hygroscopic; very unstable. (Meyer, Z. anorg. 1901, 27. 379.)
Hydroscopic; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Meyer, Z. anorg. 1901, 27. 378.)

Potassium thorium hydrogen nitrate, $3\text{KNO}_3, \text{Th}(\text{NO}_3)_4, 3\text{HNO}_3$.
Decomp. in the air. (Meyer, B. 1900, 33. 2140.)
+ $4\text{H}_2\text{O}$. Sol. in HNO_3 of 1.2 sp. gr. Effloresces in the air. (Meyer, Z. anorg. 1901, 27. 380.)

Potassium uranyl nitrate, $\text{K}(\text{UO}_2)(\text{NO}_3)_3$.
Decomp. by H_2O . Sol. in conc. HNO_3 . (Meyer, B. 1903, 36. 4057.)

Solubility in H_2O at t° .

t°	In 100 pts. of the solution			Solid phase
	Pts. by wt. UO_2	Pts. by wt. K	Pts. by wt. NO_3	
0.5	31.98	1.72	...	Double salt + KNO_3
13.0	33.40	2.74	...	"
25.0 a)	37.08	4.05	23.49	"
b)	37.06	3.98	23.46	"
45.0	42.18	5.16	...	"
59.0	41.65	6.03	...	"
80.6 a)	43.72	6.42	...	Double salt
b)	43.70	6.34	...	"

Potassium uranyl nitrate is decomp. by H_2O at temp. below 60° ; above 60° it is sol. in H_2O without decomp.
(Rimbach, B. 1904, 37. 473.)

Potassium nitrate barium sulphate, KNO_3 , 2BaSO_4 .

Easily decomp. Sol. in conc. H_2SO_4 . (Silberberger, M. 1904, 25. 251.)

Potassium nitrate phosphomolybdate.

See Phosphomolybdate nitrate, potassium.

Potassium nitrate sulphate, KNO_3 , KHSO_4 .

Decomp. by H_2O and alcohol. (Jacquelin.)

Potassium nitrate sulphotungstate, 2KNO_3 , K_2WS_4 (?).

Very sol. in hot or cold H_2O . Insol. in alcohol. (Berzelius.)

Potassium nitrate tungstate (?).

100 pts. boiling H_2O dissolve 5 pts. salt. (Storer's Dict., p. 393.)

Potassium nitrate zinc iodide.

Permanent. Easily sol. in H_2O . Insol. in alcohol. (Anthon.)

Praseodymium nitrate, $\text{Pr}(\text{NO}_3)_3 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (von Schule, Z. anorg. 1898, 18. 355.)

Praseodymium rubidium nitrate,

$[\text{Pr}(\text{NO}_3)_3]\text{Rb}_2 + 4\text{H}_2\text{O}$.

Hygroscopic. (Jantsch, Z. anorg. 1911, 69. 230.)

Praseodymium sodium nitrate, $\text{Pr}(\text{NO}_3)_3$, $2\text{NaNO}_3 + \text{H}_2\text{O}$.

Sol. in H_2O . (von Schule, Z. anorg. 1898, 18. 356.)

Praseodymium zinc nitrate, $2\text{Pr}(\text{NO}_3)_3$, $3\text{Zn}(\text{NO}_3)_2 + 24\text{H}_2\text{O}$.

1 l. sat. solution in $\text{HNO}_3 + \text{Aq}$ (sp. gr. 1.325) contains 14.69 g. hydrous salt at 16° . (Jantsch, Z. anorg. 1912, 76. 321.)

Radium nitrate.

Has apparently the same solubility in H_2O as the corresponding Ba comp. (Curie, Dissert. 1903.)

Rhodium nitrate, $\text{Rh}(\text{NO}_3)_3 + 2\text{H}_2\text{O}$ (?).

Deliquescent. Sol. in H_2O . Insol. in alcohol. (Claus.)

Rhodium uranyl nitrate,

$2(\text{UO}_2)(\text{NO}_3)_2 \cdot \text{Rh}_2(\text{NO}_3)_6 + 10\text{H}_2\text{O}$.

Sol. in H_2O and acids; insol. in aq. alkalies. (Lancien, C. C. 1912, I. 208.)

Rubidium nitrate, RbNO_3 .

100 pts. H_2O dissolve 20.1 pts. at 0° ; 43.5 pts. at 10° . (Bunsen.)

Solubility in H_2O at t° .

t°	G. RbNO_3 per 100 g.		t°	G. R per H_2O
	H_2O	Solu- tion		
0	19.5	16.3	60	200
10	33.0	24.8	70	251
20	53.3	34.6	80	309
30	81.3	44.8	90	375
40	116.7	53.9	100	452
50	155.6	60.9	118.3	617

(Berkeley, Trans. Roy. Soc. 1904, 207.)

100 g. H_2O dissolve 66.855 g. Rb at 25° . (Haigh, J. Am. Chem. Soc. 1914, 36. 1148.)

Sp. gr. $20^\circ/4^\circ$ of a normal solution = 1.100835; of a 0.5 normal solution = 1.00835. (Haigh, J. Am. Chem. Soc. 1912, 34. 24.)

Sp. gr. of $\text{RbNO}_3 + \text{Aq}$.

G.-equiv. RbNO_3 per l.

at 18° = 0.5035

Sp. gr. at $6^\circ/6^\circ$ = 1.05342

Sp. gr. at $18^\circ/18^\circ$ = 1.05226

Sp. gr. at $30^\circ/30^\circ$ = 1.05154

G.-equiv. RbNO_3 per l.

at 18° = 2.000

Sp. gr. at $6^\circ/6^\circ$ = 1.20654

Sp. gr. at $18^\circ/18^\circ$ = 1.20302

Sp. gr. at $30^\circ/30^\circ$ = 1.20034

(Clausen, W. Ann. 1914, (4) 44.)

Easily sol. in HNO_3 . (Schultz, (2) 5. 531.)

Sol. in acetone. (Eidmann, C. C. 1014; Naumann, B. 1904, 37. 4328.)

Insol. in methyl acetate. (Nau, 1909, 42. 3790.)

Rubidium hydrogen nitrate, RbNO_3 .

Fairly stable in air. (Wells, Ar 1901, 26. 273.)

Rubidium dihydrogen nitrate, RbNH_4NO_3 .

Decomp. rapidly in air. (Wells, Ar 1901, 26. 273.)

$2\text{RbNO}_3 \cdot 5\text{HNO}_3$. Decomp. Known only in solution in HCl . (Ditte, A. ch. (5) 18. 320.)

Rubidium silver nitrate, $\text{RbNO}_3 \cdot \text{AgNO}_3$.

Sol. in H_2O . (Russell and Maskell, Soc. Proc. 26. 357.)

Rubidium thorium nitrate, $\text{Rb}_2\text{Th}(\text{NO}_3)_6$.

Sl. sol. in HNO_3 ; decomp. by H_2O . (Z. anorg. 1901, 27. 384.)

yl nitrate, $\text{Rb}(\text{UO}_2)(\text{NO}_3)_3$.
 H_2O . Sol. in conc. HNO_3 .
 13, 36. 4057.)

ibility of H_2O at t° .

pts. of the solution			Solid phase
Pts. wt. NO_3	Pts. by wt. Rb	Pts. by wt. total salt	
9.72	4.63	59.57	Double salt + RbNO_3
9.76	4.67	59.64	"
...	11.01	69.46	Double salt
...	11.01	69.52	"

iranyl nitrate is decomp. by
 mp.; at 80° it is sol. in H_2O
 ip.

ach, B. 1904, 37. 476.)

ate, $\text{Sm}(\text{NO}_3)_3 + 6\text{H}_2\text{O}$.

1 H_2O . (Cleve, C. N. 48. 74.)
 scopic. (Demarçay, C. R.
 7.)

c nitrate, $2\text{Sm}(\text{NO}_3)_3$,
 $+ 24\text{H}_2\text{O}$.

lution in $\text{HNO}_3 + \text{Aq}$ (sp. gr.
 s 36.47 g. hydrous salt at 16° .
 norg. 1912, 76. 321.)

ate, basic.

(Nilson, B. 13. 1444.)
 $+ \text{H}_2\text{O}$. (Crookes, Roy. Soc.
 A. 518.)
 (Crookes.)

ate, $\text{Sc}(\text{NO}_3)_3$.

oy. Soc. Proc. 1908, 80. A, 518.)
 ery sol. in H_2O . (Crookes.)

AgNO_3 .

H_2O at 11° dissolve 127.7 pts.
 ch. Pharm. (2) 82. 260.)
) dissolve at:

54° 85° 110°
 500 714 1111 pts. AgNO_3 .

mers, Pogg. 92. 497.)

O dissolve 1622.5 pts. at 125° ,
 s. at 133° . (Tilden and Shen-
 ans. 1884. 23.)
 1 boils at 125° . (Kremers.)

Solubility in H_2O at t° .
 Sat. $\text{AgNO}_3 + \text{Aq}$ contains % AgNO_3 at t° .

t°	% AgNO_3	t°	% AgNO_3
—7	46.2	36.5	75.7
—7	46.0	40.5	76.8
—5	47.6	45	77.1
—1	52.4	48	78.5
—1	51.9	73	84.0
+5	56.3	122	88.7
10	61.2	134	92.1
15.5	66.1	135	92.8
20	67.8	135	92.7
26	71.1	148	93.3
29	73.0	160	95.2
31	73.8	182	96.9

(Étard, A. ch. 1894, (7) 2. 526.)

100 g. sat. $\text{AgNO}_3 + \text{Aq}$ at 15.5° contain
 65.5 g. AgNO_3 . (Greenish and Smith, Pharm.
 Jour. 1903, 71. 881.)

Solubility of AgNO_3 in H_2O at $30^\circ = 10.31$
 mol.-litre. (Masson, Chem. Soc. 1911, 99.
 1136.)

100 g. $\text{AgNO}_3 + \text{Aq}$ sat. at 30° contain 73.0
 g. AgNO_3 . (Schreinemakers and de Baat,
 Arch. Néer. Sc. 1911, (2) 15. 415.)

100 g. sat. $\text{AgNO}_3 + \text{Aq}$ contain 53.5 g.
 AgNO_3 at 0° ; 66.7 g. at 18° . (Mylius, Z.
 anorg. 1912, 74. 411.)

Sp. gr. of aqueous solution, according to
 C. K. = Chemiker Kalender; K. M. = Kohl-
 rausch by Mendelejeff (Z. anal. 27. 284); and
 K = Kohlrausch (W. Ann. 1879. 1), contain-
 ing:

	5	10	15	20% AgNO_3
G. K.	1.041	1.080	1.125	1.160
K. M.	1.0440	1.0901	...	1.1969
K.	1.0422	1.0893	1.1404	1.1958

	25	30	35	40% AgNO_3
C. K.	1.206	1.251
K. M.	1.4791
K.	1.2555	1.3213	1.3945	1.4773

	45	50% AgNO_3
K.	1.5705	1.6745

Sp. gr. of $\text{AgNO}_3 + \text{Aq}$ at 25° .

Concentration of AgNO_3	Sp. gr.
1-normal	1.1386
$\frac{1}{2}$ - " "	1.0692
$\frac{1}{4}$ - " "	1.0348
$\frac{1}{8}$ - " "	1.0173

(Wagner, Z. phys. Ch. 1890, 5. 40.)

Sol. in 500 pts. HNO_3 ; 30 pts. 2HNO_3 ,
 $3\text{H}_2\text{O}$ at 20° ; and 6 pts. 2HNO_3 , $3\text{H}_2\text{O}$ at 100° .
 (Schultz, Zeit. Ch. 1869. 531.)

Insol. in conc. HNO_3 . (Warren, C. C.
 1897. I, 438.)

Solubility of AgNO_3 in HNO_3 + Aq at 25° .

G. mol. per l.		G. AgNO_3 per l.	Sp. gr. 25°
HNO_3	AgNO_3		
0	10.31	1752	2.3921
0.404	9.36	1591	2.2754
0.962	8.08	1373	2.1243
1.698	6.54	1111	1.9402
2.834	4.526	769.1	1.7052
4.497	2.590	440.1	1.4980
5.992	1.698	288.6	1.4195
8.84	0.843	143.2	1.3818
12.53	0.347	58.90	1.3976

(Masson, Chem. Soc. 1911, **99**. 1132.) $\text{AgNO}_3 + \text{NH}_4\text{NO}_3$. Solubility of AgNO_3 in NH_4NO_3 + Aq. See under NH_4NO_3 . $\text{AgNO}_3 + \text{KNO}_3$. Solubility of AgNO_3 + KNO_3 in H_2O . See under KNO_3 . $\text{AgNO}_3 + \text{AgNO}_2$.Solubility of $\text{AgNO}_3 + \text{AgNO}_2$ at 18° .

G. per l.		G. per l.	
AgNO_3	AgNO_2	AgNO_3	AgNO_2
0.000	3.184	3.512	2.201
0.439	3.042	7.024	1.799
0.878	2.926	14.048	1.480
1.756	2.601		

(Naumann and Rucker, B. 1905, **38**. 2293.)See also under AgNO_2 .Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, **54**. 674.)Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 829.)

Sol. in 4 pts. boiling alcohol.

Sol. in 10 pts. alcohol. (Dumas.)

Sol. in 11 pts. alcohol of 90%. (Hager.)

Solubility in 100 pts. alcohol of given vol. % at t° .

t°	95%	80%	70%	60%
15	3.8	10.3	22.1	30.5
50	7.3	58.1
75	18.3	42.0	...	89.0

t°	50%	40%	30%	20%	10%
15	35.8	56.4	73.7	107	158
50	98.3	214	...
75	160	340	...

(Eder, J. pr. (2) 17. 44.)

100 pts. absolute methyl alcohol dissolve 3.72 pts. at 19° ; 100 pts. absolute ethyl alcohol dissolve 3.1 pts. at 19° . (de Bruyn, Z. phys. Ch. 10. 783.)Only traces are sol. in absolute alcohol ether. 100 pts. of a mixture of 1 vol. of (95 vol. %) + 1 vol. pure ether dissolve 2.3 pts. AgNO_3 at 15° ; 100 pts. of 2 vols. of + 1 vol. ether dissolve 2.3 pts. AgNO_3 (Eder, J. pr. 1878, (2) 17. 45.)Solubility of AgNO_3 in ethyl alcohol at 30° .

% by wt. H_2O	% by wt. alcohol	% by wt.
27.0	...	73.0
27.71	2.8	69.4
30.80	13.67	55.5
32.10	20.13	47.7
31.40	25.85	42.7
28.95	37.26	33.7
27.91	44.54	27.5
20.92	64.42	14.6
6.83	86.54	6.0

(Schreinemakers, Z. phys. Ch. 1909, **65**Solubility of $\text{AgNO}_3 + \text{KNO}_3$ in a See under KNO_3 .Sol. in methyl, ethyl, and isobutyl alcohol, CCl_4 , CHCl_3 , acetone and pyridine. (V J. phys. Chem. 1910, 14. 587.)100 pts. H_2O sat. with ether dissolve 2.3 pts. AgNO_3 at 15° . (Eder, l. c.)

Sol. in glycerine.

Sol. in benzonitrile. 100 g. benzene dissolve about 105 g. AgNO_3 at 18° . (Naumann and Schier, B. 1914, **47**. 1369.)1 pt. acetonitrile dissolves about 1 g. AgNO_3 . (Scholl and Steinkopf, B. 1914, **43**. 4398.)

Easily sol. in methyl, ethyl, and propylamine. (Shinn, J. phys. Chem. 1907, 11. 1014.)

Sol. in acetone. (Krug and M'Elroy, Anal. Ch. 6. 184.)

0.35 pts. are sol. in 100 pts. acetone

0.35 " " " " 100 " "

(Laszyński, B. 1894, **27**. 2287.)

Sol. in acetone and in methylal. (Eder, J. pr. 1878, II. 1014.)

1 g. AgNO_3 is sol. in 227 g. acetone. Sp. gr. of sat. solution $18^\circ/4^\circ = 0.798$. (Naumann, B. 1904, **37**. 4339.)Insol. in CS_2 . (Arctowski, Z. anorg. Ch. 1906, **6**. 257.)Difficultly sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)Insol. in methyl acetate. (Naumann, B. 1909, **42**. 3790); ethyl acetate. (Naumann, B. 1904, **37**. 3601.)Sol. in urethane. (Castoro, Z. anorg. Ch. 1906, **20**. 61.)100 g. C_2H_5 dissolve 0.022 g. AgNO_3 at 40.5° . (Linebarger, Am. J. Chem. 1895, **49**. 48.)Mol. weight determined in pipidine and benzonitrile. (Werner, Z. phys. Ch. 1897, **15**. pp. 17, 23 and 32.)

Solubility of AgNO_3 in pyridine at t° .

	G. AgNO_3 per 100 g. $\text{C}_5\text{H}_5\text{N}$	Solid phase
5°	0	$\text{C}_5\text{H}_5\text{N}$
5	3	"
	6	"
	9	"
25	11.1	$\text{C}_5\text{H}_5\text{N} + \text{AgNO}_3, 6\text{C}_5\text{H}_5\text{N}$
	11.7	$\text{AgNO}_3, 6\text{C}_5\text{H}_5\text{N}$
	12.2	"
	12.6	"
	13.9	"
	17.6	"
	18.8	" + $\text{AgNO}_3, 3\text{C}_5\text{H}_5\text{N}$
	20.03	$\text{AgNO}_3, 3\text{C}_5\text{H}_5\text{N}$
	22.34	"
	27.21	"
	33.64	"
	40.86	"
	53.52	"
	62.26	"
	63.06	"
	66.35	"
	70.85	"
5	69.85	" + $\text{AgNO}_3, 2\text{C}_5\text{H}_5\text{N}$
	72.25	$\text{AgNO}_3, 2\text{C}_5\text{H}_5\text{N}$
	78.60	"
	89.10	"
	121.21	"
	215.02	"
	228.5	"
	230.6	"
	235.4	"
	230.4	"
	237.1	"
	241.9	"
	253.8	"
	271.4	"

.. pt.

Stürenberg and Brewer, J. phys. Ch. 1908, 12. 283.)

Silver nitrate acetylde, $\text{AgNO}_3, \text{AgHC}_2$.

Illgerodt, B. 1895, 28. 2108.)

 $\text{NO}_2, \text{Ag}_2\text{C}_2$. Ppt. (Chavastelon, C. R. 124. 1365.) $\text{NO}_2, \text{Ag}_2\text{C}_2$. (Chavastelon.)Silver nitrate ammonia, $\text{AgNO}_3, \text{NH}_3$.Easily sol. in H_2O ; rather sol. in alcohol. Sol. in ether. (Reychler, B. 16. 990.)Easily sol. in liquid ammonia below -10° . (Mitscherlich, C. R. 1894, 118. 1151.) $\text{NO}_2, 2\text{NH}_3$. Easily sol. in H_2O . (Mitscherlich.)

Ethyl alcohol dissolves 0.0383 g. mols. (Kurisch, C. 1903. II, 97.)

 $\text{NO}_2, 3\text{NH}_3$. Completely sol. in H_2O . (Pogg. 20. 153.)Silver nitrate antimonide, $\text{AgNO}_3, \text{Ag}_3\text{Sb}$.Decomp. at once by H_2O . (Poleck and Thümmel, B. 16. 2435.)Silver nitrate arsenide, $\text{AgNO}_3, \text{Ag}_3\text{As}$.Decomp. at once by H_2O . (Poleck and Thümmel.)Silver nitrate bromide, $\text{AgNO}_3, \text{AgBr}$.Decomp. immediately by H_2O or alcohol, with separation of AgBr . (Risse, A. 111. 39.)Silver nitrate chloride, $\text{AgNO}_3, \text{AgCl}$.Quickly decomp. with H_2O ; more slowly with absolute alcohol; not decomp. by ether-alcohol. (Reichert, J. pr. 92. 237.)Silver nitrate cyanide, $2\text{AgNO}_3, \text{AgCN}$.Decomp. by H_2O , not by alcohol. (Hellwig, Z. anorg. 1900, 25. 177.)Silver nitrate iodide, $\text{AgNO}_3, \text{AgI}$.Cold H_2O separates AgI , which redissolves on heating. (Stürenberg, Arch. Pharm. (2) 143. 12.) Sol. in little H_2O without decomp.; more H_2O separates AgI . (Kremers, J. pr. 71. 54.) Insol. in absolute alcohol. Sol. in conc. $\text{AgNO}_3 + \text{Aq}$. $2\text{AgNO}_3, \text{AgI}$. Sol. in little but decomp. by more boiling H_2O . (Risse, A. 111. 39.)Silver nitrate mercuric oxide, $\text{AgNO}_3, 2\text{HgO}$.Decomp. by H_2O . Sol. in HNO_3 and H_2SO_4 . (Finci, Gazz. ch. it. 1910, 41. (2) 548.)Silver nitrate phosphide, $3\text{AgNO}_3, \text{Ag}_3\text{P}$.

(Warren, C. N. 56. 113.)

Silver nitrate silicide, $4\text{AgNO}_3, \text{AgSi}$.

(Büchner, Ch. Ztg. 9. 484.)

Silver nitrate silicate, $2\text{AgNO}_3, 3\text{Ag}_4\text{SiO}_4$.Sol. in dil. $\text{HNO}_3 + \text{Aq}$, but SiO_2 separates out after heating. (Rousseau and Tite, C. R. 114. 294.)Silver nitrate sulphide, $\text{AgNO}_3, \text{Ag}_2\text{S}$.Decomp. by H_2O . (Poleck and Thümmel, B. 16. 2435.)Silver nitrate sulphocyanide, $2\text{AgNO}_3, \text{AgSCN}$.Decomp. by H_2O , not by alcohol. (Hellwig, Z. anorg. 1900, 25. 178.)Sodium nitrate, NaNO_3 .Deliquescent in moist air. Sol. in H_2O with absorption of heat. 75 pts. NaNO_3 mixed with 100 pts. H_2O at 13.2° lower the temperature 18.5° . (Rüdorff, B. 2. 68.)

Sol. in 1.58 pts. H_2O at -6° .		
" 0.46 "	" $+119^\circ$	(Marx.)
" 2.89 "	" 2°	
" 1.12 "	" 28°	(Osann.)
" 0.79 "	" 47°	
" 1.14 "	" 18.5°	(Kopp.)
" 1.136 "	" 18.75°	(Karsten.)
" 1.16 "	" 20°	(Schiff, A. 100. 328.)
" 2 "	" 18.75°	(Abl.)

100 pts. H ₂ O at t° dissolve pts. NaNO ₃ .				Solubility in 100 pts. H ₂ O at t°.			
t°	Pts. NaNO ₃	t°	Pts. NaNO ₃	t°	Pts. NaNO ₃	t°	Pts. NaNO ₃
-6	68.80	50	111.13	0	72.9	60	122
0	79.75	60	119.94	1	74.7	61	124
10	84.30	70	129.63	2	75.4	62	125
16	87.63	80	140.72	3	76.0	63	126
20	89.55	90	153.63	4	76.7	64	127
33	95.37	100	168.20	5	77.4	65	128
40	102.31	120	225.30	6	78.1	66	130
(Poggiale, A. ch. (3) 8. 469.)				7	78.7	67	131
100 pts. H ₂ O at 119° dissolve 150 pts. NaNO ₃ . (Griffiths.)				8	79.4	68	132
NaNO ₃ +Aq sat. at 18.75° has 1.3769 sp. gr., and 100 pts. H ₂ O have dissolved 88.001 pts. NaNO ₃ . (Karsten.)				9	80.1	69	133
NaNO ₃ +Aq sat. in cold contains 33.3% NaNO ₃ . (Fourcroy)				10	80.8	70	134
NaNO ₃ +Aq sat. at 12.5° contains 34% NaNO ₃ . (Hassensfratz.)				11	81.4	71	136
100 pts. H ₂ O at 15.5° dissolve 33 pts., at 52°, 100 pts. NaNO ₃ . (Ure's Dict.)				12	82.0	72	137
100 pts. H ₂ O dissolve pts. NaNO ₃ at t°.				13	82.7	73	138
t°	Pts. NaNO ₃	t°	Pts. NaNO ₃	14	83.4	74	139
0	73.0	60.65	125.5	15	84.0	75	140
13.9	81.6	99.9	173.6	16	84.7	76	141
44.65	110.5	119.7	211.4	17	85.4	77	143
(Nordenskjöld, Pogg. 136. 312.)				18	86.1	78	145
100 pts. H ₂ O dissolve pts. NaNO ₃ at t°.				19	86.8	79	146
t°	Pts. NaNO ₃	t°	Pts. NaNO ₃	20	87.5	80	148
0	70.94	70	142.31	21	88.3	81	149
10	78.57	80	153.72	22	89.0	82	151
20	87.97	90	165.55	23	89.7	83	153
30	98.26	100	178.18	24	90.3	84	153
40	109.01	110	194.26	25	91.0	85	155
50	120.00	119.4	213.43	26	91.8	86	156
60	131.11	.	.	27	92.5	87	158
(Maumené, C. R. 58. 81.)				28	93.2	88	159
100 pts. NaNO ₃ +Aq sat. at 14° contain 43.88 pts. NaNO ₃ ; at 15°, 44.53 pts. NaNO ₃ . (v. Hauer, J. pr. 98. 137.)				29	94.0	89	161
100 pts. H ₂ O dissolve 84.21-84.69 pts. NaNO ₃ at 15.6°, and sat. solution has sp. gr. 1.337-1.378. (Page and Keightley, Chem Soc. (2) 10. 556.)				30	94.9	90	162
100 pts. H ₂ O dissolve pts. NaNO ₃ at t°.				31	96.0	91	164
t°	Pts. NaNO ₃	t°	Pts. NaNO ₃	32	96	92	166
0	66.69	18	83.62	33	97	93	168
2	70.97	21	85.73	34	98	94	169
4	71.04	26	90.33	35	99	95	171
8	75.65	29	92.93	36	100	96	173
10	76.31	36	99.39	37	100	97	173
13	79.00	51	113.63	38	101	98	177
15	80.60	68	125.07	39	102	99	178
Solubility is constant from 0° to -15.7°, when NaNO ₃ +7H ₂ O separates out. (Ditte, C. R. 80. 1164)				40	102	100	180
				41	103	101	183
				42	104	102	184
				43	105	103	186
				44	106	104	188
				45	107	105	190
				46	108	106	192
				47	109	107	194
				48	110	108	196
				49	111	109	198
				50	112	110	200
				51	113	111	203
				52	114	112	204
				53	115	113	207
				54	116	114	209
				55	117	115	211
				56	118	116	212
				57	119	117	215
				58	120	117.5	216.4
				59	211

(Mulder, Scheik. Verhandl. 1864. 81.)

solution at b.-pt. contains 216.4 pts. NaNO₃ (Marx); 218.5 pts. NaNO₃ (Marx); 213.4 pts. NaNO₃ (Nordenskjöld); 224.8 pts. NaNO₃ (Legrand); 150 pts. NaNO₃ (Griffiths).

NaNO₃+Aq contains at:
130° 172° 180° 199°
67.5 77.1 78.1 82.0% NaNO₃,

250° 255° 290° 313° (mpt.).
89.5 91.5 97.5 100% NaNO₃.
(Étard, A. ch. 1894, (7) 2. 527.)

s. sat. NaNO₃+Aq contain 42.47 g. at 0°. (Coppadoro, Rass. Min. 1911, 15. 123.)

s. sat. NaNO₃+Aq contain 49.16 g. at 30°. (Coppadoro, Rass. Min. 1911, 15. 7.)

H₂O dissolve 92.14 g. NaNO₃ at 25°. J. Am. Chem. Soc. 1912, 34. 1148.)

solubility of crystals on different faces determined by Lebrun. (Belg. Acad. 18. 953.)

p. gr. of NaNO₃+Aq at 19.5°.

O ₃	Sp. gr.	% NaNO ₃	Sp. gr.
7	1.0844	39.860	1.3176
8	1.1667	46.251	1.3805
7	1.2450

(Kremers, Pogg. 95. 120.)

p. gr. of NaNO₃+Aq at 20.2°.

O ₃	Sp. gr.	% NaNO ₃	Sp. gr.
	1.0065	26	1.1904
	1.0131	27	1.1987
	1.0197	28	1.2070
	1.0264	29	1.2154
	1.0332	30	1.2239
	1.0399	31	1.2325
	1.0468	32	1.2412
	1.0537	33	1.2500
	1.0606	34	1.2589
	1.0676	35	1.2679
	1.0746	36	1.2770
	1.0817	37	1.2863
	1.0889	38	1.2958
	1.0962	39	1.3055
	1.1035	40	1.3155
	1.1109	41	1.3225
	1.1184	42	1.3355
	1.1260	43	1.3456
	1.1338	44	1.3557
	1.1418	45	1.3659
	1.1498	46	1.3761
	1.1578	47	1.3864
	1.1659	48	1.3968
	1.1740	49	1.4074
	1.1822	50	1.4180

calculated by Gerlach, Z. anal. 8. 280.)

Sp. gr. of NaNO₃+Aq at 18°.

% NaNO ₃	Sp. gr.	% NaNO ₃	Sp. gr.
5	1.0327	20	1.1435
10	1.0681	30	1.2278

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of NaNO₃+Aq at 20°, containing mols. NaNO₃ in 100 mols. H₂O.

Mols. NaNO ₃	Sp. gr.
2	1.05980
5	1.13813

(Nicol, Phil. Mag. (5) 16. 122.)

The saturated solution boils at 117.5° (Mulder.)
" " " 118.9° (Griffiths.)
" " " 119° (Marx.)
" " " 119.4° (Maumené.)
" " " 119.7° (Nordenskjöld.)
" " " 121° (Legrand.)
" " " 122-123° (Kremers.)

NaNO₃+Aq forms a crust at 118°, and contains 194 pts. NaNO₃ to 100 pts. H₂O; highest temp. observed, 120.5°. (Gerlach, Z. anal. 26. 427.)

B.-pt. of NaNO₃+Aq containing pts. NaNO₃ to 100 pts. H₂O. G=according to Gerlach (Z. anal. 26. 433); L=according to Legrand (A. ch. (2) 59. 431).

B.-pt.	G	L	B.-pt.	G	L
101°	9	9.3	112°	121.5	120.3
102	18.5	18.7	113	133	131.3
103	28	28.2	114	144.5	142.4
104	38	37.9	115	156	153.7
105	48	47.7	116	168.5	165.2
106	58	57.6	117	181	176.8
107	68	67.7	118	194	188.6
108	78.5	77.9	119	207.5	200.5
109	89	88.3	120	222	212.6
110	99.5	98.8	121	...	224.8
111	110.5	109.5

50 pts. NaNO₃ mixed with 100 pts. snow at -1° give a temp. of -17.5°. (Rüdorff, Pogg. 122. 337.)

Sp. gr. of NaNO₃+Aq at t°.

G. NaNO ₃ dissolved in 100 g. H ₂ O	G. NaNO ₃ in 100 g. of the solution	t°	Sp. gr.
4.166	4	17.8°	1.0276
11.111	10	13.9°	1.0704
25.000	20	12°	1.1441

(de Lannoy, Z. phys. Ch. 1895, 18. 465.)

Sp. gr. of $\text{NaNO}_3 + \text{Aq}$ at 20.1° , when p = per cent strength of sol.; d = observed density; w = volume conc. in grams per cc. ($\frac{pd}{100} = w$)

p	d	w
42.05	1.3380	0.56267
35.65	1.2765	0.45510
31.72	1.2407	0.39365
23.24	1.1696	0.27180
17.370	1.1228	0.19505
11.915	1.0819	0.12888
9.665	1.0656	0.10300
7.039	1.0468	0.07369
4.241	1.0273	0.04357
1.589	1.0096	0.01604

(Barnes, J. phys. Chem. 1898, 2. 545.)

Sp. gr. $20^\circ/4^\circ$ of a normal solution of NaNO_3 = 1.05386; of a 0.5 normal solution = 1.02646. (Haigh, J. Am. Chem. Soc. 1912, 34. 1151.)

Sp. gr. of sat. $\text{NaNO}_3 + \text{Aq}$ at t° .

t°	G. NaNO_3 sol. in 100 g. H_2O	Sp. gr.
-10	68.0	1.342
0	43.0	1.358
10	80.5	1.377
20	88.0	1.387
30	96.2	1.406
40	104.9	1.418
50	114.0	1.437
60	124.6	1.456
70	136.0	1.467

(Tschernaj, J. Russ. Phys. Chem. Soc. 1912, 44. 1565.)

Sol. in 66 pts. HNO_3 ; in 32 pts. 2HNO_3 , $3\text{H}_2\text{O}$ at 32° ; in 4 pts. 2HNO_3 , $3\text{H}_2\text{O}$ at 123° . (Schultz, Zeit. Ch. (2) 5. 531.)

Solubility in $\text{HNO}_3 + \text{Aq}$ at 0° .

G. per 100 cc. of solution		Sp. gr.
NaNO_3	HNO_3	
56.5	0.00	1.341
54.2	1.67	1.338
51.48	3.59	1.331
48.42	5.55	1.324
44.88	7.92	1.312
41.44	10.65	1.308
33.61	17.02	1.291
29.86	20.33	1.285
26.46	23.48	1.282
20.00	30.26	1.276
15.32	36.09	1.276
10.97	44.76	1.291

(Engel, C. R. 1887, 104. 911.)

Solubility of NaNO_3 in $\text{NH}_4\text{OH} + \text{Aq}$

G. per 100 g. H_2O		%
NH_3	NaNO_3	
13.87	75.03	1.
17.28	73.99	1.
20.38	73.18	1.

(Fedotieff and Koltunoff, Z. anorg. 251.)

$\text{NaNO}_3 + \text{Na}_2\text{CO}_3$. (See Na_2CO_3 .)
Sol. in sat. $\text{NH}_4\text{Cl} + \text{Aq}$.
Very rapidly sol. in sat. $\text{BaCl}_2 +$ pptn. of $\text{Ba}(\text{NO}_3)_2$.
Sol. in sat. $\text{KCl} + \text{Aq}$, with form KNO_3 .
Sol. in sat. $\text{NH}_4\text{NO}_3 + \text{Aq}$. (See NH_4NO_3 .)
Sol. in sat. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$, with pptn. of $\text{Ba}(\text{NO}_3)_2$. (See $\text{Ba}(\text{NO}_3)_2$.)
Sol. in sat. $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$, with pptn. of $\text{Pb}(\text{NO}_3)_2$. (See $\text{Pb}(\text{NO}_3)_2$.)
 $\text{NaNO}_3 + \text{KNO}_3$.
Sol. in sat $\text{KNO}_3 + \text{Aq}$; solution at 18° contains 54.33% mixed salt, H_2O dissolve 118.98 pts. mixed, 89.53 pts. NaNO_3 and 29.45 pts (See KNO_3 .)
 $\text{NaNO}_3 + \text{Sr}(\text{NO}_3)_2$.
If $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ sat. at 14.5° is NaNO_3 , 100 pts. H_2O dissolve:

NaNO_3	83.7	66.4
$\text{Sr}(\text{NO}_3)_2$	51.0
		117.4

(Mulder.)

$\text{NaNO}_3 + \text{NaNO}_2$. See under $\text{NaNO}_2 + \text{NaCl}$.
100 pts. H_2O dissolve 24.91 pts 54.55 pts. $\text{NaNO}_3 = 79.46$ pts. of the at 20° . (Nicol, Phil. Mag. (5) 31. 3

100 pts. H_2O dissolve at 18.7

	1	2	3	4
NaCl	36	25.22	24.96	24.98
NaNO_3	..	52.89	52.84	52.82

2. Sat. $\text{NaCl} + \text{Aq}$ treated with 1
3. Sat. $\text{NaNO}_3 + \text{Aq}$ treated with
4. Simultaneous treatment of the by H_2O . (Karsten.)
6. Excess of both salts + Aq was cooled to 20° . (Rüdorff, B. 6. 484.)

ty of NaCl with addition of NaNO₃ at 15.5°.

	100 ccm. contain in g.		
	NaCl	H ₂ O	NaNO ₃
5	31.78	88.47	0.00
5	27.89	87.63	7.53
0	26.31	86.25	13.24
0	23.98	82.66	21.58
0	22.30	80.42	28.18
5	20.40	79.25	33.80
5	19.40	77.37	37.88
5	19.67	77.34	37.64

O₂ separated in last two solutions.

y of NaNO₃ with addition of NaCl at 15°.

	100 ccm. contain in g.		
	NaNO ₃	H ₂ O	NaCl
0	62.38	74.82	0
5	56.56	75.69	4.00
5	52.09	75.71	7.24
0	47.08	76.86	11.36
5	42.66	76.96	15.33
5	39.90	77.14	17.81
5	38.73	77.15	18.97
5	38.02	77.49	19.34

l separated in last two solutions.
odländer, Z. phys. Ch. 7. 360.)

y of NaNO₃ in NaCl+Aq at 15°.

	G. per 100 cc. sat. solution		
	NaCl	NaNO ₃	H ₂ O
)	0	62.38	74.82
5	4.0	56.76	75.69
5	7.24	52.09	75.71
)	11.36	47.08	76.86
5	15.33	42.66	76.96
5	17.81	39.90	77.14
5	18.97*	38.73*	77.15
5	19.34*	38.02*	77.49

tions sat. with both salts.
änder, Z. phys. Ch. 1891, 7. 361.)

y of NaNO₃+NaCl (g. in 100 g. H₂O) at 25°.

	NaCl	Solid phase
	8.39	NaNO ₃
	16.32	"
	23.74	NaNO ₃ +NaCl
	27.56	NaCl
	31.48	"

Mem. Col. Sc. Kyoto, 1910, 2. 245.)

Solubility in NaCl+Aq at 20°, 30°, 40° and 91°. Tables given in the original show that each salt diminishes the solubility of the other. (Leather, Mem. Dept. Agric. India, 1914, 3. 177; Chem. Soc. 1915, 106. (2) 13.)
See also under NaCl.

NaNO₃+NaOH.
Solubility in NaOH+Aq at 0°. NaNO₃= mols. NaNO₃ (in mg.) in 10 ccm. of solution; Na₂O=mols. Na₂O (in mg.) in 10 ccm. of solution.

NaNO ₃	Na ₂ O	NaNO ₃ +Na ₂ O	Sp. gr.
66.4	0	66.4	1.341
62.5	2.875	65.375	1.338
57.15	6.1	63.25	1.333
47.5	12.75	60.25	1.327
29.5	26.	55.5	1.326
17.5	39	56.5	1.332
13.19	45.875	59.065	1.356
6.05	60.875	66.925	1.401

(Engel, Bull. Soc. (3) 6. 16.)
Solubility in NaOH+Aq at 0°.

G. per 100 cc. solution		Sp. gr.
NaOH	NaNO ₃	
0.0	56.50	1.341
2.30	53.19	1.338
4.89	48.63	1.333
10.21	40.42	1.327
20.83	25.10	1.326
31.25	14.89	1.332
36.76	11.22	1.356
48.75	5.15	1.401

(Engel, l. c.)
Easily sol. in K₂SO₄+Aq without pptn.
Easily sol. in Na₂SO₄+Aq without pptn.
Sol. in MgSO₄+Aq, at first to a clear solution, but afterwards NaNO₃ is pptd.
Very sol. in sat. CuSO₄+Aq, but double sulphate separates out.
Very sol. in ZnSO₄+Aq with pptn. of double sulphate. (Karsten.)

Solubility of NaNO₃ in Na₂S₂O₃+Aq at t°.

t°	% NaNO ₃	% Na ₂ S ₂ O ₃	Solid phase
9	33.31	12.26	NaNO ₃
	22.57	23.41	" +Na ₂ S ₂ O ₃ , 5H ₂ O
	4.22	34.77	Na ₂ S ₂ O ₃ , 5H ₂ O
25	35.42	12.72	NaNO ₃
	25.40	24.25	"
	19.90	31.81	" +Na ₂ S ₂ O ₃ , 5H ₂ O
	18.02	32.83	Na ₂ S ₂ O ₃ , 5H ₂ O
	4.33	40.50	"

(Kremann and Rothmund, Z. anorg. 1914, 86. 373.)

Very sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Easily sol. in liquid HF. (Franklin, Z. anorg. 1905, 46. 2.)

Hydrazine dissolves 26.6 pts. NaNO₃ at 12.5–13°. (de Bruyn, R. t. c. 1899, 18. 297.)

100 pts. alcohol of 0.9 sp. gr. dissolve 10.5 pts. NaNO₃; 0.872 sp. gr., 6 pts.; 0.834 sp. gr., 0.38 pt.; insol. in alcohol of 0.817 sp. gr. (Kirwan.)

100 pts. alcohol of 61.4% by weight dissolve 21.2 pts. NaNO₃ at 26°. (Pohl, W. A. B. 6. 600.)

100 pts. alcohol of 62° Tr. dissolve 7.4 pts. NaNO₃ at 19.5°.

100 pts. alcohol of 93° Tr. dissolve 0.93 pt. NaNO₃ at 19.5°. (Wittstein.)

100 pts. alcohol containing % alcohol by weight dissolve pts. NaNO₃ at 15°, or 100 pts. solution contain % NaNO₃:

10	20	30	40	60	80% alcohol.
65.3	48.8	35.5	25.8	11.4	2.8 pts. NaNO ₃ .
39.5	32.8	26.2	20.5	10.2	2.7% NaNO ₃ .

(Schiff.)

100 pts. wood-spirit of 40% dissolve 32.3 pts. NaNO₃. (Schiff, A. 118. 365.)

Solubility in alcohol at 16.5°.

Sp. gr.	100 ccm. contain in g.		
	Alcohol	Water	NaNO ₃
1.3745	0	75.25	62.20
1.3162	6.16	70.82	54.64
1.2576	11.60	68.10	46.06
1.2140	16.49	65.04	39.87
1.1615	22.17	61.67	32.31
1.0855	32.22	52.92	23.41
1.0558	37.23	48.50	19.85
1.0050	43.98	42.78	13.74
0.9420	52.60	32.13	9.47
0.9030	60.00	25.65	4.65
0.8610	63.16	21.31	1.63

(Bodländer, Z. phys. Ch. 7. 317.)

100 pts. absolute methyl alcohol dissolve 0.41 pt. at 25°.

100 pts. absolute ethyl alcohol dissolve 0.036 pt. at 25°. (de Bruyn, Z. phys. Ch. 10. 783.)

Solubility in alcohol at 40°.

Wt. % alcohol	G. NaNO ₃ per 100 g. alcohol + Aq.
0	104.5
8.22	90.8
17.4	73.3
26.0	61.6
36.0	48.4
42.8	40.6
55.3	27.1
65.1	18.1
77.0	9.4
87.2	4.2

(Bathrick, J. phys. Ch. 1896, 1. 162.)

Solubility in alcohol at 30°.

Wt. % alcohol in solvent	G. NaNO ₃ per 100	
	Solution	Wt.
0	49.10	96
5	46.41	91
10	43.50	85
20	37.42	74
30	31.31	65
40	25.14	55
50	18.94	46
60	12.97	37
70	7.81	28
90	1.21	12

(Taylor, J. phys. Ch. 1897, 1. 72

Solubility in ethyl alcohol at 21°

(Concentration of alcohol in g. : 1000 g. H₂O.)

Normality	Solubility in 1000 g. H ₂ O	Mol. %
1/4	920.30	10
1/2	908.80	10
1	896.60	10
1	870.95	10
2	825.35	9

(Armstrong and Eyre, Proc. R. Soc. 184. 127.)

Very sl. sol. in acetone. (Krug and roy, J. Anal. Ch. 6. 184.)

Solubility of NaNO₃ in acetone at

Wt. % acetone	G. NaNO ₃ per acetone + A
0.0	105
8.47	91.2
16.8	78.3
25.2	66.4
34.3	57.9
44.1	46.2
53.9	32.8
64.8	23.0
76.0	10.8
87.6	3.2

(Bathrick, J. phys. Ch. 1896, 1. 16

Solubility of NaNO_3 in acetone at 30° .

State of	G. NaNO_3 per 100 g.	
	Solution	Water
	49.10	96.45
	46.96	93.20
	45.11	90.40
	40.10	83.70
	35.08	77.20
	29.80	70.75
	24.34	64.40
	18.55	59.95
	13.15	50.50
	7.10	38.20
	1.98	20.20

(*Ann. Chem. Phys.* 1897, 2, 723.)

glycerine.

in ethylamine. (Shinn, *J. phys. Chem.* 1907, 11, 538.)

in methyl acetate. (Naumann, *B. Chem.* 1879, 12, 3790); ethyl acetate. (Naumann, *B. Chem.* 1879, 12, 314.)

in benzonitrile. (Naumann, *B. Chem.* 1914, 47, 1914.)

thorium nitrate, $\text{NaTh}(\text{NO}_3)_6 + 6\text{H}_2\text{O}$.

soluble; sol. in dil. HNO_3 and Aq . (*Z. anorg. Chem.* 1901, 27, 381.)

nitrate sulphate, $\text{NaNO}_3, \text{Na}_2\text{SO}_4 + \text{H}_2\text{O}$.

(*Marignac, Ann. Min.* (5) 12.

Min. Darapskite.

nitrate tungstosilicate, $3\text{Na}_2\text{W}_{12}\text{SiO}_{40} + 45\text{H}_2\text{O}$.

(*Boff, Chem. Soc.* 1897, 72, (2) 174.)

1 nitrate, $\text{Sr}(\text{NO}_3)_2$.

pts. cold, and 0.5 pt. boiling H_2O . (Dumas.)

" at 18.75° (Abl.) (Wittstein)

sat. $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at $19-20^\circ$ contain 45.49 g. (*v. Hauer, J. pr.* 98, 137)

$\text{Sr}(\text{NO}_3)_2$ dissolves in pts. H_2O at t°

H_2O	t°	Pts. H_2O	t°	Pts. H_2O
32	25	1.10	75	0.99
73	50	1.02	100	0.94

(Kremers, *Pogg.* 92, 499.)

100 pts. H_2O dissolve at 0° , 39.5 pts. $\text{Sr}(\text{NO}_3)_2$ (Mulder); at 0° , 40.16 pts. $\text{Sr}(\text{NO}_3)_2$ (Poggiale); at 0° , 43.1 pts. $\text{Sr}(\text{NO}_3)_2$ (Kremers); at 100° , 101.1 pts. $\text{Sr}(\text{NO}_3)_2$ (Mulder); at 100° , 106.5 pts. $\text{Sr}(\text{NO}_3)_2$ (Kremers, *Pogg.* 92, 499); at 100° , 119.25 pts. $\text{Sr}(\text{NO}_3)_2$ (Poggiale).

Solubility in 100 pts. H_2O at t° .

t°	Pts. $\text{Sr}(\text{NO}_3)_2$	t°	Pts. $\text{Sr}(\text{NO}_3)_2$	t°	Pts. $\text{Sr}(\text{NO}_3)_2$
0	39.5	36	90.7	73	96.0
1	41.2	37	90.8	74	96.2
2	42.8	38	91.0	75	96.4
3	44.3	39	91.1	76	96.5
4	45.8	40	91.3	77	96.7
5	47.3	41	91.4	78	96.8
6	48.8	42	91.5	79	97.0
7	50.3	43	91.6	80	97.2
8	51.8	44	91.8	81	97.4
9	53.4	45	91.9	82	97.5
10	54.9	46	92.1	83	97.7
11	56.5	47	92.2	84	97.9
12	58.0	48	92.3	85	98.0
13	59.6	49	92.5	86	98.2
14	61.2	50	92.6	87	98.4
15	62.8	51	92.8	88	98.6
16	64.4	52	92.9	89	98.8
17	66.0	53	93.1	90	99.0
18	67.6	54	93.2	91	99.2
19	69.2	55	93.4	92	99.4
20	70.8	56	93.5	93	99.6
21	72.5	57	93.6	94	99.8
22	74.1	58	93.8	95	100.0
23	75.8	59	93.9	96	100.2
24	77.4	60	94.0	97	100.4
25	79.0	61	94.2	98	100.6
26	80.7	62	94.3	99	100.9
27	82.4	63	94.5	100	101.1
28	84.1	64	94.6	101	101.3
29	85.8	65	94.8	102	101.6
30	87.6	66	94.9	103	101.8
31	89.5	67	95.1	104	102.0
31.3	90.0	68	95.2	105	102.3
32	90.2	69	95.4	106	102.5
33	90.3	70	95.6	107	102.7
34	90.5	71	95.7	107.9	102.9
35	90.6	72	95.9		

(Mulder, *Scheik. Verhand.* 1864, 114.)

Sat. $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ contains at:

-6° $+14^\circ$ 20° 32°
24.5 35.9 39.8 46.9% SrNO_3

53° 56° 76° 94° 110°
47.2 47.8 49.1 50.4 50.2% SrNO_3

(Étard, *A. ch.* 1894, (7) 2, 528.)

79.27 g. anhydrous $\text{Sr}(\text{NO}_3)_2$ are sol. in 100 g. H_2O at 25° . (Parsons and Carson, *J. Am. Chem. Soc.* 1910, 32, 1385.)

Solubility of $\text{Sr}(\text{NO}_3)_2$ in H_2O at t° .		
t°	G. $\text{Sr}(\text{NO}_3)_2$ in 100 g. H_2O	Sp. gr.
0.58	40.124	1.2856
14.71	60.867	1.3938
26.40	82.052	1.4883
29.06	87.648	1.5110
30.28	88.577	1.5144
32.58	88.943	1.5141
39.74	90.086	1.5128
47.73	91.446	1.5115
61.34	93.856	1.5105
68.96	95.576	1.5106
78.98	97.865	1.5109
88.94	100.136	1.5117

(Berkeley and Appleby, Proc. R. Soc. 1911, (A) 85. 503.)

100 g. of the sat. solution contain at 20° , 41.43 g. $\text{Sr}(\text{NO}_3)_2$. (Findlay, Chem. Soc. 1914, 105. 782.)

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at 19.5° .			
$\text{Sr}(\text{NO}_3)_2$	Sp. gr.	$\text{Sr}(\text{NO}_3)_2$	Sp. gr.
1	1.009	21	1.192
2	1.017	22	1.202
3	1.025	23	1.213
4	1.034	24	1.223
5	1.041	25	1.233
6	1.049	26	1.246
7	1.059	27	1.257
8	1.068	28	1.268
9	1.076	29	1.280
10	1.085	30	1.292
11	1.095	31	1.304
12	1.103	32	1.316
13	1.113	33	1.330
14	1.122	34	1.340
15	1.131	35	1.354
16	1.140	36	1.367
17	1.150	37	1.381
18	1.160	38	1.395
19	1.170	39	1.410
20	1.181	40	1.422

(Kremers, calculated by Gerlach, Z. anal. 8. 286.)

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at 23.4° . a=no. of grms. $\times \frac{1}{2}$ mol. wt. dissolved in 1000 grms. H_2O ; b=sp. gr. if a is $\text{Sr}(\text{NO}_3)_2$, $4\text{H}_2\text{O}$, $\frac{1}{2}$ mol. wt. = 142; c=sp. gr. if a is $\text{Sr}(\text{NO}_3)_2$, $\frac{1}{2}$ mol. wt. = 106.

a	b	c	a	b	c
1	1.078	1.081	5	1.303	1.350
2	1.146	1.155	6	1.345	1.407
3	1.205	1.224	7	1.383	...
4	1.257	1.284

(Favre and Valson, C. R., 79. 968.)

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at 17.5°		
% $\text{Sr}(\text{NO}_3)_2$	Sp. gr.	% $\text{Sr}(\text{NO}_3)_2$
10	1.083	40
20	1.180	Sat. sol.
30	1.294

(Gerlach, Z. anal. 27. 283.)

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at t°		
t°	% $\text{Sr}(\text{NO}_3)_2$	Sp.
14.0°	5	1.0
14.3°	10	1.0
14.5°	15	1.1
14.5°	20	1.1
14.5°	25	1.2
14.4°	34.33	1.3

(Long, W. Ann. 1880, 11. 39

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at room containing:
10.29 21.19 32.61% $\text{Sr}(\text{NO}_3)_2$
1.0885 1.124 1.3067
(Wagner, W. Ann. 1883, 18. 2

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at 2	
Concentration of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$	Sp. gr.
1-normal	1.08
$\frac{1}{2}$ " "	1.04
$\frac{1}{4}$ " "	1.02
$\frac{1}{8}$ " "	1.01

(Wagner, Z. phys. Ch. 1890, 6.

$\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ containing 10.50% has sp. gr $20^\circ/20^\circ = 1.0905$.
 $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ containing 25.51% has sp. gr. $20^\circ/20^\circ = 1.2440$.
(Le Blanc and Rohland, Z. phys. 19. 279.)

Sp. gr. of $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ at 20° c
M g. mols. of salt per liter.
M 0.01 0.025 0.05
Sp. gr. 1.001525 1.004207 1.008391

M 0.10 0.25 0.50
Sp. gr. 1.016834 1.04201 1.08312

M 1.00
Sp. gr. 1.16354
(Jones and Pearce, Am. Ch. J. 1907,

Sr(NO₃)₂ + Aq, containing pts. Sr(NO₃)₂ to 100 pts. H₂O.

Pts. Sr(NO ₃) ₂	B.-pt.	Pts. Sr(NO ₃) ₂
12	104°	81.4
24	104.5	89.6
34.8	105	97.6
45	105.5	105
54.4	106	112.2
63.6	106.3	116.5
72.6

Gerlach, Z. anal. 26. 448.)

NO₃)₂ + Aq boils at 106.8°, and con- 9 pts. salt to 100 pts. H₂O. (Grif-

(NO₃)₂ + Aq boils at 107.5–108° ; 107.9° (Mulder).

NO₃)₂ + Aq forms a crust at 106.3°, ins 116.5 pts. Sr(NO₃)₂ to 100 pts. hest temp. observed was 107°. Z. anal. 26. 427.)

sol. in conc. HNO₃ or HCl + Aq.

1 HNO₃ + Aq. (Schultz, Zeit. Ch.)

ty in Sr(OH)₂, 8H₂O + Aq at 25°.

/25°	G. SrO as Sr(OH) ₂ in 100 g. H ₂ O	G. Sr(NO ₃) ₂ in 100 g. H ₂ O
	0.38	79.47
	0.78	80.83

J. Am. Chem. Soc. 1910, 32. 1388.)

l. in liquid NH₃. (Franklin, Am. 8, 20. 829.)

3500 pts. absolute alcohol. Sol. in . of a mixture of 1 pt. ether and 1 l. (Rose, Pogg. 110. 296.)

1189 pts. abs. alcohol and in 199.87 ary rectified spirits. (Hill, Pharm.) 19. 420.)

lity in ethyl alcohol + Aq at 25°.

I in nt	% C ₂ H ₅ OH in the solution	% Sr(NO ₃) ₂ in the solution
	99.38	0.02
	77.15	2.60
	53.6	10.5
	32.35	20.5
	13.8	33.2
	12.35	34.3
	10.45	35.7
	9.5	36.7
	6.0	40.05
	3.45	42.7
	0	46.6

the solutions which contain small amounts of alcohol.

(D'Ans and Siegler, Z. phys. Ch. 1913, 82. 39.)

Not completely insol. in boiling amyl alcohol, 30 ccm. dissolving about 1 mg. (Browning, Sill. Am. J. 143. 52.)

Perfectly anhydrous Sr(NO₃)₂ is sol. in 83044 pts. absolute ether-alcohol (1 : 1). (Fresenius, Z. anal. 32. 190.)

Solubility in organic solvents.

Solvent	% Sr(NO ₃) ₂ in the solution at 25°
Methyl alcohol	1.26
Ethyl alcohol	0.02
Propyl alcohol	0.02
Isobutyl alcohol	0.01
Amyl alcohol	0.003
Acetone	0.02

(D'Ans and Siegler, Z. phys. Ch. 1913, 82. 44.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Sol. in acetone. (Eidmann, C. C. 1899. II, 1014.)

Difficultly sol. in acetone. (Naumann, B. 1904, 37. 4328.)

The composition of the hydrates formed by Sr(NO₃)₂ at different dilutions is calculated from determinations of the lowering of the fr.-pt. produced by Sr(NO₃)₂ and of the conductivity and sp. gr. of Sr(NO₃)₂ + Aq. (Jones, Am. Ch. J. 1905, 34. 305.)

+4H₂O. Efflorescent.

Solubility in ethyl alcohol + Aq at 25°.

% C ₂ H ₅ OH in the solvent	% C ₂ H ₅ OH in the solution	% Sr(NO ₃) ₂ in the solution
0	0	44.25
4	1.7	42.8
6	2.6	42.1
10.8	4.95	40.4
16.0	7.95	37.6
20	12.35	34.3

(D'Ans and Siegler, Z. phys. Ch. 1913, 82. 39.)

Tellurium nitrate, basic, 4TeO₂, N₂O₅ + 1½H₂O.

Very hygroscopic. Easily decomp. by H₂O. Sol. in HNO₃ + Aq, but more sol. when dil. than conc. (Klein and Morel, Bull. Soc. (2) 43. 205.)

Tellurium nitrate, TeNO₃.

Sol. in acetone. (Eidmann, C. C. 1899. IX, 1014.)

is the solid phase in the solutions rich in alcohol; Sr(NO₃)₂ + 4H₂O in

Terbium nitrate, $Tb(NO_3)_3 \cdot 6H_2O$.
Sol. in H_2O . Sol. in alcohol. (Urbain, C. R. 1908, 146. 128.)

Thallous nitrate, $TlNO_3$.
1 pt. $TlNO_3$ dissolves, according to C = Crookes; L = Lamy:
at 15° 18° 58° 107°
in 9.4 10.3 2.3 0.17 pts. H_2O .
C L L L

Sat. $TlNO_3$ + Aq contains at:
3.5° 18° 32° 58° 95°
4.2 8.8 13.2 30.4 74.5% $TlNO_3$

107° 135° 145° 150° 155°
85 95 95.2 96.5 97% $TlNO_3$.
(Étard, A. ch. 1894, (7) 2. 527.)

Solubility in H_2O at t°.

t°	G. $TlNO_3$ in 100 g. H_2O	g. mol. $TlNO_3$ in 1 l.
0	3.91	0.149
10	6.22	0.230
20	9.55	0.357
25	0.433
30	14.3	0.522
40	20.9	0.755
50	30.4	1.07
60	46.2	1.58
70	69.5	2.29
80	111	3.40
90	200	5.32
100	414	8.29
105	594	10.25

(Berkeley, Trans. Roy. Soc. 1904, 203. A, 211.)

Sp. gr. of $TlNO_3$ + Aq at 25°.

Concentration of $TlNO_3$ + Aq	Sp. gr.
$\frac{1}{5}$ -normal	1.0562
$\frac{1}{5}$ " "	1.0283

(Wagner, Z. phys. Ch. 1890, 5. 40.)

Solubility of $TlNO_3$ + KNO_3 . (See KNO_3 .)
Insol. in alcohol. (Lamy.)
Sol. in acetone. (Eidmann, Dissert. 1899;
Naumann, B. 1904, 37. 4328.)

Thallous hydrogen nitrate, $TlNO_3 \cdot 2HNO_3$.
(Wells, Am. Ch. J. 1901, 26. 273.)
 $TlNO_3 \cdot 3HNO_3$. (Ditte.)

Thallic nitrate, $Tl(NO_3)_3 \cdot 3H_2O$.
Effloresces in the air. (Meyer, Z. anorg. 1900, 24. 361.)
+ $6H_2O$, or $8H_2O$. Deliquescent. Sol. in H_2O .

Thallic thallic nitrate, $2TlNO_3 \cdot Tl$.
Decomp. by H_2O . (Wells, A. 1901, 26. 278.)

Thallic uranyl nitrate, $Tl(UO_2)_2$.
Decomp. in moist air. Decom. (Meyer, B. 1903, 36. 4058.)

Thorium nitrate, $ThO_2 \cdot 2N_2O_5$ + 6
Crystallized. Sl. hygroscopic. (Zeit. angew. Ch. 1897, 10. 116.)
+ $12H_2O$. Very deliquescent, H_2O and alcohol.
Difficultly sol. in acetone. (N. 1904, 37. 4328.)

Thorium zinc nitrate, $ZnTh(NO_3)_6$.
Sol. in HNO_3 ; very hygroscopic. (Z. anorg. 1901, 27. 386.)

Thulium nitrate, $Tm_2(NO_3)_6 \cdot 8H_2O$.
Deliquescent. Can readily be decomposed by HNO_3 . (James, J. Am. Chem. Soc. 1904, 26. 1344.)

Tin (stannous) nitrate, basic, $2Sn(NO_3)_2 \cdot H_2O$.
Difficultly sol. with partial decomposition. (Weber, J. pr. (2), 26. 121.)

Tin (stannous) nitrate, $Sn(NO_3)_2$.
Deliquescent, and easily decomposed. (J. pr. (2) 26. 121.)

Tin (stannic) nitrate, basic, $4Sn(NO_3)_2 \cdot 4H_2O$.
(Thomas, Bull. Soc. 1896 (3) 121.)

Tin (stannic) nitrate, $Sn(NO_3)_4$.
Sol. in H_2O , but decomp. on standing. Stable in presence of conc. Aq at 90°, but decomp. at 100°. (Martini, Gazz. ch. it. 22. 384.)
Insol. in moderately conc. H_2O . Decomp. by H_2O . (Engel, C. R. 1891, 710.)

Titanium nitrate, $5TiO_2 \cdot N_2O_5$ + 6
Sol. to a slight milkiness in H_2O . Decomp. on boiling. (Menz, J. pr. 1901, 26. 121.)

Uranyl nitrate, basic.
Sol. in H_2O . (Ordway, Sill. A. 1901, 209.)

Uranyl nitrate, $UO_2(NO_3)_2$.
+ H_2O . (de Forcrand, C. R. 1891, 1046.)
Sol. in fuming HNO_3 , from which it can be cryst.
+ $2H_2O$. 52.39% is sol. in dry H_2O .
54.25% " " " " "
(Lebeau, Bull. Soc. 1911, (4) 121.)

+2H₂O. (Vasilieff, C. C. 1910, II. 1527.)
+3H₂O. Mpt. 121.5°. (Vasilieff.)
Cryst. out of hot HNO₃+Aq. (Ditte.)
100 pts. HNO₃ dissolve 39 pts. at 14°. (Ditte, A. ch. 1879, (5) 18. 337.)
+4H₂O. (de Coninck, C. C. 1901, I. 1354.)
+6H₂O. Deliquescent in moist, and efflorescent in dry air. Sol. in 0.5 pt. cold H₂O, in 3 pt. absolute alcohol, and in 4.0 pts. ether. (Bucholz.)
Melts in crystal H₂O at 59.4°. (Ordway.)
1 pt. is sol. in 2 pts. H₂O at 12.9°-14.2°. (de Coninck, C. R. 1900, 131. 1220.)

Solubility in H₂O at t°.

t°	% by wt. UO ₂ (NO ₃) ₂ ·6H ₂ O
-18.1	54.90
-12.1	58.00
- 2.2	62.13
0	63.01
+12.3	67.36
25.6	72.83
36.7	78.05
45.2	82.96
71.8	86.32

Vasilieff, J. Russ. Phys. Chem. Soc. 1910, 42. 570.)

Sp. gr. of (UO₂)(NO₃)₂+Aq at t°.

t°	% salt	Sp. gr.
11.5	1	1.0049
12.4	2	1.0096
15.1	3	1.01401
14.1	4	1.0187
16.7	5	1.0230
14.1	6	1.8281
15.7	7	1.0236
15.2	8	1.0378
16.5	9	1.0410
15.2	10	1.0462
13.7	11	1.0504
11.5	12	1.0550
14.5	13	1.0594
11.3	14	1.0643
12.5	15	1.0680
13.2	16	1.0718

(de Coninck, C. R. 1900, 131. 1219.)

Sp. gr. of a sat. aq. solution = 1.7536 at 17°, containing 54.77% UO₂(NO₃)₂.
Sp. gr. = 1.0257 when 2.80% UO₂(NO₃)₂ present. (Vassiliev, C. C. 1912. I, 1430.)

Sp. gr. of solution in HNO₃+Aq.

Pts. of salt in 100 pts. HNO ₃ sp. gr. 1.153	1	2	3	4	5
emp.	11.0°	11.8°	11.3°	12.0°	11.6°
p. gr. of the solution	1.1585	1.1614	1.1663	1.1698	1.1751

(de Coninck, C. R. 1901, 132. 90.)

Sp. gr. of solution in H₂SO₄+Aq.

Pts. of salt in 100 pts. H ₂ SO ₄ sp. gr. 1.138	1 pt.	2 pts.	3 pts.	4 pts.	5 pts.
Temp.	11.2°	11.8°	10.7°	12.0°	11.4°
Sp. gr. of solution	1.1427	1.1450	1.1511	1.1540	1.1576

(de Coninck.)

Very sol. in dil. HBr and selenic acid (d=1.4). Sol. in conc. H₂SO₄, HNO₃, dil. HCl and less sol. in conc. HCl. (de Coninck, C. R. 1900, 131. 1220.)

Sp. gr. of solution in HBr+Aq of sp. gr. 1.21.

Sp. gr.	% salt dissolved
1.2122	1
1.2168	2
1.2198	3
1.2250	4
1.2305	5

(de Coninck, Belg. Acad. Bull. 1901. 222.)

Insol. in KOH+Aq, NaOH+Aq or NH₄OH+Aq. Sol. in lime water. (de Coninck, C. R. 1900, 131. 1220.)

Sl. attacked by liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 830.)

At 15°, uranyl nitrate is sol. in comm. methyl alcohol, dil. and conc. ethyl alcohol, propyl and isobutyl alcohol, comm. amyl alcohol, acetone, ether, ethyl acetate, dil. and conc. formic acid and dil. acetic acid; sl. sol. in comm. essence of terebenthine; insol. in benzene (cryst.), comm. toluene and xylene, ligroin, CHCl₃, glycerine and CS₂. (de Coninck, C. R. 1900, 131. 1220.)

1 pt. is sol. in 55 pts. methyl alcohol at ca. 11.8°.

1 pt. is sol. in 30 pts. ethyl alcohol (85°) at ca. 12.9°.

1 pt. is sol. in 65 pts. acetone at ca. 12.0°.

1 pt. is sol. in 5.6 pts. acetic acid (d = 1.035) at ca. 14.25°.

(de Coninck, C. R. 1900, 131. 1304.)

1 pt. sol. in 23.5 pts. methyl alcohol at 11.2°.

1 " " " 16.0 " ether " 11.9°.

1 " " " 18.4 " ethyl acetate " 10.3°.

1 " " " 5.3 " conc. formic acid at 15.1°.

(de Coninck, C. R. 1901, 132. 91.)

Sp. gr. of solution in comm. methyl alcohol at t°.

t°	% salt	Sp. gr. referred to H ₂ O
11	1	0.8902
12.9	2	0.8938
12.2	3	0.9003
10.7	4	0.9068
12.8	5	0.9108

(de Coninck, C. R. 1900, 131. 1304.)

Sp. gr. of solution in ethyl alcohol (85°) at t°.

d₁ = sp. gr. referred to alcohol.

d₂ = sp. gr. referred to H₂O.

t°	% salt	d ₁	d ₂
11.9	1	0.8918	1.0060
12.2	2	0.8979	1.0127
11.6	3	0.9023	1.0177
13.1	4	0.9056	1.0227
11.7	5	0.9131	1.0280

(de Coninck, C. R. 1900, 131. 1219.)

Sp. gr. of solution in acetic acid (d = 1.055) at t°.

d₁ = sp. gr. referred to H₂O.

d₂ = sp. gr. referred to acetic acid.

t°	% salt	d ₁	d ₂
14.0	1	1.0387	1.0034
13.8	2	1.0434	1.0080
14.8	3	1.0469	1.0100
16.9	4	1.0505	1.0148
14.6	5	1.0564	1.0205
10.4	6	1.0626	1.0265
11.7	7	1.0662	1.0300

(de Coninck, C. R. 1900, 131. 1304.)

When excess of UO₂(NO₃)₂ is shaken with ether at 7°, two layers are formed, the ether layer containing 59 g. salt per 100 g. solution and the aqueous layer 62.5 g. salt per 100 g. solution. (Lebeau, C. R. 1911, 152. 440.)

Sol. in nearly all proportions in glycerine. (Postans, Pharm. J. 1883, (3) 13. 752.)

Sol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Sol. in acetone. (Eidmann, C. C. 1899. II, 1014; Naumann, B. 1904, 37. 4328.)

+18H₂O. Sat. aq. solution has D_{17°/17°} = 1.7536. (Vasilieff, J. Russ. phys. Chem. Soc. 1911, 43. 1183.)

Uranyl nitrate ammonia, (UO₂)(NO₃)₂, 2NH₃.

(v. Unruh, Dissert. 1909.)

(UO₂)(NO₃)₂, 3NH₃. (v. Unruh.)

(UO₂)(NO₃)₂, 4NH₃. (v. Unruh.)

Uranyl nitrate phosphate, UO₂H₄(PO₄)₂, UO₂(NO₃)₂ + 14H₂O.

Easily sol. in warm H₂O, with gradual decomp. Easily sol. in HNO₃, HCl, or H₂SO₄ + Aq. Sol. in acetic acid with decomp. (Heints, A. 151. 216.)

Divanadyl nitrate (?).

Known only in solution. Decomp. on evaporation.

Ytterbium nitrate, basic.

Easily sol. in H₂O.

Ytterbium nitrate.

Very sol. in H₂O.

+3H₂O. Ppt. (Cleve, Z. anorg. 1 32. 140.)

+4H₂O. (Cleve.)

Yttrium nitrate, basic, 2Y₂O₃, 3N₂O₅ + 4H₂O.

Deliquescent in moist air. Decomp. cold or boiling H₂O. Sol. in a solution of yttrium nitrate without decomp. (Behr Bunsen, A. 137. 1.)

Yttrium nitrate, Y(NO₃)₃ + 6H₂O.

Easily sol. in H₂O, alcohol, or (Cleve.)

141.6 grams are sol. in 100 grams H₂O at 25°. (James, J. Am. Chem. Soc. 1911 876.)

Zinc nitrate, basic, 8ZnO, N₂O₅ + 2H₂O.

Insol. in H₂O (Grouvelle, A. ch. 12.

6ZnO, N₂O₅ + 8H₂O = Zn(NO₃)₂, 5ZnO + 3H₂O. (Bertels, J. B. 1784. 274.)

5ZnO, N₂O₅ + 5½H₂O. Insol. in somewhat sol. in hot H₂O. (Haven

+6H₂O. Slowly decomp. by cold (Rousseau and Tite.)

9ZnO, 2N₂O₅. Decomp. by H₂O. (and Reischauer, N. Jahrb. Pharm. 11.

4ZnO, N₂O₅ + 2H₂O. (Schindler.)

+3H₂O. (Ordway, Sill. Am. J. (2) 3 Gerhardt, J. Pharm. (3) 12. 61.)

Insol. in H₂O; sol. in dil. acids. (A asasco, Bull. Soc. 1896, 15. 1080.)

2ZnO, N₂O₅ + 3H₂O. Decomp. by H₂O slowly by alcohol. (Wells, Am. Ch. J. 2

7ZnO, 4N₂O₅ + 14H₂O = 4Zn(NO₃)₂, 3Zn(OH)₂ + 11H₂O. (Bertels.)

Zinc nitrate, Zn(NO₃)₂.

Very deliquescent. Easily sol. in H₂O and alcohol.

Sp. gr. of Zn(NO₃)₂ + Aq. F. = accord Franz (J. pr. (2) 6. 274) at 17.5°; O. = according to Oudemans (Z. anal. 7. 410) at 14°:

	5	10	15%Zn(N	
F.	1.0496	1.0604	1.1476	
O.	1.0425	1.067	1.1355	
	20	25	30%ZnO	
F.	1.2024	1.2040	1.2208	
O.	1.1875	1.245	1.206	
	35	40	45	50%ZnO
F.	1.906	1.4572	1.5258	1.5984
O.	1.1875	1.245	1.206	1.206

Calculated for Zn(NO₃)₂ + 6H₂O:

	10	20	30	40	50%
F.	1.05361	1.1131	1.1782	1.2496	1.3

(Oudemans.)

Zn(NO₃)₂ + Aq when heated soon decomposes, with formation of an insol. base (Ordway.)

of $\text{Zn}(\text{NO}_3)_2 + \text{Aq}$ at room temp.
:
30.626 44.5% $\text{Zn}(\text{NO}_3)_2$.
1.2291 1.4367
gner, W. Ann. 1883, 18. 270.)
gr. of $\text{Zn}(\text{NO}_3)_2 + \text{Aq}$ at 25°.

ration of $\text{Zn}(\text{NO}_3)_2 + \text{Aq}$	Sp. gr.
ormal	1.0758
"	1.0404
"	1.0191
"	1.0096

gner, Z. phys. Ch. 1890, 5. 40.)
of $\text{Zn}(\text{NO}_3)_2 + \text{Aq}$ at 16°.

$\text{Zn}(\text{NO}_3)_2$ g. per 1000 solution	Sp. gr. 16°/16°
0000	1.000000
9950	1.000814
0061	1.001646
1535	1.003413
1824	1.006733
7760	1.014702
5920	1.028890
6780	1.058644

en, Z. phys. Ch. 1897, 24. 108.)
of $\text{Zn}(\text{NO}_3)_2 + \text{Aq}$ at 17.3°, when p =
strength of solution; d = observed
w = volume conc. in grams per cc.
)

	d	w
	1.5504	0.73310
	1.4579	0.60240
	1.3136	0.40535
	1.2933	0.37780
	1.1830	0.23246
	1.1284	0.16232
	1.0988	0.12478
	1.0597	0.07515
	1.0491	0.06213
	1.0118	0.01593
	1.0087	0.01221

es, J. Phys. Chem. 1898, 2. 545.)
asily sol. in liquid NH_3 . (Franklin,
J. 1898, 20. 830.)
 H_2O . 100 pts. HNO_3 dissolve 28
3°; 55 pts. at 55°. (Ditte, A. ch.
18. 335.)
ity in H_2O .
ution contains at:
0° 41° 43° 45.5° mpt.
.42 68.21 69.26 77.77% $\text{Zn}(\text{NO}_3)_2$.
unk, Z. anorg. 1899, 20. 401.)

+6 H_2O .
Solubility in H_2O .
Sat. solution contains at:
-18° -15° -13° -12°
44.63 45.26 45.51 45.75% $\text{Zn}(\text{NO}_3)_2$,
0° +12.5° 18° 25°
48.66 52.00 53.50 55.90% $\text{Zn}(\text{NO}_3)_2$,
36.4° (mpt.) 36° 33.5°
63.63 64.73 65.83% $\text{Zn}(\text{NO}_3)_2$.
(Funk, Z. anorg. 1899, 20. 400.)

100 g. $\text{Zn}(\text{NO}_3)_2 + \text{Aq}$ sat. at 0° contain
48.7 g. $\text{Zn}(\text{NO}_3)_2$; at 18°, 53.5 g. $\text{Zn}(\text{NO}_3)_2$.
(Mylius, Z. anorg. 1910, 74. 411.)
Melts in its crystal H_2O at 36.4° (Ordway),
50° (Pierre); boils at 131° (Ordway).
Sp. gr. of solution sat. at 18° = 1.664, and
contains 53.9% $\text{Zn}(\text{NO}_3)_2$. (Mylius, B.
1897, 30. 1718.)
Sol. in methyl acetate. (Naumann, B.
1909, 42. 3790); ethyl acetate. (Naumann,
B. 1910, 43. 314.)
+9 H_2O .
Solubility in H_2O .
Sat. solution contains at:
-25° -22.5° -20° -18°
40.12 40.75 42.03 43.59% $\text{Zn}(\text{NO}_3)_2$.
Cryohydrate is formed at -29°.
(Funk, Z. anorg. 1899, 20. 401.)

Zinc nitrate ammonia, $\text{Zn}(\text{NO}_3)_2 \cdot 4\text{NH}_3$.
Ppt. (Ephraim, B. 1915, 48. 638.)
+ $\frac{2}{3}\text{H}_2\text{O}$.
Deliquescent. Sol. in H_2O . (André, C. R.
100. 639.)
13 ZnO , 3 N_2O_5 , 2 $\text{NH}_3 + 18\text{H}_2\text{O}$.
Insol. in cold, decomp. by warm H_2O .
(André, C. R. 1885, 100. 640.)

Zinc nitrate cupric oxide,
 $\text{Zn}(\text{NO}_3)_2 \cdot 3\text{CuO} + 3\text{H}_2\text{O}$.
(Mailhe, A. ch. 1902, (7) 27. 169.)

Zinc nitrate hydrazine, $\text{Zn}(\text{NO}_3)_2 \cdot 3\text{N}_2\text{H}_4$.
Decomp. by hot H_2O .
Sol. in NH_4OH . (Franzen, Z. anorg.
1908, 60. 279.)

Zirconium nitrate, basic, 3 ZrO_2 , 2 N_2O_5 .
Insol. in H_2O .
 ZrO_2 , N_2O_5 . Easily sol. in H_2O and alcohol.
+ H_2O . As above.

Zirconium nitrate, $\text{Zr}(\text{NO}_3)_4 + 5\text{H}_2\text{O}$ (?).
Deliquescent, and sol. in H_2O .

Nitric oxide, NO .
See Nitrogen dioxide.

Nitrilobromosmic acid.

Ammonium nitrilopentabromosmate,
 $[\text{OsNBr}_5](\text{NH}_4)_2 + \text{H}_2\text{O}.$

Very sol. in H_2O . Decomp. in dil. aq. solution.

Stable in $\text{HBr} + \text{Aq}.$

Insol. in organic solvents. (Werner, B. 1906, 39. 501.)

Cæsium hydrogen nitrilopentabromosmate,
 $[\text{OsNBr}_5]_2\text{Cs}_2\text{H}.$

Sl. sol. in H_2O . (Werner.)

Potassium nitrilotetrabromosmate,
 $[\text{OsNBr}_4]\text{K} + 2\text{H}_2\text{O}.$

Very sol. in H_2O . Decomp. in aq. solution. Stable in $\text{HBr} + \text{Aq}.$ Insol. in organic solvents. (Werner.)

Rubidium nitrilopentabromosmate,
 $[\text{OsNBr}_5]\text{Rb}_2.$

Sol. in H_2O . Decomp. in dil. aq. solution after a short time. (Werner.)

Nitrilochlorosmic acid.

Ammonium nitrilopentachlorosmate,
 $(\text{OsNCl}_5)(\text{NH}_4)_2.$

Sol. in H_2O ; insol. in conc. $\text{HCl} + \text{Aq}.$ (Werner, B. 1901, 34. 2702.)

Cæsium nitrilopentachlorosmate,
 $(\text{OsNCl}_5)\text{Cs}_2.$

Sol. in H_2O . (Werner.)

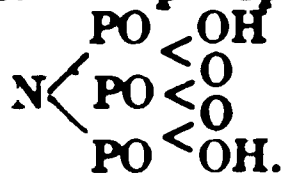
Potassium nitrilopentachlorosmate,
 $(\text{OsNCl}_5)\text{K}_2.$

Sol. in H_2O ; pptd. by HCl ; insol. in organic solvents. (Werner.)

Rubidium nitrilopentachlorosmate,
 $(\text{OsNCl}_5)\text{Rb}_2.$

Sol. in H_2O ; decomp. in dil. neutral solution. (Werner.)

Nitrilotrimetaphosphoric acid, $\text{H}_2\text{NP}_3\text{O}_7 =$



Known only in solution. (Mente, A. 248. 260.)

Aluminum nitrilotrimetaphosphate.

Insol. in H_2O , conc. HCl , or $\text{HNO}_3 + \text{Aq}.$ Slowly sol. in boiling conc. H_2SO_4 . Sol. in warm $\text{NaOH} + \text{Aq}$ or $\text{Na}_2\text{CO}_3 + \text{Aq}$ without decomp. Insol. in $\text{NH}_4\text{OH} + \text{Aq}.$ (Mente.)

Barium —, $\text{BaNP}_3\text{O}_7.$

Insol. in dil. or conc. acids. Decomp. by boiling NaOH or $\text{Na}_2\text{CO}_3 + \text{Aq}.$ Insol. in $\text{NH}_4\text{OH} + \text{Aq}.$ (Mente.)

Cadmium nitrilotrimetaphosphate.

Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}.$ or bcd $(\text{NH}_4)_2\text{CO}_3$, or $\text{NaOH} + \text{Aq}.$ (Mente.)

Calcium —, $\text{CaNP}_3\text{O}_7 + \text{H}_2\text{O}.$

Sol. in conc. $\text{HCl} + \text{Aq}$ by long boiling, more easily in fuming $\text{HNO}_3 + \text{Aq}.$ Ins. NH_4OH or $\text{NaOH} + \text{Aq}.$ (Mente.)

Chromium —.

Slowly sol. in dil. acids. Easily s ammonia. Sol. in cold $\text{NaOH} + \text{Aq}.$ (M)

Cobalt —, $\text{CoNP}_3\text{O}_7 + \text{H}_2\text{O}.$

Insol. in H_2O . Sl. sol. in dil. acids. l sol. in $\text{NH}_4\text{OH} + \text{Aq}.$ Decomp. by Na $\text{Na}_2\text{CO}_3 + \text{Aq}.$ (Mente.)

Copper —.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}.$ Decomp. by ! $+ \text{Aq}.$ (Mente.)

Ferric —, $\text{Fe}_2(\text{NP}_3\text{O}_7)_3.$

Insol. in conc. acids. Easily sol. in N $+ \text{Aq}$ or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}.$ Decomp NaOH or $\text{Na}_2\text{CO}_3 + \text{Aq}.$ (Mente.)

Lead —.

Insol. in dil. acids. Sol. in fuming ! Insol. in $\text{NH}_4\text{OH} + \text{Aq}.$ Sol. in NaOH (Mente.)

Magnesium —, $\text{MgNP}_3\text{O}_7 + \text{H}_2\text{O}.$

Slowly sol. in $\text{HCl} + \text{Aq}.$ Sol. in H_2 fuming HNO_3 with addition of Br_2 in NH_4OH or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}.$ (Men)

Manganous —, $\text{MnNP}_3\text{O}_7 + \text{H}_2\text{O}.$

Insol. in dil. acids. Very sl. sol. in N $\text{Aq}.$ Insol. in Na_2CO_3 or $(\text{NH}_4)_2\text{CO}_3$ Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}.$ (Mente.)

Mercurous —, $\text{Hg}_2\text{NP}_3\text{O}_7.$

Insol. in dil. acids, NH_4OH , Na $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}.$ Easily sol. in HNO_3 . (Mente.)

Nickel —, $\text{NiNP}_3\text{O}_7 + \text{H}_2\text{O}.$

Insol. in dil. acids, NH_4OH , or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}.$ (Mente.)

Zinc —, $\text{ZnNP}_3\text{O}_7 + 2\text{H}_2\text{O}.$

Easily sol. in NH_4OH , NaOH $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}.$ (Mente.)

Nitrilosulphonic acid, $\text{N}(\text{SO}_3\text{H})_2.$

Not known in free state. (Raschig, 161.)

Potassium nitrilosulphonate, $\text{N}(\text{SO}_3\text{K})_2 + 2\text{H}_2\text{O}.$

Soluble in H_2O . (Raschig, A. 241. Is identical with "potassium aminephosphate" of Claus.

1 cold H_2O (Claus); sol. in 50 pts. H_2O (Fremy); in H_2O at scarcely 40° range. Decomp. by boiling. (Claus.)

sodium nitrilosulphonate, $(K)_2(SO_3Na)$.

insol. in cold H_2O . (Raschig, A.

trilosulphonate, $N(SO_3Na)_3$.

lated on account of its extreme in H_2O . (Raschig, A. 241. 161.)

sulphophosphoric acid, H_3 .

by H_2O . (Stock, B. 1906, 39.

n nitrilodisulphophosphate, NH_4 .

ol. in H_2O . Not decomp. by boiling i. Decomp. by acid. (Stock.)

ol. in liquid NH_3 . (Stock, B. 1903,

n hydrogen nitriloisulphophos-, $SHP(SNH_4)N$.

omp. by boiling with alkali. De- acids. (Stock, B. 1906, 39. 1999.)

trilodisulphophosphate, $BaNPS_2 +$

H_2O with decomp. Decomp. by Not decomp. by warming with decomp. by acid. (Stock.)

odisulphophosphate, NPS_2Pb .

liquid NH_3 . Solution decomp. th separation of PbS . (Stock.)

trilodisulphophosphate, $NPSNa_2$.

omp. by boiling with alkali. De- acid. (Stock.)

phuric acid.

n nitrilosulphate, $N(SO_3NH_4)_3 +$

al. sol. in H_2O , but much more sol. lt. (Divers and Haga, Chem. Soc. 094.)

trilosulphate, $N(SO_3Na)_3 + 5H_2O$.

ol. in H_2O . (Divers and Haga, 1901, 79. 1097.)

phurous acid.

n nitrilosulphite, $NH(SO_3NH_4)_2$.

at deliquescent. Very sol. in H_2O . comp. in solution. Decomp. by h HCl . (Divers, Proc. Chem. Soc. 63.)

Nitritocobaltic chloride.

Sol. in 200 pts. cold H_2O . (Jørgensen, Z. anorg. 5. 172.)

Nitritoplatindiamine nitrate, $(NO_2)_2Pt(N_2H_4NO_2)_2$.

Sol. in cold H_2O with decomp.; violently decomp. on warming. (Hadow, Chem. Soc. (2) 4. 345.)

Nitritopurplecobaltic comps.

See **Xanthocobaltic comps.**

Nitritopurpleorhodium comps.

See **Xanthorhodium comps.**

Nitrocarbamic acid.

Potassium nitrocarbamate, $NO_2.NK$. COOK.

Decomp. by H_2O . (Thiele, B. 1894, 27. 1909.)

Nitro cobalt, Co_2NO_2 .

Decomp. by H_2O . (Sabatier and Senderens, C. R. 115. 236.)

Nitro copper, $CuNO_2$.

Violently decomp. by H_2O . (Sabatier and Senderens, C. R. 116. 756.)

Nitroferricyanhydric acid.

See **Nitroprussic acid**.

Nitrogen, N_2 .

Nearly insol. in all known solvents.

1 vol. recently boiled H_2O absorbs 0.0147 vol. N at 15.5° . (Henry, 1893.)

1 vol. recently boiled H_2O absorbs 0.025 vol. N. (Dalton.)

1 vol. recently boiled H_2O absorbs 0.0156 vol. N at ord. temp. (Dalton.)

1 vol. H_2O at t° and 760 mm. absorbs V vols. N gas reduced to 0° and 760 mm.

t°	V	t°	V	t°	V
0	0.02035	7	0.01713	14	0.01500
1	0.01981	8	0.01675	15	0.01478
2	0.01932	9	0.01640	16	0.01458
3	0.01884	10	0.01607	17	0.01441
4	0.01838	11	0.01577	18	0.01426
5	0.01794	12	0.01549	19	0.01413
6	0.01752	13	0.01523	20	0.01403

(Bunsen.)

Coefficient of absorption = $0.020346 - 0.00053887t + 0.000011156t^2$. (Bunsen.)

1 l. H₂O absorbs ccm. N from atmospheric air at 760 mm. pressure and t°.

t°	ccm. N	t°	ccm. N
0	19.29	15	13.95
5	17.09	20	12.80
10	15.36	25	11.81

(Dittmar, Challenger Exped. Report, vol. i.)

t°	ccm. N	t°	ccm. N
0	19.14	15	13.73
5	16.93	20	12.63
10	15.14	25	11.80

(Hamberg, 1885.)

Absorption of N by H₂O at t° and 760 mm.
β = coefficient of absorption.

t°	β	t°	β	t°	β
0	0.02388	18	0.01696	36	0.01252
1	2337	19	1667	37	1233
2	2288	20	1639	38	1215
3	2241	21	1611	39	1198
4	2196	22	1584	40	1182
5	2153	23	1557	41	1166
6	2111	24	1530	42	1151
7	2070	25	1504	43	1137
8	2031	26	1478	44	1124
9	1993	27	1453	45	1111
10	1956	28	1428	46	1099
11	1920	29	1404	47	1088
12	1885	30	1380	48	1078
13	1851	31	1357	49	1069
14	1818	32	1334	50	1061
15	1786	33	1312	60	1000
16	1755	34	1291	100	1000
17	1725	35	1271

(Bohr and Bock, W. Ann. 44. 318.)

Absorption of N by H₂O at t° and 760 mm.
β = coefficient of absorption; β₁ = "Solubility" (see under Oxygen).

t°	β	β ₁
0	0.02348	0.02334
1	2291	2276
2	2236	2220
3	2182	2166
4	2130	2113
5	2081	2063
6	2032	2013
7	1986	1966
8	1941	1920
9	1898	1877
10	1857	1834
11	1819	1795
12	1782	1758
13	1747	1722
14	1714	1687

Absorption of N by H₂O at t°.—Continued

t°	β	A
15	0.1682	0 1654
16	1651	1632
17	1622	1591
18	1594	1562
19	1567	1534
20	1542	1507
21	1519	1482
22	1496	1457
23	1473	1433
24	1452	1410
25	1432	1387
26	1411	1365
27	1392	1344
28	1374	1323
29	1356	1303
30	1340	1284
31	1321	1263
32	1304	1243
33	1287	1224
34	1270	1204
35	1254	1185
36	1239	1167
37	1224	1149
38	1210	1131
39	1196	1114
40	1183	1097
41	1171	1082
42	1160	1067
43	1149	1052
44	1139	1037
45	1129	1023
46	1120	1009
47	1111	0995
48	1102	0982
49	1094	0968
50	1087	0955
52	1072	0929
54	1058	0902
56	1045	0876
58	1033	0849
60	1022	0822
62	1011	0794
64	1001	0765
66	0992	0736
68	0983	0707
70	0976	0678
72	0970	0645
74	0965	0614
76	0961	0581
78	0959	0546
80	0957	0510
82	0956	0473
84	0955	0433
86	0954	0398
88	0953	0343
90	0952	0294
92	0951	0242
94	0950	0187
96	0949	0128
98	0948	0066
100	0947	0000

(Winkler, B. 24. 3606.)

efficient of absorption for $H_2O = 0.01432$ at 0° ; 0.01621 at 20° ; 0.01789 at 15° ; 0.01973 at 10° ; 0.02173 at 5° . (Braun, Z. Ch. 1900, 33, 730.)

solubility in H_2O at various pressures.

volume of the absorbing liquid.

Hg-pressure in metres.

coefficient of solubility.

V	t°	P	λ
1 ccm.	19.4	0.8010	0.01617
		1.0453	0.01616
		1.2488	0.01611
		1.4764	0.01608
		1.8111	0.01602
		2.3961	0.01597
		2.9074	0.01585
		3.3411	0.01579
		4.1382	0.01561
		5.0059	0.01554
		5.1103	0.01546
		6.1110	0.01528
		6.2767	0.01515
		7.1059	0.01499
		7.5815	0.01487
		8.1074	0.01473
2 ccm.	24.9	0.8977	0.01498
		1.0129	0.01493
		1.1887	0.01491
		1.5573	0.01487
		1.9846	0.01482
		2.5171	0.01478
		2.8781	0.01473
		3.2956	0.01455
		4.0947	0.01440
		4.5581	0.01434
		5.0529	0.01426
		5.5935	0.01413
		6.1956	0.01406
		7.0333	0.01382
		7.5596	0.01377
		7.4. 1846	0.01369

Cassuto, Phys. Zeit. 1904, 5, 236.)

efficient of absorption for $H_2O = 0.01565$ at 18° . (Hufner, Z. phys. Ch. 1907, 57.)

Absorption of N_2 by distilled H_2O at t° .
a = ccm. of N_2 absorbed by 1 l. of H_2O at t° and 760 mm.

t°	a	t°	a	t°	a
0	23.00	17	16.29	34	12.93
1	22.50	18	16.03	35	12.79
2	22.02	19	15.78	36	12.65
3	21.55	20	15.54	37	12.52
4	21.09	21	15.29	38	12.39
5	20.64	22	15.06	39	12.27
6	20.20	23	14.84	40	12.15
7	19.77	24	14.63	41	12.04
8	19.35	25	14.43	42	11.92
9	18.94	26	14.23	43	11.80
10	18.54	27	14.04	44	11.68
11	18.16	28	13.87	45	11.57
12	17.80	29	13.71	46	11.46
13	17.46	30	13.55	47	11.35
14	17.14	31	13.39	48	11.24
15	16.84	32	13.23	49	11.13
16	16.56	33	13.08	50	11.02

(Fox, Trans. Faraday Soc. 1909, 5, 73.)

Solubility in H_2O at $25^\circ = 0.1561$. (Drucker and Moles, Z. phys. Ch. 1910, 75, 418.)

Solubility of N_2 in H_2O at $25^\circ = 0.0231$. (Calculated according to special formula, for which see original article.) (Findlay and Creighton, Chem. Soc. 1911, 99, 1315.)

Coefficient of absorption for $H_2O = 0.01689$ at 15° ; 0.01670 at 16.2° ; 0.01622 at 17.2° . (Müller, Z. phys. Ch. 1912, 81, 493.)

1 l. sea water (sp. gr. 1.027) absorbs ccm. N_2 from atmosphere at t° and 760 mm. pressure—

t°	According to Törnøe	According to Dittmar	According to Hamberg
0	14.40	15.60	14.85
5	13.25	13.86	13.32
10	12.10	12.47	12.06
15	10.95	11.34	11.04
20	9.80	10.41	10.25
25	8.65	9.62	9.62

No. of ccm. of N_2 (containing 1.185% argon) absorbed by a l. of sea-water from a free dry atmosphere of 760 mm. pressure at given temperatures.

Cl per 1000	0°	4°	8°	12°	16°	20°	24°	28°
0	18.64	17.02	15.63	14.45	13.45	12.59	11.86	11.25
4	17.77	16.27	14.98	13.88	12.94	12.15	11.46	10.89
8	16.90	15.51	14.32	13.30	12.44	11.70	11.07	10.52
12	16.03	14.75	13.66	12.72	11.93	11.25	10.67	10.16
16	15.18	14.00	13.00	12.15	11.73	10.81	10.27	9.80
20	14.31	13.24	12.34	11.57	10.92	10.36	9.87	9.44

(Fox, Trans. Faraday Soc. 1909, 5, 71.)

Absorption of N_2 by $H_2SO_4 + Aq$ at t° . α = coefficient of absorption.

Normality of the acid	t°	α
0	20.9	0.0156
4.9	20.9	0.0091
8.9	20.9	0.0072
10.7	21.2	0.0066
20.3	21.1	0.0049
24.8	21.5	0.0048
29.6	20.8	0.0051
34.3	20.9	0.0100
35.8	21.1	0.0129

(Bohr, Z. phys. Ch. 1910, 71. 49.)

Absorption of N_2 by $BaCl_2 + Aq$. α_{t° = coefficient of absorption at t° .

Per cent of $BaCl_2$ in the solution	α_{25°	α_{20°	α_{15°	α_{10°	α_5°
13.830	0.00783	0.00923	0.01036	0.01166	0.01270
11.927	0.00855	0.00978	0.01139	0.01249	0.01368
6.903	0.01044	0.01184	0.01317	0.01474	0.01598
6.738	0.01036	0.01182	0.01340	0.01494	0.01628
3.870	0.01137	0.01323	0.01460	0.01660	0.01802
3.325	0.01190	0.01346	0.01502	0.01681	0.01826

(Braun, Z. phys. Ch. 1900, 33. 733.)

Absorption of N_2 by $NaCl + Aq$. α_{t° = coefficient of absorption at t° .

Per cent of $NaCl$ in the solution	α_{25°	α_{20°	α_{15°	α_{10°	α_5°
11.732	0.00470	0.00657	0.00810	0.00930	0.01016
10.945	0.00545	0.00703	0.00824	0.00912	0.01052
8.135	0.00749	0.00872	0.01014	0.01131	0.01260
8.033	0.00720	0.00871	0.00995	0.01121	0.01248
6.595	0.00802	0.00872	0.01120	0.01252	0.01380
6.400	0.00826	0.00975	0.01134	0.01259	0.01375
4.196	0.00900	0.01151	0.01294	0.01451	0.01579
3.800	0.01005	0.01198	0.01316	0.01475	0.01615
2.120	0.01131	0.01311	0.01460	0.01638	0.01795
2.100	0.01133	0.01314	0.01467	0.01650	0.01805
0.886	0.01295	0.01477	0.01640	0.01838	0.01994
0.671	0.01304	0.01484	0.01642	0.01845	0.02000

(Braun, *l. c.*)

At -191.5° liquid oxygen dissolves 458 times its vol. or 50.7 per cent of its weight of gaseous nitrogen. (Erdmann, B. 1904, 37. 1191.)

At 18° and 760 mm. 100 vols. H_2O or alcohol of 0.84 sp. gr. absorb 4.2 vols. N gas. (de Saussure, 1814.)

1 vol. alcohol at t° and 760 mm. dissolve vols. N gas reduced to 0° and 760 mm.

t°	V	t°	V
0	0.12634	13	0.1219
1	0.12593	14	0.1216
2	0.12553	15	0.1214
3	0.12514	16	0.1211
4	0.12476	17	0.1208
5	0.12440	18	0.1205
6	0.12405	19	0.1203
7	0.12371	20	0.1203
8	0.12338	21	0.1202
9	0.12306	22	0.1200
10	0.12276	23	0.1199
11	0.12247	24	0.1197
12	0.12219

(Bunsen's Gasometry.)

1 vol. alcohol absorbs 0.126338–0.00041 0.000060 t° vols. N gas. (Carius, A. 84. 1)

Solubility in alcohol at 25° .

Vol. H_2O	Vol. % alcohol	Solubility
100	0	0.0163
80	20	0.0153
67	33	0.0171
0	100	0.1432

(Just, Z. phys. Ch. 1901, 37. 361.)

1 vol. ether absorbs 0.15 vol. N (Döbereiner). caoutchouc absorbs 5 vols. N in 5 weeks (Humb.)

Solubility of N_2 in ether = 0.2580 at 0.2561 at 10° . (Christoff, Z. phys. Ch. 179. 459.)

Solubility in organic solvents

Solvent	Solubility at $25^\circ C$	Solubility at $20^\circ C$	
Glycerine	Not measurable		
Water	0.01634	0.01703	–0
Aniline	0.03074	0.02992	+0
Carbon bisulphide	0.05860	0.05290	+0
Nitrobenzene	0.06255	0.06052	+0
Benzene	0.1159	0.114	+0
Glacial acetic acid	0.1190	0.1172	+0
Xylene	0.1217	0.1185	+0
Amyl alcohol	0.1225	0.1206	+0
Toluene	0.1235	0.1186	+0
Chloroform	0.1348	0.1282	+0
Methyl alcohol	0.1415	0.1346	+0
Ethyl alcohol (99.8%)	0.1432	0.1400	+0
Acetone	0.1460	0.1382	+0
Amyl acetate	0.1542	0.1512	+0
Ethyl acetate	0.1727	0.1676	+0
Isobutyl acetate	0.1734	0.1701	+0

(Just, Z. phys. Ch. 1901, 37. 361.)

Coefficient of absorption for petrol 0.117 at 20° ; 0.135, at 10° . (Gnielwa Walsz, Z. phys. Ch. 1. 70.)

ion of N_2 by propionic acid + Aq.
coefficient of absorption at t° .

α_{25}°	α_{20}°	α_{15}°	α_{10}°	α_5°
0.01301	0.01483	0.01593	0.01779	0.01951
0.01295	0.01447	0.01585	0.01800	0.01977
0.01336	0.01471	0.01634	0.01823	0.02040
0.01329	0.01469	0.01630	0.01845	0.02026
0.01335	0.01476	0.01637	0.01855	0.02077
0.01338	0.01480	0.01648	0.01872	0.02089
0.01365	0.01541	0.01688	0.01919	0.02098
0.01371	0.01547	0.01674	0.01915	0.02087

n, Z. phys. Ch. 1900, 33. 732.)

of N_2 in isobutyric acid + Aq at t° .
ected pressure at end of experi-
m. Hg at 0° .
bility of N_2 .

at	t°	P	S
utyric	25.05	262.6	0.1609(?)
		388.3	0.1640
		566.1	0.1647
		662.4	0.1656
		783.5	0.1656
		832.2	0.1656
rtion yric l sure = m.	23.02	246.2	0.0393
		492.2	0.0393
		563.6	0.0393
		836.3	0.0400
		867.3	0.0401
sure = m.	29.02	231	0.0373
		468.4	0.0384
		480.7	0.0383
		536	0.0385
		656	0.0384
		720	0.0386

nd Moles, Z. phys. Ch. 1910, 76.
434.)

on of N_2 by chloralhydrate + Aq.
of the solution.
loralhydrate in the solution.
ficient of absorption at t° .
efficient of absorption at 15° .

P	β_{t°	β_{15°
15.8	0.01574	0.01580
28.2	0.01418	0.01422
37.25	0.01288	0.01300
47.0	0.01260	0.01275
56.52	0.01230	0.01245
71.5	0.01415	0.01420
78.8	0.01447	0.01495

r, Z. phys. Ch. 1912, 81. 499.)

Absorption of N_2 by organic substances + Aq
at 15° .

P = % of the organic substance in the sol-
vent.

β_{15° = coefficient of absorption at 15° .

S $_{15^\circ}$ = Solubility at 15° .

Organic substance used	P	β_{15°	S $_{15^\circ}$
Chloralhydrate	0	0.01725	...
	0	0.01675	0.01796
	0	0.01706	...
	6.9	0.0164	0.0173
	14.0	0.0154	0.0162
	15.0	0.0152	0.0160
	23.6	0.0134	0.0141
	26.1	0.0141	0.0149
	37.6	0.0123	0.0130
	48.9	0.0115	0.0121
	49.3	0.0118	0.0124
	61.3	0.0114	0.0120
	70.9	0.0131	0.0138
	71.2	0.0130	0.0137
	78.3	0.0152	0.0160
	79.1	0.0156	0.0165
Glycerine	0	0.01707	...
	0	0.01708	...
	15.7	0.01425	...
	15.7	0.01376	...
	29.9	0.01087	...
	46.6	0.00840	...
	57.6	0.00698	...
	67.1	0.00635	...
	72.8	0.00552	...
	74.7	0.00597	...
	77.0	0.00527	...
	85.1	0.00482	...
	87.3	0.00492	...
	88.5	0.00536	...
	99.25	0.00524	...

(Hammel, Z. phys. Ch. 1915, 90. 121.)

Absorption of N_2 by glycerine + Aq.

t° = temp. of the solution.

P = % glycerine in the solution.

β_{t° = coefficient of absorption at t° .

β_{15° = coefficient of absorption at 15° .

t°	P	β_{t°	β_{15°
16.1	25.0	0.01240	0.01286
15.6	42.2	0.00966	0.00976
14.7	51.5	0.00759	0.00759
14.9	58.0	0.00703	0.00703
15.9	80.25	0.00520	0.00530
16.2	90.0	0.00570	0.00583
18.0	95.0	0.00578	0.00716

(Müller, Z. phys. Ch. 1912, 81. 496.)

Solubility of N₂ in glycerine + Aq at 25°.
G = % by wt. of glycerine in the solvent.
S = solubility of N₂.
P = corrected pressure at end of experiment in mm. Hg at 0°.

G	P	S
16	598.4	0.0103
"	915.5	0.0103
29.7	556.5	0.0067
"	846.5	0.0068
48.9	617.7	0.0052
"	859.8	0.0051
74.5	588.5	0.0025
84.1	637.3	0.0024
"	757.0	0.0024

(Drucker and Moles, Z. phys. Ch. 1910, 75. 418.)

Absorption of N₂ by sucrose + Aq.
t° = temp. of the solution.
P = % sucrose in the solution.
β_{t°} = coefficient of absorption at t°.
β_{15°} = coefficient of absorption at 15°.

t°	P	β _{t°}	β _{15°}
16.2		0.01670	0.01700
17.2		0.01622	0.01688
16.8	11.38	0.01432	0.01480
16.9	20.00	0.01233	0.01280
17	29.93	0.01025	0.01053
17.8	30.12	0.01033	0.01090
18	47.89	0.00742	0.00785
17.7	48.57	0.00658	0.00700

(Müller, Z. phys. Ch. 1912, 81. 493.)

Absorption of N₂ by organic substances + Aq at t°.
V = absorbed volume reduced to 0° and 760 mm.
α = coefficient of absorption.

Solution	Vol. of solution ccm.	t°	V ccm.	α
N-dextrose	409.94	20.18	4.55	0.01215
1/2 N-dextrose	409.94	20.21	5.14	0.01380
1/4 N-dextrose	409.94	20.2	5.51	0.01480
N-levulose	409.94	20.25	4.27	0.01221
N-arabinose	409.94	20.21	4.40	0.01203
N-erythritol	409.94	20.25	4.87	0.01321
N-alanine	409.94	20.19	4.445	0.01213
N-glycocoll	409.94	20.16	4.47	0.01212
N-urea	409.94	20.18	5.37	0.01477
N-acetamide	409.94	20.22	5.385	0.01475

(Hüfner, Z. phys. Ch. 1907, 57. 618-621.)

Nitrogen bromide, NBr₃.
Decomp. under H₂O.

Nitrogen bromophosphide, PBr₃N.
Insol. in H₂O. Sol. in ether, less in CHCl₃. (Besson, C. R. 114. 147)

Nitrogen bromosulphide.
See Nitrogen sulphobromide.

Nitrogen chloride, NCl₃.
Very unstable. Explodes when 93° or by contact with other s
Insol. in H₂O, but is decomp. then hours by cold H₂O). Sol. in CS₂, S₂Cl₂. (H. Davy, Phil. Trans. 181
Sol. in C₆H₆, CS₂, CHCl₃, CCl₄. (B. 1897, 30. 1434.)

Nitrogen chlorophosphide, N₂P₂Cl₄.
Insol. in H₂O, but slowly decomp
Insol. in hot H₂SO₄, HCl, or H
Decomp. by hot fuming HNO₃.
alcohol; very sol. in ether, but these gradually decompose. Sol. in CS
C₆H₆, and oil of turpentine.
Sol. in POCl₃. (Gladstone, Che 138.)

Nitrogen chlorosulphide.
See Nitrogen sulphochloride.

Nitrogen fluoride.
Very explosive. (Warren, C. N.

Nitrogen monoiodamine, NH₂I.
Very rapidly decomp. by H₂O in (Raschig, A. 230. 212.)

Nitrogen diiodamine, NHI₂.
Properties as triioddiamine.

Nitrogen triioddiamine, NH₂, NI₃.
Decomp. by H₂O. (Raschig, A. Insol. in absolute alcohol. Sol comp. in HCl + Aq. (Bunsen.)

Nitrogen iodide, N₂I.
See Triazoiiodide.

Nitrogen iodide, NI₃.
Insol. in H₂O, but slowly decomp
Sol. in HCl + Aq. Sol. in KCN + lon, J. pr. 17. 1.)
Sol. in Na₂S₂O₃ + Aq. (Guyard, 526.)
Sol. in KSCN + Aq. (Raschig, A

Nitrogen iodide ammonia, NI₃, 2 2NH₃; and NI₃, NH₃.
(Hugot, C. R. 1900, 130. 507.)
NI₃, 12NH₃. Ppt.; insol. in eth (B. 1900, 33. 3028.)

gen monoxide, N₂O.

Liquid. Miscible with alcohol or ether.
Gas.

l. H₂O absorbs 0.78–0.86 vol. N₂O at ordinary (Henry); 0.80 vol. at ordinary temp. (Dalton); d. at ordinary temp. (de Saussure); 0.708 vol. at eisch); 0.54 vol. (Davy).

H₂O at t° and 760 mm. absorbs V vols. N₂O, reduced to 0° and 760 mm.

V	t°	V
1.3052	13	0.8304
1.2605	14	0.8034
1.2172	15	0.7778
1.1752	16	0.7535
1.1346	17	0.7306
1.0954	18	0.7090
1.0575	19	0.6888
1.0210	20	0.6700
0.9858	21	0.6525
0.9520	22	0.6364
0.9196	23	0.6216
0.8885	24	0.6082
0.8588

(Bunsen's Gasometry.)

ol. H₂O absorbs 1.30521–0.0453620t + 8430t² vols. N₂O at t° and 760 mm. en.)

fficient of absorption by H₂O = 0.01883 . (Steiner, Z. phys. Ch. 1895, 18. 14.)

fficient of absorption by H₂O = 0.600 5°; 0.773 at 15.5°; 0.951 at 8.1°. (Gor. phys. Ch. 1895, 18. 4.)

Absorption of N₂O by H₂O at t°.

t°	Coefficient of absorption
25	0.5752
20	0.6654
15	0.7896
10	0.9479
5	1.1403

Roth, Z. phys. Ch. 1897, 24. 123.)

ibility in H₂O at 25° = 0.5942; at 20° = ; at 15° = 0.7784; at 10° = 0.9101; at 067. (For formula for "solubility," der oxygen.) (Geffcken, Z. phys. Ch. 19. 278.)

ibility of N₂O in H₂O = 0.592 at 25° and 762 mm. pressure. (Findlay and ton, Chem. Soc. 1910, 97. 538.)

vols. H₂SO₄ (sp. gr. = 1.84) absorb 75.7 N₂O; 100 vols. H₂SO₄ + Aq (sp. gr. = absorb 66.0 vols. N₂O; 100 vols. H₂SO₄ (sp. gr. = 1.705) absorb 39.1 vols. N₂O; 100 vols. H₂SO₄ + Aq (sp. gr. = 1.45) absorb 33.0 vols. N₂O.

l₂ + Aq, and NaCl + Aq absorb con- le amounts of N₂O. (Lunge, B. 14.

Absorption by acids + Aq.

M = content in gram-equivalents per liter.

S = solubility (see under Oxygen).

Absorption of N₂O by HNO₃ + Aq.

M	S 25°	S 15°
0.610	0.5969	0.7770
0.614	0.5980	0.7766
1.253	0.6045	0.7767
1.254	0.6061	0.7767
2.405	0.6156	0.7735
2.435	0.6149	0.7737

(Geffcken, Z. phys. Ch. 1904, 49. 278.)

Absorption of N₂O by HCl + Aq.

M	S 25°	S 15°
0.549	0.5775	0.7550
0.550	0.5759	0.7528
1.089	0.5670	0.7360
1.093	0.5657	0.7347
2.300	0.5546	0.7103
2.340	0.5564	0.7122

(Geffcken.)

Absorption of N₂O by $\frac{H_2SO_4}{2}$ + Aq.

M	S 25°	S 15°
0.523	0.5648	0.7328
0.526	0.5657	0.7340
1.050	0.5426	0.6997
1.054	0.5419	0.6984
2.042	0.5083	0.6440
2.047	0.5087	0.6428
2.971	0.4819	0.6024
2.963	0.4820	0.6030
3.897	0.4569	0.5648
3.973	0.4577	0.5640

(Geffcken.)

Absorption of N₂O by H₃PO₄ + Aq at t°.

t°	% of H ₃ PO ₄				
	3.38%	4.72%	8.84%	9.89%	13.35%
5	1.057	1.0365	0.9883	0.9635	0.9171
10	0.8827	0.8665	0.8296	0.8101	0.7711
15	0.7388	0.7258	0.6977	0.6826	0.6505
20	0.6253	0.6147	0.5926	0.5810	0.5555
25	0.5427	0.5329	0.5143	0.5054	0.4860

(Roth, Z. phys. Ch. 1897, 24. 134.)

100 vols. conc. FeSO₄ + Aq absorb 19.5 vols. N₂O.

Solubility of N₂O in a solution containing 47.7 g. Fe(OH)₃ per litre at 25° = 0.5799; 47.9 g. Fe(OH)₃ per litre at 25° = 0.5787. (Geffcken, Z. phys. Ch. 1904, 49. 299.)

100 vols. KOH+Aq (sp. gr. = 1.12) absorb 18.7 vols. N_2O ; 100 vols. KOH+Aq sat. with pyrogallol absorb 18.1 vols. N_2O ; 100 vols. NaOH+Aq (sp. gr. = 1.1) (7% NaOH) absorb 23.1 vols. N_2O ; 100 vols. NaOH+Aq sat. with pyrogallol absorb 28.0 vols. N_2O .

Absorption of N_2O by KOH+Aq.

M = content in gram-equivalents per litre.

S = solubility (see under oxygen).

M	S 25°	S 15°
0.541	0.5087	0.6591
0.542	0.5093	0.6595
1.074	0.4252	0.5427
1.082	0.4221	0.5392

(Geffcken, Z. phys. Ch. 1904, 49, 278.)

Coefficient of solubility of N_2O in salts+Aq at t° .

Salt	Concentration of salt		Coeff. of absorption at			
	G per 100 g. solution	G mol. per l	5°	10°	15°	20°
CaCl ₂	5.79	0.547	0.819	0.697	0.591	0.500
	9.86	0.964	0.608	0.586	0.509	0.435
	13.99	1.416	0.510	0.441	0.380	0.328
LiCl	1.35	0.319	0.986	0.831	0.700	0.599
	3.85	0.928	0.878	0.743	0.629	0.536
	11.48	2.883	0.606	0.512	0.437	0.382
Li ₂ SO ₄	2.37	0.219	0.934	0.792	0.670	0.569
	5.46	0.521	0.795	0.665	0.557	0.474
	8.56	0.836	0.646	0.555	0.477	0.415
MgSO ₄	5.90	0.521	0.766	0.664	0.561	0.471
	7.66	0.687	0.708	0.586	0.486	0.414
	10.78	0.997	0.569	0.491	0.417	0.346
KCl	4.90	0.676	0.879	0.751	0.643	0.555
	7.64	1.037	0.799	0.693	0.591	0.494
	14.58	2.187	0.654	0.574	0.500	0.430
	22.08	3.114	0.544	0.459	0.390	0.339
K ₂ SO ₄	2.62	0.154	0.986	0.831	0.701	0.605
	4.78	0.285	0.918	0.763	0.637	0.542
NaCl	6.20	1.107	0.800	0.682	0.585	0.509
	8.88	1.614	0.713	0.603	0.510	0.434
	12.78	2.391	0.634	0.532	0.449	0.386
Na ₂ SO ₄	5.76	0.427	0.808	0.677	0.584	0.495
	8.53	0.646	0.692	0.574	0.482	0.416
	12.44	0.974	0.559	0.486	0.417	0.354
SrCl ₂	3.31	0.215	0.928	0.788	0.671	0.578
	5.73	0.380	0.848	0.709	0.610	0.556
	13.24	0.939	0.644	0.547	0.463	0.390

(Gordon, Z. phys. Ch. 1895, 18, 5.)

Absorption of N_2O by salts+Aq at 15°.
M = number of molecules of salt per litre
s = coefficient of absorption.

Salt	M	s
KCl	3.554	0.0802
	2.909	0.1012
	1.755	0.1279
	1.051	0.1489
	0.526	0.1667
KNO ₃	2.430	0.1180
	1.820	0.1311
	1.541	0.1391
	0.879	0.1559
	0.482	0.1683
K ₂ CO ₃	4.352	0.0160
	2.939	0.0288
	2.156	0.0462
	1.376	0.0761
	0.690	0.1183
	0.341	0.1501
	0.209	0.1628
NaCl	4.815	0.0606
	2.801	0.0925
	2.049	0.1130
	0.825	0.1548
NaNO ₃	5.711	0.0578
	3.980	0.0610
	2.656	0.1052
	1.413	0.1370
Na ₂ CO ₃	0.679	0.1603
Na ₂ SO ₄	1.218	0.0639
	0.819	0.1082
	0.438	0.1383
	0.207	0.1639
Na ₂ SO ₄	1.364	0.0775
	0.638	0.1254
	0.335	0.1519
LiCl	3.734	0.0990
	1.800	
	0.835	0.1619
MgSO ₄	2.501	0.0499
	1.631	0.0797
	0.936	0.1159
	0.433	0.1501
ZnSO ₄	2.180	0.0605
	1.277	
	0.899	0.1173
	0.397	0.1525
CaCl ₂	2.962	0.0519
	2.556	0.0519
	1.827	0.0839
	1.122	0.1138
	0.578	0.1450
	0.321	0.1619

(Steiner, Z. phys. Ch. 1895, 18, 14-6.)

ent of absorption of N₂O by NaCl+
Aq at t°.

Per cent of NaCl			
0.990	1.808	3.886	5.865
1.0609	1.0032	0.9131	0.8428
0.8812	0.8383	0.7699	0.7090
0.7339	0.7026	0.6495	0.5976
0.6191	0.5962	0.5520	0.5088
0.5363	0.5190	0.4775	0.4424

oth, Z. phys. Ch. 1897, 24. 139.)

rpption of N₂O by salts+Aq at 20°.
ncentration of the solution in terms
al.
efficient of absorption.
ption of N₂O by KNO₃+Aq at 20°.

	C	a
163	0.1061	0.6270
720	0.2764	0.6173
389	0.5630	0.6002
577	1.1683	0.5713
		0.5196

otion of N₂O by NaNO₃+Aq at 20°.

	C	a
24	0.1336	0.6270
31	0.3052	0.6089
77	0.6286	0.5876
01	1.1200	0.5465
		0.4926

ropp, Z. phys. Ch. 1904, 48. 107.)

bsorption of N₂O by salts+Aq.
ontent in gram-equivalents per litre.
lubility.

	M	S 25°	S 15°
1	0.598	0.5532	0.7203
	0.600	0.5504	0.7185
	1.158	0.5223	0.6800
	1.166	0.5200	0.6775
	0.550	0.5367	0.6950
	0.557	0.5344	0.6916
	0.886	0.5025	0.6466
	0.913	0.5012	0.6442
	0.514	0.5428	0.7074
	0.545	0.5406	0.7036
	0.558	0.5276	0.6884
	0.561	0.5278	0.6877
	1.057	0.4760	0.6163
	1.059	0.4773	0.6146

Absorption of N₂O by salts+Aq.—
Continued.

Salt	M	S 25°	S 15°
KBr	0.546	0.5306	0.6877
	0.550	0.5318	0.6892
	0.937	0.4908	0.6352
	0.959	0.4899	0.6334
RbCl	0.439	0.5399	0.7050
	0.444	0.5386	0.7053
	0.977	0.4873	0.6306
	0.993	0.4846	0.6276
	0.558	0.5218	0.6782
	0.559	0.5217	0.6787
	1.070	0.4673	0.6046
	1.102	0.4639	0.6020

(Geffcken, Z. phys. Ch. 1904, 49. 278.)

Solubility of N₂O in a solution containing
39.6 g. As₂S₃ per litre at 25°=0.5819; 42.4 g.
As₂S₃ per litre at 25°=0.5833. (Geffcken.)

1 vol. alcohol at t° and 760 mm. absorbs V
vols. N₂O gas reduced to 0° and 760 mm.

t°	V	t°	V
0	4.1780	13	3.3734
1	4.1088	14	3.3200
2	4.0409	15	3.2678
3	3.9741	16	3.2169
4	3.9085	17	3.1672
5	3.8442	18	3.1187
6	3.7811	19	3.0714
7	3.7192	20	3.0253
8	3.6585	21	2.9805
9	3.5990	22	2.9368
10	3.5408	23	2.8944
11	3.4838	24	2.8532
12	3.4279

(Bunsen's Gasometry.)

Coefficient of absorption=4.17805-
0.0698160t+0.0006090t². (Carius.)

At 18° and 760 mm., 100 vols. H₂O absorb 76 vols.
N₂O; 100 vols. alcohol of 0.840 sp. gr. absorb 153 vols.;
100 vols. rectified naphtha of 0.784 sp. gr. absorb 254
vols.; 100 vols. oil of lavender of 0.880 sp. gr. absorb
275 vols.; 100 vols. olive oil of 0.915 sp. gr. absorb 150
vols.; 100 vols. sat. KCl+Aq (26% KCl) of 1.212 sp.
gr. absorb 29 vols. (de Saussure, 1814.)

1 vol. oil of turpentine absorbs 2.5-2.7 vols. N₂O.
(de Saussure.)

Absorption of N₂O by glycerine+Aq at t°.

t°	% by weight of glycerine			
	3.460%	6.726%	12.120%	16.244%
25	0.5558	0.5415	0.5268	0.5083
20	0.6468	0.6303	0.6050	0.5851
15	0.7672	0.7454	0.7098	0.6857
10	0.9172	0.8871	0.8411	0.8102
5	1.0967	1.0552	0.9990	0.9586

(Roth, Z. phys. Ch. 1897, 24. 128.)

Absorption of N ₂ O by urea + Aq at t°.					
t°	% by weight of urea				
	3.312%	4.974%	6.366%	7.296%	9.966%
25	0.5686	0.5669	0.5588	0.7502	0.5689
20	0.6533	0.6558	0.6539	0.6553	0.6508
15	0.7708	0.7732	0.7605	0.7722	0.7614
10	0.9209	0.9201	0.9086	0.9208	0.9007
5	1.1040	1.0964	1.0880	1.1012	1.0685

(Roth, Z. phys. Ch. 1897, 24. 124.)

Absorption of N ₂ O by sugar + Aq at 15°.	
Number of molecules of C ₁₂ H ₂₂ O ₁₁ per litre	Coefficient of absorption
1.699	0.0892
0.993	0.1284
0.520	0.1561

(Steiner, Z. phys. Ch. 1895, 18. 15.)

Absorption of N₂O by organic substances + Aq.
C = concentration of the solution in terms of normal.
a = coefficient of absorption.
Absorption of N₂O by chloral hydrate + Aq at 20°.

P	C	a
0	...	0.6270
2.947	0.184	0.6182
6.848	0.445	0.6128
13.48	0.942	0.5960
16.15	1.165	0.5891
19.60	1.474	0.5793
24.02	1.911	0.5675

(Knopp, Z. phys. Ch. 1904, 48. 106.)

P	C	a
0	...	0.6270
1.492	0.2045	0.6323
5.702	0.816	0.6369
13.680	2.140	0.6504
15.011	2.385	0.6534
25.589	4.645	0.7219

(Knopp, Z. phys. Ch. 1904, 48. 107.)

Absorption of N ₂ O by oxalic acid + Aq at t°		
t°	Coeff. of abs. in H ₂ C ₂ O ₄ + Aq of given % strength	
	8.122%	3.699%
25	0.5786	0.5643
20	0.6694	0.6538
15	0.7940	0.7745
10	0.9526	0.9264
5	1.1450	1.1004

(Roth, Z. phys. Ch. 1897, 24. 130.)

Coefficient of absorption for petroleum 2.11 at 20°; 2.49 at 10°. (Gniewas Walfisz, Z. phys. Ch. 1. 70.)
The solubility of N₂O in various collo solutions has been determined by Fm and Creighton (Chem. Soc. 1910, 97. 538) which see original article.

Nitrogen dioxide, NO.

1 vol. H₂O absorbs 0.1 vol. NO gas at ordinary t (Davy); 1 vol. absorbs 0.05 vol. (Henry); 1 vol ab 1/27 vol. (Dalton.)
Absorption of NO by H₂O at 760 mm pressure.
β = Coefficient of absorption.
β' = "Solubility."

t°	β	β'	t°	β	β'
0	0.07381	0.07337	55	0.03040	0.0
5	6461	6406	60	2954	
10	5709	5640	65	2877	
15	5147	5061	70	2810	
20	4706	4599	75	2751	
25	4323	4189	80	2700	
30	4004	3838	85	2665	
35	3734	3529	90	2648	
40	3507	3254	95	2638	
45	3311	3000	100	2628	
50	3152	2771			

(Winkler, B. 1901, 34. 1414.)

205.69 cc. H₂O absorb 9.6798 cc. NO a and 760 mm. (Hufner, Z. phys. Ch. 1 59. 420.)
Sol. in conc. HNO₃ + Aq.
100 vols. HNO₃ + Aq of 1.3 sp. gr. agi with NO gas take up 20 vols. NO. If a twice as strong or one-half as strong, quantity NO is proportional to the am of HNO₃. Very dil. HNO₃ + Aq ab scarcely more NO than pure H₂O. (Dal
100 pts. HN(O)₃ + Aq of 1.4 sp. gr. absorb 90 pt (Dalton); sol. in Br₂ and very sl. sol. in conc. l (Berthelot.)
1 ccm. conc. H₂SO₄ of 1.84 sp. gr. ab 0.035 ccm. NO; of 1.50 sp. gr., 0.017 NO. (Lunge, B. 18. 1391.)

of NO by $\text{H}_2\text{SO}_4 + \text{Aq}$ at 18° and 760 mm.

cient of solubility.

α	H_2SO_4	α
not constant	70%	0.0113
0.0193	60%	0.0118
0.0117	50%	0.0120

er, Z. anorg. 1906, 50. 387.)

in aqueous solutions of ferrous ally the sulphate. (Priestley.) $\text{SO}_4 + \text{Aq}$ of 1.081 sp. gr., contain- FeSO_4 to 6 grains H_2O , absorbs 6 (Dalton.)
n by ferrous salts + Aq is propor- e amount of Fe present, irrespec- acid or concentration of the solu- een 0° and 10° , about 2 mols. NO d for each atom of Fe; between , 1 mol. NO for 2 atoms of Fe; only 1 mol. NO for $2\frac{1}{2}$ to 3 atoms e amount of NO absorbed also the pressure. The sp. gr. of the solution is greater after the ab- NO than before. The solutions . by heat, and at 100° all NO is (Gay, A. ch. (6) 5. 145.)

n of NO by $\text{FeSO}_4 + \text{Aq}$ at 25° .
 I_2O (in litres) containing 1 mol.
IO (in litres) absorbed.

V	A	V
1.47	7.2	5.52
2.01	12.0	6.46
2.55	18.6	8.01
4.40	36.0	10.40

schütter, B. 1907, 40. 877.)

on of NO by $\text{FeSO}_4 + \text{Aq}$ at t° .
 $\text{FeSO}_4 + \text{Aq}$ contain 0.0221 g. Fe.
t of absorption = 0.06067 at 20.09° .

Pressure mm.	NO absorbed ccm.
704.9	14.42
683.5	14.10
668.6	13.80
651.9	13.58
632.9	13.15
613.7	12.98

Absorption of NO by $\text{FeSO}_4 + \text{Aq}$ at t° .—
Continued.

205.69 cc. of $\text{FeSO}_4 + \text{Aq}$ contain 0.0296 g. Fe.
Coefficient of absorption = 0.06505.

t°	Pressure mm.	NO absorbed ccm.
20.05	677.5	14.30
20.05	655.3	14.07
20.04	639.1	13.81
20.00	620.2	13.39
20.15	600.5	13.20
20.14	581.2	12.92

205.69 cc. of $\text{FeSO}_4 + \text{Aq}$ contain 0.0409 g. Fe.
Coefficient of absorption = 0.06684.

t°	Pressure mm.	NO absorbed ccm.
20.04	667.6	16.79
20.02	650.6	16.65
20.00	613.1	15.71
20.00	594.6	15.41
20.10	577.1	15.32

205.69 cc. of $\text{FeSO}_4 + \text{Aq}$ contain 0.0513 g. Fe.
Coefficient of absorption = 0.07981.

t°	Pressure mm.	NO absorbed ccm.
20.10	644.8	18.82
20.10	623.8	18.47
20.08	606.4	18.02
20.10	589.7	17.56
20.10	571.1	17.19
20.10	553.1	16.95

205.69 cc. of $\text{FeSO}_4 + \text{Aq}$ contain 0.0663 g. Fe.
Coefficient of absorption = 0.08059.

t°	Pressure mm.	NO absorbed ccm.
20.10	697.3	21.91
20.10	678.9	21.60
20.10	660.4	21.18
20.08	638.2	20.71
20.04	620.7	20.28
20.00	602.5	19.87

205.69 cc. of $\text{FeSO}_4 + \text{Aq}$ contain 0.099 g. Fe.
Coefficient of absorption = 0.11661.

t°	Pressure mm.	NO absorbed ccm.
20.10	649.9	34.26
20.15	631.1	33.82
20.20	618.4	33.26
20.00	603.3	32.76
19.85	588.6	32.34
19.85	574.2	31.95

(Hüfner, Z. phys. Ch. 1907, 59. 419.)

Absorption of NO by $\text{NiSO}_4 + \text{Aq}$ at t° .
205.69 cc. $\text{NiSO}_4 + \text{Aq}$ contain 0.0506 g. Ni.
Coefficient of absorption = 0.08311.

t°	Pressure mm.	NO absorbed ccm.
20.2	654.7	23.00
20.2	629.8	22.54
20.2	609.5	22.03
20.15	591.7	21.65
20.14	573.4	21.18

(Hüfner, *l. c.*)

Absorption of NO by $\text{CoSO}_4 + \text{Aq}$ at t° .
205.69 cc. $\text{CoSO}_4 + \text{Aq}$ contain 0.0598 g. Co.
Coefficient of absorption = 0.09146.

t°	Pressure mm.	NO absorbed ccm.
20.15	678.3	23.47
20.16	653.5	23.01
20.20	636.6	22.55
20.30	615.9	21.99
20.40	600.0	21.56

(Hüfner, *l. c.*)

Absorption of NO by $\text{MnCl}_2 \cdot 4\text{H}_2\text{O} + \text{Aq}$ at t° .
205.69 cc. $\text{MnCl}_2 \cdot 4\text{H}_2\text{O} + \text{Aq}$ contain 0.0697 g. Mn.
Coefficient of absorption = 0.06111.

t°	Pressure mm.	NO absorbed ccm.
20.0	711.96	14.25
20.05	686.5	13.99
20.2	657.4	13.49
20.3	638.9	13.05
20.45	621.0	12.81

(Hüfner, *l. c.*)

Coefficient of absorption for $\text{FeSO}_4 + \text{Aq}$ of concentration used by Hüfner (*Z. phys. Ch.* 1907, 59. 417) = 0.180 at 20° . Hüfner's results are incorrect because he assumed that the absorption-coefficient of NO always had the same value, whereas it does not. NO is reduced by $\text{FeSO}_4 + \text{Aq}$. (Usher, *Z. phys. Ch.* 1908, 62. 624.)
Coefficient of absorption for $\text{CoSO}_4 + \text{Aq}$ sat. at 20° = 0.0288. (Usher, *Z. phys. Ch.* 1908, 62. 624.)
Coefficient of absorption for $\text{NiSO}_4 + \text{Aq}$ of the concentration used by Hüfner (*cf. Z. phys. Ch.* 1907, 59. 422) = 0.048 at 20° .
Coefficient of absorption for $\text{NiSO}_4 + \text{Aq}$ sat. at 20° = 0.0245. (Usher, *l. c.*)
Coefficient of absorption for $\text{MnCl}_2 + \text{Aq}$ sat. at 20° = 0.0082. (Usher, *Z. phys. Ch.* 1908, 62. 624.)

Absorption of NO by $\text{FeCl}_2 + \text{Aq}$ at 20° .
 A = vol. H_2O (in litres) containing 1 FeCl_2 .
 V = vol. NO (in litres) absorbed.

A	V
2.5	3.30
5.18	4.83
10.35	6.56
20.7	8.32
51.8	11.89

(Kohlschütter, *B.* 1907, 40. 878.)

Absorption by $\text{HCl} + \text{FeCl}_2 + \text{Aq}$
10.37 l. 30% HCl containing 1 mol. in solution absorb 15.64 l. NO.
10.37 l. 10% HCl containing 1 mol. in solution absorb 6.17 l. NO.
(Kohlschütter, *l. c.*)
Absorption by salts + $\text{FeCl}_2 + \text{Aq}$
10.37 l. sat. $\text{NaCl} + \text{Aq}$ containing FeCl_2 in solution absorb 6.549 l. NO.
10.37 l. sat. $\text{NH}_4\text{Cl} + \text{Aq}$ containing FeCl_2 in solution absorb 6.549 l. NO.
(Kohlschütter, *l. c.*)

Solubility of NO in $\text{Fe}(\text{NO}_3)_3 + \text{Aq}$ at 20° .
 A = vol. H_2O (in litres) containing $\text{Fe}(\text{NO}_3)_3$.
 V = vol. NO (in litres) absorbed.

A	V
3.25	2.77
6.50	4.16
13.00	5.54
26.00	6.61

(Kohlschütter, *l. c.*)

Absorption of NO by $\text{CuCl}_2 + \text{Aq}$ at 20° .
 A = vol. H_2O (in litres) containing CuCl_2 .
 V = vol. NO (in litres) absorbed.

A	V
0.231	0.120
0.277	0.098
0.371	0.052

(Kohlschütter, *l. c.*)

Absorption of NO by $\text{CuCl}_2 + \text{conc. HCl}$ at 20° .
 A = vol. conc. HCl (in litres) containing 1 mol. CuCl_2 .
 V = vol. NO (in litres) absorbed.

A	V	A	V
0.389	0.801	7.499	3.
0.410	0.933	12.500	3.
0.840	2.838	18.750	3.
1.230	3.426	28.650	1.
2.462	3.989		

(Kohlschütter, *l. c.*)

on of NO by CuCl₂+acetic acid.
acetic acid (in litres) containing 1
NO (in litres) absorbed.

	V
52	51.77
14	39.67
19	81.60

(Kohlschütter, l. c.)

on of NO by CuCl₂+98% formic
acid.
98% formic acid (in litres) contain-
CuCl₂.
NO (in litres) absorbed.

	V
9	12.76
10	13.17
10	14.34
10	18.68
10	27.29

(Kohlschütter, l. c.)

ion of NO by CuCl₂+acetone.
acetone (in litres) containing 1
NO (in litres) absorbed.

V	A	V
14.04	291.60	40.99
24.01	583.20	67.22
24.60	1166.40	81.96

(Kohlschütter, l. c.)

of NO by CuCl₂+methyl alcohol.
methyl alcohol (in litres) contain-
CuCl₂.
NO (in litres) absorbed.

V	A	V
3.30	20.50	6.15
5.60	82.25	4.90

(Kohlschütter, l. c.)

i of NO by CuCl₂+ethyl alcohol.
ethyl alcohol (in litres) containing
l₂.
NO (in litres) absorbed.

V	A	V
8.70	38.41	18.15
12.38	76.83	18.05
15.43	192.10	15.92

(Kohlschütter, l. c.)

Absorption of NO by CuBr₂+Aq.
A=vol. H₂O (in litres) containing 1 mol.
CuBr₂.
V=vol. NO (in litres) absorbed.

A	V
0.37	0.515
0.62	0.120
0.925	0.000

(Kohlschütter, l. c.)

Absorption of NO by CuBr₂+ethyl alcohol.
A=vol. alcohol (in litres) containing 1 mol.
CuBr₂.
V=vol. NO (in litres) absorbed.

A	V	A	V
2.625	16.02	131.20	22.23
5.25	19.26	262.50	23.46
13.12	20.51	656.10	30.46
43.74	21.13		

(Kohlschütter, l. c.)

Sol. in stannous and chromous salts+Aq.
(Peligot.)
Not absorbed by Fe₂(SO₄)₃+Aq. (Dalton.)
1 vol. absolute alcohol absorbs 0.31606-
0.003487t+0.000049t² vols. NO between 0°
and 25°. (Bunsen.)

1 vol. alcohol at t° and 760 mm. absorbs V
vols. NO gas reduced to 0° and 760 mm.

t°	V	t°	V
0	0.31606	13	0.27901
1	0.31262	14	0.27685
2	0.30928	15	0.27478
3	0.30604	16	0.27281
4	0.30290	17	0.27094
5	0.29985	18	0.26917
6	0.29690	19	0.26750
7	0.29405	20	0.26592
8	0.29130	21	0.26444
9	0.28865	22	0.26306
10	0.28609	23	0.26178
11	0.28363	24	0.26060
12	0.28127

(Bunsen's Gasometry.)

Abundantly absorbed by CS₂. (Friedburg,
C. N. 48. 97.)

Nitrogen trioxide, N₂O₃.

Sol. in H₂O at 0°. If large amt. of H₂O is
present, the solution is quite stable at ordi-
nary temp. (Fremy, C. R. 79. 61.)

Sol. in HNO₃+Aq.
Sol. in conc. H₂SO₄ to form HNOSO₃.
Sol. in ether.

Nitrogen trioxide stannic chloride, N_2O_3 , $SnCl_4$.

Decomp. by H_2O . (Weber, Pogg. 118. 471.)

Nitrogen tetroxide, NO_2 or N_2O_4 .

Sol. in H_2O at 0° with decomp. Miscible with very conc. HNO_3 . Absorbed abundantly by CS_2 , $CHCl_3$, and C_6H_6Cl . (Friedburg, C. N. 47. 52.)

Sol. in $C_6H_5NO_2$.

Sl. sol. in $H_2S + Aq$.

Sol. in H_2SO_4 or conc. $HNO_3 + Aq$.

H_3PO_4 absorbs some liquid NO_2 . (Frankland, Chem. Soc. 1901, 79. 1362.)

Nitrogen pentoxide, N_2O_5 .

Very deliquescent. Combines with H_2O to form HNO_3 with evolution of heat.

Nitrogen hexoxide, NO_3 .

Decomposes upon air or with H_2O . (Hautefeuille and Chappins, C. R. 92. 80, 134; 94. 1111, 1306.)

Nitrogen oxybromide.

See Nitrosyl and Nitroxyl bromide.

Nitrogen oxychloride.

See Nitrosyl and Nitroxyl chloride.

Nitrogen oxyfluoride.

See Nitrosyl fluoride and Nitroxyl fluoride.

Nitrogen phosphochloride, $P_2N_2Cl_4$.

See Nitrogen chlorophosphide.

Nitrogen selenide, NSe .

Very explosive. Insol. in H_2O . Sol. in $HNO_3 + Aq$, and $NaClO + Aq$. (Espenschied, A. 113. 101.)

Insol. in H_2O , ether, absolute alcohol; very sl. sol. in CS_2 , C_6H_6 , and glacial acetic acid. Decomp. by HCl or $KOH + Aq$. (Verneuil, Bull. Soc. (2) 38. 548.)

Nitrogen sulphide, N_4S_4 .

Insol. in H_2O . Decomp. by hot H_2O . Sl. sol. in alcohol, ether, wood alcohol, oil of turpentine. Easily sol. in CS_2 . Slowly decomp. by $HCl + Aq$ or $KOH + Aq$, rapidly by $HNO_3 + Aq$. 15 g. dissolve in 1 kilo. of CS_2 . (Fordos and Gélis, C. R. 31. 702.)

Sol. in $CHCl_3$. (Demarçay, C. R. 91. 854.)

Sol. in warm glacial acetic acid with decomp. on boiling. (Ruff and Geisel, B. 1904, 37. 1591.)

Nitrogen pentasulphide, N_2S_5 .

Sol. in ether and most organic solvents; insol. in H_2O ; fairly stable in ethereal solution, but decomp. by light. (Muthmann, Z. anorg. 1897, 13. 206.)

Nitrogen sulphobromide, N_2S_4Br .

Decomp. by boiling H_2O and by dil. al also by boiling with alcohol. (Muth B. 1897, 30. 630.)

$N_2S_4Br_4$. Decomp. by moist air. (C. B. 1896, 29. 340-341.)

$N_4S_4Br_8$. Decomp. by moist air. unstable. (Clever.)

$N_4S_4Br_2$. Insol. most solvents; unstable. (Clever.)

Nitrogen sulphochloride, $N_2S_4Cl_4$.

Unstable on air. Sol. in warm C crystallizes out on cooling. (Demarçay, 91. 854, 1066.)

Demarçay calls this comp. thiazyl chl

Sol. in hot dry benzene, and in decomp. by moist air. (Andreocci, Z. 1897, 14. 249.)

$N_2S_4Cl_2$. Partly sol. in H_2O . (Demarçay, C. R. 92. 726.)

Demarçay calls this compound dithiazyl dichloride.

$N_2S_4Cl_2 = N_2S_2, SCl_2$. Decomp. on air. (Fordos and Gélis.)

Demarçay (C. R. 92. 726) calls this thiodithiazyl dichloride.

$N_2S_4Cl_2$. Sol. in H_2O with subsequent decomp. More sol. than S in CS_2 . (Soubeiran, A. ch. 67. 71.)

Is a mixture of S_2Cl_2 and N_2S_4 . (Fordos and Gélis, C. R. 31. 702.)

$N_2S_4Cl_2$. Sl. sol. in warm, insol. in $CHCl_3$. (Demarçay, C. R. 92. 726.)

"Thiothiazyl chloride." (Demarçay.)

$N_2S_4Cl_2$. Sol. in H_2O . Insol. in most solvents. Sl. sol. in $CHCl_3$. Easily converted to thionyl chloride. (Demarçay, C. R. 91. 1066.)

Demarçay calls the compound thiazyl chloride $= (NS)_2 \equiv S-Cl$.

$N_4S_4Cl_2 = 2N_2S_2, SCl_2$. Decomp. on air. (Michaelis.)

$N_6S_6Cl_2 = 3N_2S_2, SCl_2$. Not decomposed by H_2O containing ammonia.

Nitrogen sulphotiodide, N_2S_4I .

Readily decomp. by H_2O . (Muth and Seitter, B. 1897, 30. 627.)

Nitrohydroxylaminic acid, $H_2N_2O_4$.

Known only in solution. (Angeli, Gazz. chim. it. 1897, 27 (2) 357.)

Barium nitrohydroxylamine, $Ba.N_2O_4$.

Ppt. More stable in the air than the sodium salt. Not decomp. by prolonged boiling with H_2O . (Angeli, Gazz. chim. it. 1896, 26. 17-25.)

Cadmium nitrohydroxylamine, $Cd.N_2O_4$.

As Ba salt. (Angelico and Fanara, Gazz. chim. it. 1901, 31. (2) 21.)

nitrohydroxylamine, $\text{CaN}_2\text{O}_3 + \text{H}_2\text{O}$.
Gazz. ch. it. 1900, 30. (1) 593.)

nitrohydroxylamine, $\text{CaN}_2\text{O}_3 + \text{H}_2\text{O}$.
O and Fanara, Gazz. ch. it. 1901, 31. (2) 15.)

hydroxylamine, PbN_2O_3 .
Gazz. ch. it. 1900, 30. (1) 593.)

nitrohydroxylamine, $\text{K}_2\text{N}_2\text{O}_3$.
salt. More hygroscopic. (Angeli, Gazz. ch. it. 1897, 27. (2) 357.)
 H_2O . (Angeli, Gazz. ch. it. 1900, 30. (1) 593.)

hydroxylamine, $\text{Ag}_2\text{N}_2\text{O}_3$.
Angeli, C. C. 1901, I. 1192.)

hydroxylamine, $\text{Na}_2\text{N}_2\text{O}_3$.
sol. in H_2O . Pptd. by alcohol. solution is readily decomp. by boiling. (Angeli, Gazz. ch. it. 1896, 26. (2) 17.)

nitrohydroxylamine, $\text{SrN}_2\text{O}_3 + \text{H}_2\text{O}$.
Gazz. ch. it. 1900, 30. (1) 593.)
 H_2O . (Angelico and Fanara, Gazz. ch. it. 1901, 31. (2) 15.)

acid, $\text{I}_2\text{O}_4(\text{NO})_2$.
osiodic acid.

ous acid.

potassium nitronitrite, $\text{K}_2\text{Pt}(\text{NO}_2)_4$.
decomp. by heat. (Miolati, C. C. 1896, II. 1192.)

inous acid.

onitrous acid.

isic acid, $\text{H}_2\text{FeC}_6\text{N}_6\text{O} + \text{H}_2\text{O} = \text{Fe}(\text{CN})_6\text{NO} + \text{H}_2\text{O}$.

cent. Easily sol. in H_2O , alcohol, (Playfair, A. 74. 317.)

ides.

ali and alkali-earth nitroprussides
 H_2O , and the solutions are not
cohol. The others are mostly insol.

a nitroprusside,
 $\text{Fe}(\text{CN})_6(\text{NO})$.

cent. Very sol. in H_2O ; not pptd.
by alcohol. (Playfair.)

Barium nitroprusside, $\text{BaFe}(\text{CN})_6\text{NO} + 4\text{H}_2\text{O}$.

Very sol. in H_2O .
 $+6\text{H}_2\text{O}$.

Cadmium nitroprusside, $\text{CdFe}(\text{CN})_6\text{NO}$.

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. Insol. in
dil. or conc. $\text{HNO}_3 + \text{Aq}$ even when boiling.
Not attacked by NH_4OH or $\text{KOH} + \text{Aq}$.
(Norton, Am. Ch. J. 10. 222.)

Calcium nitroprusside, $\text{CaFe}(\text{CN})_6\text{NO} + 4\text{H}_2\text{O}$.

Very sol. in H_2O . (Playfair.)

Cobalt nitroprusside, $\text{CoFe}(\text{CN})_6\text{NO}$.

Ppt. (Norton, Am. Ch. J. 10. 222.)
 $+4\text{H}_2\text{O}$.

Copper nitroprusside, $\text{CuFe}(\text{CN})_6\text{NO} + 2\text{H}_2\text{O}$.

Insol. in H_2O or alcohol.

Ferrous nitroprusside, $\text{FeFe}(\text{CN})_6\text{NO} + x\text{H}_2\text{O}(?)$.

Insol. in H_2O .

Mercurous nitroprusside, $\text{Hg}_2\text{Fe}(\text{CN})_6\text{NO}$.

Insol. in H_2O . Unstable. (Norton, Am. Ch. J. 10. 222.)

Nickel nitroprusside, $\text{NiFe}(\text{CN})_6\text{NO}$.

As the Co salt. (Norton.)

Potassium nitroprusside, $\text{K}_2\text{Fe}(\text{CN})_6\text{NO} + 2\text{H}_2\text{O}$.

Sl. deliquescent. Sol. in 1 pt. H_2O at 16° .
 $\text{K}_2\text{Fe}(\text{CN})_6\text{NO}$, 2KOH . Very sol. in H_2O .

Silver nitroprusside, $\text{Ag}_2\text{Fe}(\text{CN})_6\text{NO}$.

Insol. in H_2O , alcohol, or $\text{HNO}_3 + \text{Aq}$. Sol.
in $\text{NH}_4\text{OH} + \text{Aq}$.

Sodium nitroprusside, $\text{Na}_2\text{Fe}(\text{CN})_6\text{NO} + 2\text{H}_2\text{O}$.

Sol. in $2\frac{1}{2}$ pts. H_2O at 16° , and in less hot
 H_2O .

Zinc nitroprusside, $\text{ZnFe}(\text{CN})_6\text{NO}$.

Very sl. sol. in cold, more in hot H_2O .

Nitrosulphonic acid.

Cupric nitrosulphonate, $\text{NO} \begin{smallmatrix} \text{O} \\ \diagup \text{SO}_2 \diagdown \end{smallmatrix} \text{Cu}$.

Decomp. by H_2O . (Raschig, B. 1907, 40. 4583.)

Nitrosobromoruthenic acid.

Silver nitrosobromoruthenate ammonia,
 $\text{Ag}_2\text{Ru}(\text{NO})\text{Br}_2, \text{NH}_3$.

Decomp. by H_2O . Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Very sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Brizard, Bull. Soc. 1895, (3) 13. 1093.)

Nitrosobromosmic acid.

Potassium nitrosobromosmate, $K_2Os(NO)Br_2$.
Stable in aqueous solution. (Wintrebert, A. ch. 1903, (7) 28. 132.)

Nitrosochloroplatinic acid.

Potassium nitrosochloroplatinate,
 $K_2PtCl_4(NO)$.
Sol. in H_2O . (Vèzes, C. R. 110. 757.)

Nitrosochlororuthenic acid.

Ammonium nitrosochlororuthenate,
 $(NH_4)_2Ru(NO)Cl_4$.
Sol. in H_2O . (Joly, C. R. 107. 991.)
5 pts. are sol. in 100 pts. H_2O at 25° .
22 " " " " " " " " 60° .
(Howe, J. Am. Chem. Soc. 1894, 16. 390.)

Cæsium nitrosochlororuthenate,
 $Cs_2Ru(NO)Cl_4$.
0.20 pt. is sol. in 100 pts. H_2O at 25° .
0.56 " " " " " " " " 100° .
(Howe.)
+ $2H_2O$. Very sol. in H_2O . 105.8 pts. are
sol. in 100 pts. H_2O . (Howe.)

Potassium nitrosochlororuthenate,
 $K_2Ru(NO)Cl_4$.
Sol. in H_2O . (Joly.)
12 pts. are sol. in 100 pts. H_2O at 25° .
80 " " " " " " " " 60° .
(Howe.)

Rubidium nitrosochlororuthenate,
 $Rb_2Ru(NO)Cl_4$.
Sol. in boiling H_2O without decomp.
0.57 pt. is sol. in 100 pts. H_2O at 25° .
2.13 " " " " " " " " 60° .
(Howe.)

Silver nitrosochlororuthenate ammonia,
 $Ag_2Ru(NO)Cl_4, NH_3$.
Decomp. by H_2O . Sl. sol. in $NH_4OH + Aq$.
Very sol. in $Na_2S_2O_3 + Aq$. (Brizard, Bull.
Soc. 1895, (3) 13. 1092.)

Nitrosoiodic acid, $I_2O_4(NO)_2$ (?)
Decomp. with H_2O , alcohol, ether, or acetic
ether. Slowly sol. in H_2SO_4 . (Kämmerer, J.
pr. 83. 65.)

Nitrososulphonic acid.

Potassium dinitrososulphonate, $N_2O.O.K.$
 SO_3K .
Sol. in H_2O . Very unstable. (Hantzsch,
B. 1894, 27. 3268.)

Potassium nitrosodisulphonate, $ON(SO_3K)_2$.
Sol. in H_2O . Very explosive. (Hantzsch,
B. 1895, 28. 996 and 2744.)

Potassium nitrosotrisulphonate,
+ H_2O .

Sol. in H_2O . (Hantzsch, B. 1894

Sodium nitrosotrisulphonate, NO
Decomp. by H_2O . (Traube, E
2521.)

Nitrososulphuric acid,
 $H_2N_2SO_4 = H_2SO_4(NO)_2$.
Not known in free state.

Ammonium dinitrososulphate,
 $(NH_4)_2(NO)_2SO_4$.
Sol. in H_2O . Insol. in hot alc
louze, A. 15. 240.)

Barium —, $Ba(NO)_2SO_4$.
Sol. in H_2O . (Divers and Haga,
47. 364.)

Barium potassium —, $BaK_2(SN$
Sol. in much H_2O to form a c
but the solution gradually depos
(Hantzsch, B. 1894, 27. 3271.)

Cupric nitrosodisulphate, $Cu(NO)$
(Sabatier, Bull. Soc. 1897, (3) 1'

Lead dinitrososulphate.
Insol. in H_2O . (Divers and H
Soc. 47. 364.)

Potassium —, $K_2(NO)_2SO_4$.
Decomp. by H_2O at ordinary te
in alcohol. (Pelouze, A. ch. 60. 1
Sol. in about 8 pts. H_2O at 14.5°
in presence of KOH . (Divers
Chem. Soc. 1895, 67. 455.)

Sodium —, $Na_2(NO)_2SO_4$.
More sol. than K salt. (Pelous
Sl. sol. in H_2O ; very unstable m
decomp. by H_2O . (Divers, C. N
266.)

Nitrososulphurous acid.

Ruthenium sodium nitrososulphite
 $O[Ru(SO_3)_2(NO)Na_2] + 2H_2O$
Sl. sol. in cold H_2O . (Miolati
it. 1900, 30. 511.)

Nitrosulphide of iron.

See Ferrotetranitrososulphonic a

Binitrosulphide of iron.

Roussin's comp. is ammonium
nitrososulphonate, which see.

ionic acid, $\text{HNSO}_3 = \frac{\text{HO}}{\text{NO}_2} \text{SO}_3$.

amber crystals.) Rapidly sol. in lecomp. When brought into large H_2O , no gas is evolved. (Freymy, 1.)

SO_3 without decomp. Sol. in cold of sp. gr. 1.7–1.55 (Weber, J. 1.)

$\text{H}_2\text{SO}_4 + \text{Aq}$ of 1.6 sp. gr. (Dana.) difficultly sol. in dil. than conc. q. (Müller.)

nitrosulphonate, $\text{KOSO}_2\text{NO}_2(?)$.

by H_2O . (Schultz-Sellack, B. 4.)

ionic anhydride (?), $\text{N}_2\text{O}_3, 2\text{SO}_3, (\text{NO}_2)_2$.

sol. in H_2O with decomp. Abundant in cold H_2SO_4 . (Rose, Pogg. 47.)

cold, slowly sol. in warm H_2SO_4 . (Re, A. ch. 73. 362.)

ionic chloride, $\text{NO}_2\text{SCl} = \text{O}_2\text{Cl} (?)$.

by H_2O . Sol. in fuming H_2SO_4 decomp. Decomp. by conc. H_2SO_4 . (Pogg. 123. 333.)

sulphuric acid.

nitrososulphuric acid.

bromide, NOBr .

with cold H_2O . (Landolt, A. 116.)

tribromide, NOBr_3 .

p. by H_2O or cold alcohol. e with ether. (Landolt, A. 116. 177.) e of NOBr and Br_2 . (Fröhlich, A. 1.)

platinic bromide, $2\text{NOBr}, \text{PtBr}_4$.

ascent. Decomp. by H_2O . (Toppe, 1868. 274.)

chloride, NOCl .

p. by H_2O . Absorbed by fuming without decomp.

boron chloride, $\text{NOCl}, \text{BCl}_3$.

boron nitrosyl chloride.

platinic chloride, $2\text{NOCl}, \text{PtCl}_4$.

deliquescent, and sol. in H_2O with evolution of NO . (Rogers and Boye, Phil. 7. 397.)

Nitrosyl thallium chloride, $2\text{NOCl}, \text{TlCl}, \text{TlCl}_3$.

Very deliquescent, and sol. in H_2O with decomp. (Sudborough, Chem. Soc. 59. 657.)

Nitrosyl stannic chloride, $2\text{NOCl}, \text{SnCl}_4$.

Decomp. by H_2O , chloroform, or benzene, not by carbon disulphide. (Jørgensen.)

Nitrosyl titanium chloride, $2\text{NOCl}, \text{TiCl}_4$.

Decomp. by H_2O . (Weber, Pogg. 118. 476.)

Nitrosyl zinc chloride, $\text{NOCl}, \text{ZnCl}_2$.

Very deliquescent, and sol. in H_2O with evolution of NO . (Sudborough, Chem. Soc. 59. 656.)

Nitrosyl chloride sulphur trioxide, NOCl, SO_3 .

Decomp. by H_2O . Sol. in conc. H_2SO_4 with evolution of HCl . (Weber, Pogg. 123. 233.)

Nitrosyl fluoride, NOF .

Sol. in H_2O . Solution decomp. on standing with formation of NO and HNO_3 . (Ruff and Stäuber, Z. anorg. 1905, 47. 190.)

Nitrosyl sulphate, acid, $\text{H}(\text{NO})\text{SO}_4$.

See Nitrosulphonic acid.

Nitrosyl sulphate, anhydro, $(\text{NO})_2\text{S}_2\text{O}_7$.

See Nitrosulphonic anhydride.

Nitrosyl selenic acid, $\text{SeO}_3(\text{ONO})_2$.

Decomp. by H_2O . (Lenher and Mathews, J. Am. Chem. Soc. 1906, 28. 516.)

Nitrosyl sulphuric acid, $\text{H}(\text{NO})\text{SO}_4$.

See Nitrosulphonic acid.

Nitrous acid, HNO_2 .

Known only in aqueous solution.

See Nitrogen trioxide.

Nitrites.

Normal nitrites, except AgNO_2 , are sol. in H_2O and alcohol; but, as a rule, they are less sol. than the corresponding nitrates.

Ammonium nitrite, NH_4NO_2 .

Very deliquescent, and sol. in H_2O .

H_2O solution decomp. at 50° . (Berzelius.) Very dil. solution can be evaporated on water bath without decomp. (Bohlig, A. 125. 25.) Solution containing $\frac{1}{100,000}$ pt. NH_4NO_2 can be evaporated to $\frac{1}{2}$ its vol. without decomp. Solution containing $\frac{1}{500}$ pt. gives a distillate containing 8.6% of NH_4NO_2 , while residue contains 82% of original quantity, 9.4% being lost. (Schöyen.)

Very deliquescent, sol. in H_2O ; slowly but easily sol. in alcohol; insol. in ether. (Sørensen, Z. anorg. 1894, 7. 38.)

Ammonium barium cupric nitrite,
 $(\text{NH}_4)_2\text{BaCu}(\text{NO}_2)_6$.
 Ppt.; decomp. readily. (Przibylla, Z. anorg. 1897, 15. 424.)

Ammonium bismuth silver nitrite,
 $(\text{NH}_4)_2\text{BiAg}(\text{NO}_2)_6$.
 Moderately sol. in H_2O . Rapidly hydrolysed by H_2O . (Ball and Abram, Chem. Soc. 1913, 103. 2120.)

Ammonium bismuth sodium nitrite,
 $2\text{NH}_4\text{NO}_2, \text{Bi}(\text{NO}_3)_3, \text{NaNO}_2$.
 Easily decomp. (Ball, Chem. Soc. 1905, 87. 761.)

Ammonium cadmium nitrite ammonia, basic,
 $2\text{NH}_4\text{NO}_2, \text{Cd}(\text{NO}_2)_2, \text{Cd}(\text{OH})_2, 2\text{NH}_3$.
 Decomp. by H_2O . (Morin, C. R. 100. 1497.)

Ammonium calcium cupric nitrite,
 $(\text{NH}_4)_2\text{CaCu}(\text{NO}_2)_6$.
 Ppt.; decomp. easily. Sol. in H_2O . Sl. sol. in alcohol. (Przibylla, Z. anorg. 1897, 15. 423.)

Ammonium cobaltic nitrite, $3(\text{NH}_4)_2\text{O}, \text{Co}_2\text{O}_3, 6\text{N}_2\text{O}_5 + 1\frac{1}{2}\text{H}_2\text{O}$.
 Sl. sol. in H_2O , decomp. in aq. solution on heating. (Rosenheim, Z. anorg. 1898, 17. 45.)

+ $3\text{H}_2\text{O}$.
 Somewhat sol. in cold H_2O ; decomp. by boiling. Decomp. by conc. H_2SO_4 , not by acetic or dil. mineral acids. (Erdmann, J. pr. 97. 405.)

Ammonium cupric lead nitrite,
 $\text{CuPb}(\text{NH}_4)_2(\text{NO}_2)_6$.
 Stable at ordinary temp.; sol. in HNO_3 with decomp.
 2.575 pts. are sol. in 100 pts. H_2O at 20° , or 2.51% salt in sat. solution at 20° . (Przibylla, Z. anorg. 1897, 15. 420.)

Ammonium cupric strontium nitrite,
 $(\text{NH}_4)_2\text{CuSr}(\text{NO}_2)_6$.
 Ppt.; sol. in H_2O with decomp. (Przibylla, l. c.)

Ammonium iridium nitrite.
 See Iridonitrite, ammonium.

Ammonium lead nickel nitrite,
 $(\text{NH}_4)_2\text{PbNi}(\text{NO}_2)_6(?)$.
 Ppt. (Przibylla, Z. anorg. 1897, 15. 433.)

Ammonium osmium nitrite.
 See Osminitrite, ammonium.

Ammonium osmyl oxynitrite.
 See Osmyloxynitrite, ammonium.

Ammonium platinum nitrite.
 See Platonitrite, ammonium.

Ammonium rhodium nitrite.
 See Rhodonitrite, ammonium.

Ammonium ruthenium hydrogen nitrite,
 $\text{Ru}_2\text{H}_2(\text{NO}_2)_4, 3\text{NH}_4\text{NO}_2 + 3\text{H}_2\text{O}$.
 See Rutheninitrite, ammonium hydrogen

Barium nitrite, $\text{Ba}(\text{NO}_2)_2 + \text{H}_2\text{O}$.
 Permanent. Very sol. in H_2O .

Solubility in H_2O at t° .

t°	G. in 100 cc. $\text{Ba}(\text{NO}_2)_2$	Sp. gr.
0	58	1.40
20	63	1.45
25	71	1.50
30	82	1.53
35	97	1.61

(Vogel, Z. anorg. 1903, 35. 399.)

100 pts. H_2O dissolve at:			
0°	10°	20°	30°
63.5	69.5	79.5	93 pts. $\text{Ba}(\text{NO}_2)_2 + \text{H}$
40°	50°	60°	70°
113	136	170	202 pts. $\text{Ba}(\text{NO}_2)_2 + \text{H}$
80°	90°	100°	110°
254	331	461	765 pts. $\text{Ba}(\text{NO}_2)_2 + \text{H}$

The sat. solution at 17° contains 4 $\text{Ba}(\text{NO}_2)_2$, and has sp. gr. $17^\circ/0^\circ = 1.41$ (Oswald, A. ch. 1914, (9) 1. 62.)

100 g. H_2O at 13.5° dissolve 64 g. $\text{Ba}(\text{NO}_2)_2$ + 10.2 g. AgNO_3 with excess of AgNO_3 , 75.6 g. $\text{Ba}(\text{NO}_2)_2$ + 9.5 g. AgNO_3 with excess of AgNO_3 . (Oswald.)

Sol. in 64 pts. 94% alcohol; nearly insol. in absolute alcohol. (Lang, Pogg. 112. 3)

Solubility in alcohol + Aq at t° .

t°	Solvent	100 ccm. of the solution contain $\text{Ba}(\text{NO}_2)_2 + \text{H}_2\text{O}$
19.5	10% alcohol	49.30
21.0	20% "	29.30
20.5	30% "	18.41
20.5	40% "	13.33
20.5	50% "	9.11
20.0	60% "	4.84
19.0	70% "	2.66
19.5	80% "	0.98
20.0	90% "	0.00
20.0	absolute alcohol	0.00

(Vogel, Z. anorg. 1903, 35. 390.)

Insol. in acetone. (Naumann, B. 1904, 4329.)

Insol. in ethyl acetate. (Naumann, 1910, 43. 314.)

caesium nitrite, $\text{CsBa}_2(\text{NO}_2)_6$.

H_2O . (Jamieson, Am. Ch. J. 1907, 38. 616.)
 $(\text{NO}_2)_4 + \text{H}_2\text{O}$. Very sol. in H_2O .
 n, Am. Ch. J. 1907, 38. 616.)

caesium silver nitrite, $\text{Cs}_2\text{AgBa}(\text{NO}_2)_6$, H_2O .

np. by cold H_2O . (Jamieson, Am. Ch. 38. 616.)

cobaltic nitrite, 2BaO , Co_2O_3 , $4\text{N}_2\text{O}_5$, $10\text{H}_2\text{O}$.

1 moderately warm H_2O without de-
 out not recryst. therefrom. (Rosen-
 anorg. 1898, 17. 51-54.)

, Co_2O_3 , $6\text{N}_2\text{O}_5 + \text{H}_2\text{O}$. Ppt.; very un-
 Nearly insol. in H_2O . (Rosenheim,
 1898, 17. 47.)

cobaltous potassium nitrite, $\text{Ba}(\text{NO}_2)_2$, $\text{NO}_2)_2$, 2KNO_3 .

np. by H_2O . (Erdmann, J. pr. 97.

cupric nitrite, $\text{Ba}[\text{Cu}(\text{OH})(\text{NO}_2)_2]_2$.

Insol. in H_2O . Decomp. by H_2O .
 1 alcohol, but slowly decomp. by it.
 acker, Z. anorg. 1913, 82. 208.)

cupric potassium nitrite,
 $3\text{uK}_2(\text{NO}_2)_6$.

3 when dry, easily decomp. when
 ol. in H_2O with decomp.

pts. are sol. in 100 pts. H_2O at 20° , or
 salt is contained in sat. solution at
 Przibylla, Z. anorg. 1897, 15. 424.)

cupric thallium nitrite,
 $3\text{uTl}_2(\text{NO}_2)_6$.

l. in H_2O . (Przibylla, Z. anorg. 1898,
)

iridium nitrite.

idonitrite, barium.

mercuric nitrite, $2\text{Ba}(\text{NO}_2)_2$,
 $\text{Hg}(\text{NO}_2)_2 + 5\text{H}_2\text{O}$.

sol. in H_2O and easily decomp. (Ray,
 Soc. 1910, 97. 327.)

nickel nitrite, $2\text{Ba}(\text{NO}_2)_2$, $\text{Ni}(\text{NO}_2)_2$.

what more easily sol. in H_2O than
 potassium nitrite. (Lang.)

nickel potassium nitrite, $\text{Ba}(\text{NO}_2)_2$,
 $\text{NO}_2)_2$, 2KNO_3 .

l. in cold, easily in hot H_2O without
 it decomp. (Lang.)

nickel thallium nitrite,
 $\text{BaTl}_2(\text{NO}_2)_6$.

(Przibylla, Z. anorg. 1898, 18. 462.)

Barium osmium nitrite.

See Osminitrite, barium.

Barium osmyl oxynitrite.

See Osmyloxynitrite, barium.

Barium potassium nitrite, $\text{Ba}(\text{NO}_2)_2$, 2KNO_3 ,
 $+ \text{H}_2\text{O}$.

Easily sol. in H_2O ; insol. in alcohol. (Lang,
 Pogg, 118. 293.)

Barium rhodium nitrite, $3\text{Ba}(\text{NO}_2)_2$,
 $\text{Rh}_2(\text{NO}_2)_6$.

See Rhodonitrite, barium.

Barium silver nitrite, $\text{Ba}(\text{NO}_2)_2$, $2\text{AgNO}_3 +$
 H_2O .

Resembles the potassium salt. (Fischer.)
 Less stable than the Na salt. (Oswald.)

Bismuth nitrite, basic, $(\text{BiO})\text{NO}_2 + \frac{1}{2}\text{H}_2\text{O}$.

Sol. in HCl . (Vanino, J. pr. 1906, (2) 74.
 150.)

Bismuth caesium silver nitrite, $\text{Cs}_2\text{BiAg}(\text{NO}_2)_6$.

Very sl. sol. in H_2O . Slowly decomp. by
 H_2O . (Ball and Abram, Chem. Soc. 1913,
 103. 2122.)

Bismuth potassium nitrite, $\text{Bi}(\text{NO}_2)_3$, 3KNO_3 ,
 $+ \text{H}_2\text{O}$.

Decomp. by H_2O . (Ball, Chem. Soc. 1905,
 87. 762.)

Bismuth potassium silver nitrite,
 $\text{K}_2\text{BiAg}(\text{NO}_2)_6$.

Less sol. in H_2O than NH_4 salt. (Ball and
 Abram, Chem. Soc. 1913, 103. 2121.)

Bismuth rubidium silver nitrite,
 $\text{Rb}_2\text{BiAg}(\text{NO}_2)_6$.

Sl. sol. in H_2O with slow hydrolysis. (Ball
 and Abram.)

Bismuth silver thalious nitrite, $\text{BiAgTl}_2(\text{NO}_2)_6$.

Insol. in H_2O , but decomp. thereby. (Ball
 and Abram.)

Cadmium nitrite, basic, 2CdO , N_2O_5 .

Insol. in H_2O . (Hampe, A. 125. 335.)

Cadmium nitrite, $\text{Cd}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . (Lang, J. B.
 1862. 99.)

Cadmium potassium nitrite, $\text{Cd}(\text{NO}_2)_2$, KNO_3 .

Easily sol. in H_2O . Very difficultly sol. in
 absolute alcohol, and only sl. sol. in 90%
 alcohol. (Hampe, A. 125. 334.)

$\text{Cd}(\text{NO}_2)_2$, 2KNO_3 . Easily sol. in H_2O .
 Insol. in alcohol. (Lang, J. B. 1862. 99.)

$\text{Cd}(\text{NO}_2)_2$, 4KNO_3 . More sol. in H_2O than
 the above salt. (Lang.)

Cæsium nitrite, CsNO_2 .

Very hygroscopic. Very sol. in H_2O . (Ball, Chem. Soc. 1913, 103. 2130.)

Cæsium calcium nitrite, $\text{Cs}_2\text{Ca}(\text{NO}_2)_4 + \text{H}_2\text{O}$.

Ppt. (Jamieson, Am. Ch. J. 1907, 38. 617.)

Cæsium cobaltic nitrite, $\text{Cs}_3\text{Co}(\text{NO}_2)_6 + \text{H}_2\text{O}$.

Sol. in 20,100 pts. H_2O at 17° . (Rosenblatt, B. 19. 2531.)

Cæsium lead nitrite, $\text{CsPb}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

Sol. in cold H_2O without decomp. When solution is heated, some basic lead salt separates. (Jamieson, Am. Ch. J. 1907, 38. 618.)

Cæsium lead silver nitrite, $\text{Cs}_2\text{AgPb}(\text{NO}_2)_6 + 2\text{H}_2\text{O}$.

Ppt. (Jamieson.)

Cæsium silver nitrite, $\text{CsAg}(\text{NO}_2)_2$.

Decomp. by H_2O . (Jamieson.)

Cæsium silver strontium nitrite,

Partially decomp. by H_2O . (Jamieson.)

Cæsium strontium nitrite, $\text{CsSr}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

Ppt. Sol. in H_2O . (Jamieson.)

Calcium nitrite, $\text{Ca}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

Very deliquescent. Insol. in dil. alcohol. (Fischer, Pogg. 74. 115.)

100 ccm. of the sat. solution contain 111.6 g. $\text{Ca}(\text{NO}_2)_2 + \text{H}_2\text{O}$ at 20.5° . (Vogel, Z. anorg. 1903, 35. 395.)

Solubility in H_2O at t° .

t°	% $\text{Ca}(\text{NO}_2)_2$	Solid phase
0	38.3	$\text{Ca}(\text{NO}_2)_2, 4\text{H}_2\text{O}$
18.5	43	"
42	51.8	"
44	53.5	" + $\text{Ca}(\text{NO}_2)_2, \text{H}_2\text{O}$
54	55.2	$\text{Ca}(\text{NO}_2)_2, \text{H}_2\text{O}$
64	58.4	"
70	60.3	"
73	61.5	"
91	71.2	"

(Oswald, A. ch. 1914, (9) 1. 32.)

Sat. solution of $\text{Ca}(\text{NO}_2)_2 + \text{AgNO}_2$ contains 92.4 g. $\text{Ca}(\text{NO}_2)_2$ and 11.2 g. AgNO_2 per 100 g. H_2O at 14° . (Oswald.)

Solubility in alcohol.

100 ccm. of sat. solution in 90% alcohol contain 39.0 g. $\text{Ca}(\text{NO}_2)_2 + \text{H}_2\text{O}$ at 20° .

100 ccm. of sat. solution in absolute alcohol contain 1.1 g. $\text{Ca}(\text{NO}_2)_2 + \text{H}_2\text{O}$ at 20° . (Vogel.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

+ $4\text{H}_2\text{O}$. The sat. solution at 16° contains 42.3% $\text{Ca}(\text{NO}_2)_2$ and has sp. gr. at 16° 1.4205. (Oswald, A. ch. 1914, (9) 1. 66.)

Calcium cobaltous potassium nitrite,

Decomp. by H_2O . (Erdmann.)

Calcium cupric potassium nitrite,

Ppt., insol. in alcohol; sol. in H_2O with decomp.

14.97 pts. are sol. in 100 pts. H_2O at 20° , or 13.02 per cent of salt is contained in sat. solution. (Przibylla, Z. anorg. 1897, 15. 432.)

Calcium mercuric nitrite, $\text{Ca}(\text{NO}_2)_2, \text{Hg}(\text{NO}_2)_2 + 5\text{H}_2\text{O}$.

Very sol. in H_2O . (Ray, Chem. Soc. 1914, 97. 327.)

Calcium nickel potassium nitrite, $\text{Ca}(\text{NO}_2)_2, \text{Ni}(\text{NO}_2)_2, 2\text{KNO}_2$.

Very sl. sol. in cold, easily in hot H_2O . Insol. in alcohol. Sl. sol. in dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Erdmann.)

Calcium osmium nitrite.

See Osminitrite, calcium.

Calcium potassium nitrite, $\text{CaK}(\text{NO}_2)_3 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Topsoë, W. A. B. 72, 2. 112) Deliquescent. (Lang.)

Cobaltous nitrite.

Known only in solution.

Cobaltic lead nitrite, $3\text{PbO}, \text{Co}_2\text{O}_3, 6\text{N}_2\text{O}_5 + 12\text{H}_2\text{O}$.

Insol. in H_2O . (Rosenheim, Z. anorg. 1896, 17. 48.)

Cobaltic lead potassium nitrite, $3\text{K}_2\text{O}, 3\text{PbO}, 2\text{Co}_2\text{O}_3, 10\text{N}_2\text{O}_5 + 4\text{H}_2\text{O}$.

Sol. by boiling in much H_2O . Sol. in hot acids with evolution of N_2O_5 . (Stromeyer, A. 96. 228.)

Cobaltous potassium nitrite, $2\text{Co}(\text{NO}_2)_2, 2\text{KNO}_2 + \text{H}_2\text{O}$.

Ppt. (Sadtler.)

$\text{Co}(\text{NO}_2)_2, 2\text{KNO}_2 + \text{H}_2\text{O}$. Ppt. (Sadtler.) $3\text{Co}(\text{NO}_2)_2, 6\text{KNO}_2 + \text{H}_2\text{O}$. Insol. in cold, sol. in hot H_2O . Sl. sol. in $\text{KC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Erdmann, J. pr. 97. 397.)

Insol. in ethyl acetate. (Naumann, B. 1904, 37. 3602.)

Cobaltic potassium nitrite (cobalt yellow), $\text{Co}_2(\text{NO}_2)_6, 6\text{KNO}_2 + 3\text{H}_2\text{O}$.

Very sl. sol. in cold H_2O . Insol. in alcohol and ether. Sol. in traces in CS_2 . (St. Evre,

552.) Insol. in boiling conc. K_2SO_4 , IO_3 , or $KC_2H_3O_2 + Aq$. 1120 pts. H_2O at 17° . (Rosenblatt, 19. 2535.)

sp. when heated in aq. solution. *Ann.*, Z. anorg. 1898, 17. 42.)

sol. in NH_4Cl or $NaCl + Aq$ than in Tromeyer.)

comp. by $KOH + Aq$, except when ac.; easily decomp. by $NaOH$ or $+ Aq$.

sl. sol. in $KC_2H_3O_2 + Aq$, or KNO_3 (Fresenius.) Sol. in $HCl + Aq$.

$HC_2H_3O_2$, or $H_2C_2O_4 + Aq$. (Stro-

quantity of $HC_2H_3O_2 + Aq$ does not (Fresenius.)

potassium silver nitrite, $Ag_2(NO_2)_6$, and $K_2CoAg(NO_2)_6$.

sl. sol. in H_2O . Less sol. than Na Burgess and Karum, J. Am. Chem. Soc., 34. 653.)

potassium strontium nitrite, $NO_2)_2$, $2KNO_2$, $Sr(NO_2)_2$.

p. by H_2O . (Erdmann, J. pr. 97.

rubidium nitrite, $Rb_3Co(NO_2)_6 + 19,800$ pts. H_2O . (Rosenblatt, B.)

silver nitrite, $CoAg_3(NO_2)_6$.

sol. in H_2O . (Cunningham and Chem. Soc. 1909, 95. 1568.)

Co_2O_3 , $3N_2O_5 + 3H_2O$. Sl. sol. in comp. by boiling H_2O . (Rosenheim, 1898, 17. 56.)

silver hydroxynitrite, $Ag_3(OH)_3(NO_2)_6$.

in H_2O . (Suzuki, Chem. Soc. 1910,

silver nitrite ammonia, Co_2O_3 , Ag_2O , $)_3$, $4NH_3$.

balt ammonium comps.

odium nitrite, $2Na_2O$, Co_2O_3 , $4N_2O_5$.

H_2O and alcohol. (Rosenheim, Z. 98, 17. 50.)

Ppt. (Sadtlér, Sill. Am. J. (2) 49.

Co_2O_3 , $6N_2O_5 + xH_2O$. Sol. in H_2O ; on heating; insol. in alcohol. (Rosenheim, Z. anorg. 1898, 17. 43.)

strontium nitrite, $2SrO$, Co_2O_3 , $4N_2O_5$, H_2O .

Rosenheim, Z. anorg. 1898, 17. 54.)

Cobaltic thallium nitrite, $Co_2(NO_2)_6$, $6TlNO_2$.

Sol. in 23,810 pts. H_2O at 17° . (Rosenblatt, B. 19. 2531.)

Cobaltic zinc nitrite, $2ZnO$, Co_2O_3 , $3N_2O_5 + 11H_2O$.

Sol. in dil. acetic acid. (Rosenheim, Z. anorg. 1898, 17. 56.)

Cobalt nitrite nitrate, $2CoO$, Co_2O_3 , $3N_2O_5$, $Co(NO_2)_2 + 14H_2O$.

Ppt. (Rosenheim, Z. anorg. 1898, 17. 58.)

Cupric nitrite, basic, $2CuO$, N_2O_5 .

(Hampe, A. 125. 345.)

$Cu(NO_2)_2$, $3Cu(OH)_2$. Very sl. sol. in H_2O or alcohol. Easily sol. in dil. acids or ammonia. (van der Meulen, B. 12. 758.)

Cupric nitrite.

Known only in solution.

Cupric lead potassium nitrite, $CuPbK_2(NO_2)_6$.

(van Lessen, R. t. c. 10. 13.)

3.056 pts. are sol. in 100 pts. H_2O at 20° , or 2.51% salt is contained in sat. solution at 20° p. 429. (Przibylla, Z. anorg. 1897, 15. 429.)

Cupric potassium strontium nitrite, $CuSrK_2(NO_2)_6$.

Sol. in H_2O with decomp. 10.82 pts. are sol. in 100 pts. H_2O at 20° , or 9.77 per cent salt is contained in sat. solution at 20° . (Przibylla, Z. anorg. 1897, 15. 425.)

Cupric rubidium nitrite, $Rb_3Cu(NO_2)_6$.

Easily sol. in H_2O . Sol. in alcohol. (Kurténacker, Z. anorg. 1913, 82. 206.)

Cupric nitrite ammonia, $Cu(NO_2)_2$, $2NH_3 + 2H_2O$.

Sol. in little H_2O with absorption of much heat. Decomp. by much H_2O . (Peligot, C. R. 53. 209.)

$3CuO$, N_2O_5 , $2NH_3 + H_2O$. As above. (Peligot.)

Iridium hydrogen nitrite, $Ir_2H_4(NO_2)_{12}$.

See Iridonitrous acid.

Iridium nitrite with MNO_3 .

See Iridonitrite, M.

Iron (ferrous) lead potassium nitrite, $FePbK_2(NO_2)_6$.

Ppt.; insol. in cold H_2O ; stable at ordinary temp. (Przibylla, Z. anorg. 1897, 15. 439.)

Iron (ferrous) lead thallos nitrite, $FePbTl_2(NO_2)_6$.

Ppt. (Przibylla, Z. anorg. 1898, 18. 483.)

Lead nitrite, basic, 4PbO , $\text{N}_2\text{O}_3 + \text{H}_2\text{O} = \text{Pb}(\text{OH})\text{NO}_2$, PbO .

Sol. in 143 pts. H_2O at 23° , and 33 pts. at 100° . (Chevreul.)

Sol. in 1250 pts. cold H_2O , and 34.5 pts. at 100° . (Peligot.)

Sol. in cold HNO_3 or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

Composition is 3PbO , $\text{N}_2\text{O}_3 + \text{H}_2\text{O}$. (Meissner, J. B. 1876. 194.)

Composition is as above. (v. Lorenz, W. A. B. 84, 2. 1133.)

3PbO , $\text{N}_2\text{O}_3 = \text{Pb}(\text{NO}_2)_2$, 2PbO . Sol. in H_2O . (Bromeis, A. 72. 38; v. Lorenz.)

2PbO , $\text{N}_2\text{O}_3 + \text{H}_2\text{O}$. Sl. sol. in H_2O . (Bromeis.)

+ $3\text{H}_2\text{O}$. (Meissner.)

4PbO , $3\text{N}_2\text{O}_3 + 2\text{H}_2\text{O}$. Sol. in H_2O . (Meissner, J. B. 1876. 195.)

Lead nitrite, $\text{Pb}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

Easily sol. in H_2O . (Peligot, A. ch. 77. 87.)

Lead nickel potassium nitrite, $\text{Pb}(\text{NO}_2)_2$, KNO_2 , $\text{Ni}(\text{NO}_2)_2$.

Insol. in H_2O . (Baubigny, A. ch. (6) 17. 111.)

Ppt. (Przibylla, Z. anorg. 1897, 15. 432.)

Lead nickel thalious nitrite, $\text{NiPbTl}_2(\text{NO}_2)_6$.

Ppt. (Przibylla, Z. anorg. 1898, 18. 462.)

Lead potassium nitrite, $4\text{Pb}(\text{NO}_2)_2$, $6\text{KNO}_2 + 3\text{H}_2\text{O}$.

Easily sol. in H_2O and in absolute alcohol. (Hampe, A. 125. 334.)

$\text{Pb}(\text{NO}_2)_2$, $2\text{KNO}_2 + \text{H}_2\text{O}$. Easily sol. in H_2O . Insol. in alcohol. (Lang, J. B. 1862. 102.)

Lead potassium silver nitrite, $\text{K}_2\text{AgPb}(\text{NO}_2)_6 + 2\text{H}_2\text{O}$.

Ppt. (Jamieson, Am. Ch. J. 1907, 38. 619.)

Lead nitrite nitrate.

See Nitrate nitrite, lead.

Lithium nitrite, $\text{LiNO}_2 + \frac{1}{2}\text{H}_2\text{O}$.

Deliquescent. Easily sol. in alcohol and H_2O . (Vogel, Z. anorg. 1903, 35. 403.)

Sat. solution of $\text{LiNO}_2 + \frac{1}{2}\text{H}_2\text{O}$ in H_2O contains at:

65°	81.5°	91°	96°	92.5°
63.8	68.7	72.4	91.8	94.3% LiNO_2 .

(Oswald.)

+ H_2O . Very sol. in H_2O , readily forming supersat. solutions. Very sol. in abs. alcohol. (Ball, Chem. Soc. 1913, 103. 2133.)

100 pts. H_2O dissolve at:

0°	10°	20°
125	156	189 pts. $\text{LiNO}_2 + \text{H}_2\text{O}$.

30°	40°	50°
242	316	459 pts. $\text{LiNO}_2 + \text{H}_2\text{O}$.

LiNO_2 , $\text{H}_2\text{O} + \text{Aq}$ sat. at 19° contains LiNO_2 and has sp. gr. = 1.3186. (Oswald, ch. 1914, (9) 1. 61.)

100 g. H_2O dissolve 78.5 g. $\text{LiNO}_2 + \text{AgNO}_3$ at 14° . (Oswald.)

Lithium mercuric nitrite, LiNO_2 , $\text{Hg}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

(Ray, Chem. Soc. 1907, 91. 2033.)

4LiNO_2 , $\text{Hg}(\text{NO}_2)_2 + 4\text{H}_2\text{O}$. Ext. deliquescent. (Ray.)

Magnesium nitrite, $\text{Mg}(\text{NO}_2)_2 + 2\text{H}_2\text{O}$.

Deliquescent, and sol. in H_2O . & decomp. by boiling. Easily sol. in alcohol. (Hampe, A. 125. 334.)

Insol. in absolute alcohol. (Fischer, + $3\text{H}_2\text{O}$. Sol. in H_2O and absolute. Very deliquescent. (Vogel, Z. anorg. 35. 397.)

Magnesium osmium nitrite.

See Osminitrite, magnesium.

Magnesium potassium nitrite.

Deliquescent, and easily sol. in H_2O in alcohol. (Lang.)

Magnesium silver nitrite.

Sol. in H_2O with decomp. (Spiegel, 1895, 19. 1423.)

Manganous nitrite.

Deliquescent, and sol. in H_2O . (Lich.) Not obtained in a solid state; solution decomp. on evaporation. Pogg. 118. 290.)

Mercurous nitrite, $\text{Hg}_2(\text{NO}_2)_2$.

Sol. in H_2O with partial decomp. to $\text{Hg}(\text{NO}_2)_2$. (Ray, A. 1901, 316. 252)

Sol. in cold conc. HNO_3 . Very sol. in cold dil. HNO_3 . (Ray, Chem. Soc. 71. 339.)

Decomp. by boiling H_2O and by H_2SO_4 . (Ray, Z. anorg. 1896, 12. 31 + H_2O . Slowly decomp. by H_2O . Chem. Soc. 1897, 71. 340.)

Mercuric nitrite, basic, $\text{Hg}(\text{NO}_2)_2 + \text{H}_2\text{O}$.

Ppt. (Lang.)

12HgO , $5\text{N}_2\text{O}_3 + 24\text{H}_2\text{O}$. (Ray, C. 1897, 71. 341.)

Mercuric nitrite, $\text{Hg}(\text{NO}_2)_2$.

Deliquescent. Partly sol. in boiling H_2O but the greater part is decomp. in HNO_3 . (Ray, Proc. Chem. Soc. 190

Mercuromercuric nitrite, basic.

a. $9\text{Hg}_2\text{O}$, 4HgO , $5\text{N}_2\text{O}_3 + 8\text{H}_2\text{O}$.

β. Hg_2O , 2HgO , $\text{N}_2\text{O}_3 + 2\text{H}_2\text{O}$.

(Ray, Chem. Soc. 1897, 71. 341.)

Mercurous nitrite, $\text{Hg}(\text{NO}_2)_2$,

in H_2O . Insol. in alcohol.

Obtained from $\text{K}_2\text{Hg}(\text{NO}_2)_4$,
leaving a small excess of KNO_2 .
 H_2O . (Rosenheim, Z. anorg.

$+ \text{H}_2\text{O}$. Sol. in H_2O . For-
is incorrect. (Rosenheim, Z.
. 172.)

Sodium nitrite, $\text{Na}_2\text{Hg}(\text{NO}_2)_4$.

copic. Decomp. by hot H_2O .
anorg. 1901, 28. 173.)

deliquescent. (Ray, Chem. Soc.

3NaNO_2 . Decomp. by H_2O .
oc. 1907, 91. 2032.)

Mercuric nitrite, $3\text{Hg}(\text{NO}_2)_2$,
 $+ 5\text{H}_2\text{O}$.

H_2O . (Ray, Chem. Soc. 1910,

Mercuric hydrazine, $\text{Hg}(\text{NO}_2)_2$, N_2H_4 .
up. by H_2O . (Hofmann and
99, 305. 215.)

Nickelous nitrite, 2NiO , N_2O_3 .
e, A. 125. 343.)

Nickelous nitrite, $\text{Ni}(\text{NO}_2)_2$.

and alcohol. (Lang, J. B. 1862.

Nickelous nitrite, $\text{Ni}(\text{NO}_2)_2$, 4KNO_2 .
sol. in H_2O . (Fischer, Pogg.
emely sol. in H_2O . (Hampe,
insol. in absolute alcohol.

Nickelous strontium nitrite, $\text{Ni}(\text{NO}_2)_2$,
 $(\text{NO}_2)_2$.

l, easily sol. in hot H_2O .

Nickelous ammonia, $\text{Ni}(\text{NO}_2)_2$, 4NH_3 .

H_2O . Decomp. on standing or
sol. in alcohol. Can be re-
dissolving in $\text{NH}_4\text{OH} + \text{Aq}$,
ich absolute alcohol. (Erd-
395.)

NH_3 . Decomp. in the air
, 4NH_3 . (Ephraim, B. 1913,

Osmium nitrite, $\text{Os}(\text{NO}_2)_4$.

bert, C. R. 1905, 140. 587.)

Osmium nitrite with MnO_2 .

te, M.

Osmium nitrite with MnO_2 .

ite, M.

Osmium oxynitrite with MnO_2 .

See Osmyloxynitrite, M.

Osmium nitrite ammonia, $\text{OsO}_2(\text{NO}_2)_4$, 4NH_3 .
(Wintrebert, A. ch. 1903, (7) 28. 56.)

Palladium nitrite with MnO_2 .

See Palladonitrite, M.

Platinous hydrogen nitrite, $\text{H}_2\text{Pt}(\text{NO}_2)_4$.

See Platonitrous acid.

Platinous nitrite with MnO_2 .

See Platonitrite, M.

Potassium nitrite, KNO_2 .

Deliquescent. Sol. in H_2O .

Pure KNO_2 is not deliquescent. (Oswald,
A. ch. 1914, (9) 1. 32.)

Sol. in about $1\frac{1}{2}$ its wt. of H_2O . (Divers,
Chem. Soc. 1899, 75. 86.)

100 pts. H_2O dissolve at:

0°	10°	20°	30°	40°
281	291	302	313	325

pts. KNO_2 ,

50°	60°	70°	80°	90°
337.5	351	365	380	396

pts. KNO_2 ,

100°	110°	120°	130°
413	432	451	473

pts. KNO_2 .

Bpt. of sat. $\text{KNO}_2 + \text{Aq}$ is 132° at 758.5 mm.
pressure.

(Oswald, A. ch. 1914, (9) 1. 58.)

Sp. gr. of $\text{KNO}_2 + \text{Aq}$ at 17.5° containing:

10	20	30	40% KNO_2
1.049	1.126	1.203	1.295

50	60	70	74.5% KNO_2
1.377	1.491	1.599	1.646

(Oswald.)

100 g. H_2O at 13.5° dissolve 18 g. $\text{KNO}_2 +$
2.36 g. AgNO_2 ; at 25° , 23.1 g. $\text{KNO}_2 + 5.3$ g.
 AgNO_2 with excess of AgNO_2 .

100 g. H_2O at 13.5° dissolve 276 g. $\text{KNO}_2 +$
26.3 g. AgNO_2 ; at 25° , 279 g. $\text{KNO}_2 + 39.3$ g.
 AgNO_2 with excess of KNO_2 . (Oswald.)

See also under AgNO_2 .

Very sol. in liquid NH_3 . (Franklin, Am.
Ch. J. 1898, 20. 829.)

Deliquesces in 90% alcohol; insol. in cold
94% alcohol. More sol. in H_2O than KNO_3 ,
but less sol. in alcohol. (Fischer.)

Ppt. from its conc. aq. solution by the
addition of methyl alcohol. Addition of
ethyl alcohol to a conc. aq. solution of KNO_2
causes separation into two layers, of which
the lower aq. solution contains 71.9% KNO_2 ,
while the upper alcoholic layer contains 6.9%
 KNO_2 . (Donath, Ch. Z. 1911, 35. 773.)

Very sl. sol. in acetone. (Krug and M'El-
roy, J. Anal. Ch. 6. 184.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)
Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3602.)

Potassium rhodium nitrite, 6KNO_2 , $\text{Rh}_2(\text{NO}_2)_6$.
See Rhodonitrite, potassium.

Potassium ruthenium nitrite.
See Ruthenonitrite, potassium.

Potassium silver nitrite, KNO_2 , $\text{AgNO}_2 + \frac{1}{2}\text{H}_2\text{O}$.
Completely sol. in a little H_2O , but decomp. by more H_2O . Sol. in $\text{KNO}_2 + \text{Aq}$ without decomp. Insol. in alcohol. (Lang.)

Potassium strontium nitrite, 2KNO_2 , $\text{Sr}(\text{NO}_2)_2$.
Sol. in H_2O ; insol. in alcohol. (Lang, Pogg. 118. 293.)

Potassium zinc nitrite, 2KNO_2 , $\text{Zn}(\text{NO}_2)_2 + \text{H}_2\text{O}$.
Deliquescent. Easily sol. in H_2O . (Lang, J. B. 1862. 101.)
 $\text{K}_2\text{Zn}(\text{NO}_2)_6 + 3\text{H}_2\text{O}$. Very hygroscopic. Decomp. by H_2O . (Rosenheim, Z. anorg. 1901, 28. 174.)

Rhodium nitrite with MNO_2 .
See Rhodonitrite, M.

Rubidium nitrite, RbNO_2 .
Deliquescent; very sol. in H_2O ; sl. sol. in hot alcohol, almost insol. in acetone. (Ball, Chem. Soc. 1913, 103. 2131.)

Ruthenium nitrite with MNO_2 .
See Ruthenonitrite, M.

Silver nitrite, AgNO_2 .
Sol. in 120 pts. cold H_2O (Mitscherlich), in 300 pts. (Fischer), and more abundantly in hot H_2O .
1 l. H_2O dissolves 3.1823 g. or 0.02067 g. mols. at 18° . (Naumann and Rücker, B. 1905, 38. 2294.)
1 litre H_2O dissolves at—

0°	0.0113 mol. AgNO_2
8°	0.0159 " "
14°	0.0189 " "
16°	0.0203 " "
18°	0.0216 " "
25°	0.0260 " "
33°	0.0370 " "

(Pick and Abegg, Z. anorg. 1906, 51. 3.)

1 l. H_2O dissolves 3.609 g. AgNO_2 at 21° . (Oswald, A. ch. 1914, (9) 1. 33.)

Solubility in H_2O at t° .	
t°	% AgNO_2
1	0.1589
15	0.2752
25	0.4125
35	0.6016
51	1.0240
60	1.3625

(Creighton and Ward, J. Am. Ch. 1915, 37. 2335.)

Solubility in $\text{AgNO}_3 + \text{Aq}$ at 11°	
Mols. AgNO_3 per l. of the solution	Mols. AgNO_2 per l.
0.	0.0207
0.0026	0.0194
0.0052	0.0194
0.0103	0.0164
0.0207	0.014
0.0413	0.011
0.0827	0.009

(Abegg and Pick, B. 1905, 38. 1263.)

1 l. 0.2-N $\text{NaNO}_2 + \text{Aq}$ dissolves AgNO_2 at 25° . (Ley and Schaefer, 39. 1263.)
1 l. sat. $\text{KNO}_2 + \text{Aq}$ dissolves 26% at 13.5° . (Oswald, A. ch. 1914, (9)

Solubility in salts + Aq at 25°		
Salt	Conc. of the salt mols. per l.	G. AgNO_2 of a
		0.
AgNO_3	0.00258	0.
	0.00588	0.
	0.01177	0.
	0.02355	0.
	0.04710	0.
KNO_3	0.00258	0.
	0.00588	0.
	0.01177	0.
	0.02355	0.
	0.04710	0.

(Creighton and Ward, J. Am. Ch. 1915, 37. 2336.)

See also under KNO_2 .
 $\text{AgNO}_2 + \text{NaNO}_2$.
1 l. 0.02 N- $\text{NaNO}_2 + \text{Aq}$ dissolve AgNO_2 at 25° . 0.2-N NaNO_2 , 3.016, (Ley and Schaefer, B. 1906, 39. 1263.)
100 g. H_2O sat. with AgNO_2 and contain 10.9 g. AgNO_2 and 78.3 g. at 14° . (Oswald.)

sol. in liquid NH₃. (Franklin, Am. 898, 20. 829.)
in alcohol.
n acetone. (Eidmann, C. C. 1899, 1; Naumann, B. 1904, 37. 4328.)
ts. acetonitrile dissolve 23 pts. at ord. 0 pts. at 81.6°. (Scholl and Steinkopf, 39. 4393.)
l. in methyl acetate. (Bezold, Dis- 16.)
in ethyl acetate. (Hamers, Dissert. aumann, B. 1910, 43. 314.)

odium nitrite, AgNO₃, NaNO₃.
letely sol. in a little H₂O, but decomp. H₂O. (Fischer.)
I₂O. (Oswald, A. ch. 1914, (9) 1.

itrite ammonia, AgNO₃, NH₃.
l. in H₂O; less sol. in alcohol; nearly ether. (Reychler, B. 16. 2425.)
)₂, 2NH₃. (Reychler.)
)₂, 3NH₃. Deliquescent. Sol. in H₂O. er.)

nitrite, NaNO₂.
eliquescent. Very sol. in H₂O.
sol. in H₂O than NaNO₃, but less in
H₂O dissolve 5 pts. NaNO₂ at 15°. Chem. Soc. 1899, 75. 86.)
H₂O dissolve 83.25 g. NaNO₂ at 15°. atowski and Roszkowski, J. phys. Ch. 146.)

ts. H₂O dissolve at:
° 20° 30° 40°
8 84 91.5 98.5 pts. NaNO₂,
° 70° 80°
16 125.5 136 pts. NaNO₂,
° 110° 120°
0.5 178 198.5 pts. NaNO₂.

of sat. NaNO₂+Aq=128° at 761.5
asure. Sat. solution at 20° has a sp. 585. (Oswald, A. ch. 1914, (9) 1.

olubility in NaNO₂+Aq at t°.

	100 pts. H ₂ O dissolve	
	NaNO ₂	NaNO ₃
	73	0
	68.5	19
	67.1	36.3
	64.9	41.7*
	50.3	46.8
	30.2	55.4
	0	74.2

Solubility in NaNO₂+Aq at t°.—Continued.

t°	100 pts. H ₂ O dissolve	
	NaNO ₂	NaNO ₃
21	84.75	0
	81.1	9.6
	79.7	23.5
	73.8	50.8
	73.1	54.5*
	64.2	56.7
	46.8	62.8
	21.6	74.7
	0	89.3
52	108.8	0
	107.9	6.7
	104.3	20.6
	101.8	34.5
	99.5	43.2
	98.0	62.6*
	97.8	82.0
	65.2	88.0
	44.2	92.9
	27.2	101.4
	14.7	109
	0	118
65	120.7	0
	111.5	34.8
	108.5	62.8
	107.8	90.6*
	78.3	96
	49.5	104.1
	28.4	113.4
	14.7	121.4
	0	131
81	137.1	0
	125.7	38.8
	122.7	69.8
	122.6	101.0*
	79.1	111.5
	50.0	121.0
	27.2	131.7
	0	150
92	149.7	0
	141.2	23.6
	134.6	57.6
	132.3	107.8*
	60.2	130.6
	30.3	145.0
	0	163.5
103	166	0
	153.3	33.2
	148.8	58.8
	142.4	116.0*
	100.0	126.8
	60.1	142.9
	0	181.2

* Both salts in solid phase.
(Oswald, A. ch. 1914, (9) 1. 71.)

Solubility in H_2O is decreased by presence of Na_2SO_4 . 100 pts. H_2O dissolve 11.8 pts. Na_2SO_4 + 53.9 pts. $NaNO_2$. (Oswald.)

Very sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Neither dissolved nor attacked by liquid NO_2 . (Frankland, Chem. Soc. 1901, 79. 1361.)

Sol. in warm 90% alcohol. (Hampe, A. 125. 336.)

100 pts. absolute methyl alcohol dissolve 4.43 pts. at 19.5° ; 100 pts. absolute ethyl alcohol dissolve 0.31 pt. at 19.5° . (de Bruyn, Z. phys. Ch. 10. 783.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Strontium nitrite, $Sr(NO_2)_2$.

Very sol. in H_2O , and very sl. sol. in boiling alcohol. (Lang, Pogg. 118. 287.)

Easily sol. in 90% alcohol. (Hampe, A. 125. 340.)

+ H_2O . Hygroscopic. 100 ccm. of the sat. solution contain 62.83 g. $Sr(NO_2)_2$ + H_2O at 19.5° . (Vogel, Z. anorg. 1903, 35. 393.)

100 pts. H_2O dissolve at:

0°	10°	20°	30°
58.9	67.6	75.5	84 pts. $Sr(NO_2)_2$ + H_2O ,

40°	50°	60°	70°
94	105	116	130 pts. $Sr(NO_2)_2$ + H_2O ,

80°	90°	100°
145	162	182 pts. $Sr(NO_2)_2$ + H_2O .

Bpt. of sat. solution is 112.5° at 763 mm. pressure. The sat. solution at 19° contains 39.3% $Sr(NO_2)_2$ and has sp. gr. at $19^\circ/0^\circ = 1.4461$. (Oswald, A. ch. 1914, (9) 1. 64.)

Solubility in alcohol. 100 ccm. of the solution in 90% alcohol contain 0.42 g. $Sr(NO_2)_2$ + H_2O at 20° . 100 ccm. of the solution in absolute alcohol contain 0.04 g. $Sr(NO_2)_2$ + H_2O at 20° . (Vogel, Z. anorg. 1903, 35. 393.)

Thallous nitrite, $TlNO_2$.

Sol. in H_2O . Ppt. from solution in H_2O by absolute alcohol. (Vogel, Z. anorg. 1903, 35. 404.)

Very sol. in H_2O ; insol. in alcohol. (Ball, Chem. Soc. 1913, 103. 2131.)

Zinc nitrite, basic, $2ZnO, N_2O_2$. (Hampe, A. 125. 334.)

Zinc nitrite, $Zn(NO_2)_2 + 3H_2O$.

Deliquescent. Sol. in H_2O and alcohol. (Lang, J. B. 1862. 99.)

Nitrous oxide, N_2O .

See Nitrogen monoxide.

Nitroxyl bromide, NO, Br .

Decomp. spontaneously or with (Hasenbach, J. pr. (2) 4. 1.)

Does not exist. (Fröhlich, A. 224.)

Nitroxyl chloride, NO, Cl .

Decomp. by H_2O without evolution. Probably does not exist. (Geuther, 98.)

Nitroxyl fluoride, NO, F .

Absorbed by H_2O with formation of and HF . Decomp. by H_2O , alcohol, ether. (Moissan and Lebeau, C. R. 140. 1624.)

Nitroxypyrosulphuric acid, $(HO)_2S_2O_8(NO_2), H_2O$.

Very deliquescent. Sol. in H_2O w comp. (Weber, Pogg. 142. 602.)

Nitryl chloride, NO, Cl .

See Nitroxyl chloride.

Octamine cobaltic compounds.

The formulæ of the following octamine cobaltic compounds should be reduced half, and they should be classed as tetramine cobaltic compounds. (Jö Z. anorg. 2. 279.)

Octamine cobaltic carbonate, $Co_2(NH_2)_8(CO_3)_4 + 3H_2O$.

Easily sol. in H_2O . (Vortmann and Berg, B. 22. 2654.)

See Carbonatotetramine carbonate
 $Co_2(NH_2)_8O_2(CO_3)_4 + 3H_2O$. Ratly sol. in H_2O .

— — — chloride (?), $Co_2(NH_2)_8(OH)_2 + 2H_2O$.

Ppt.
 $Co_2(NH_2)_8(OH)_2Cl_4, 2HgCl_2$.
 $Co_2(NH_2)_8(OH)_2Cl_4, PtCl_4 + H_2O$.
mann and Blasberg, B. 22. 2654.)

— — — mercuric chloride, $Co_2(NH_2)_8 + 3HgCl_2 + H_2O$.

$Co_2(NH_2)_8Cl_4, HgCl_2$. Difficultly cold H_2O , decomp. on warming. (Vogel, Z. anorg. 1903, 35. 393.)

— — — chlorosulphite, $Co_2(NH_2)_8 + 4H_2O$.

Sol. in H_2O . (Vortmann and Mann, B. 22. 2635.)

— — — chromate,
 $Co_2(NH_2)_8(CrO_4)_2(H_2O)_2 + 2H_2O$.

Sol. in H_2O or acetic acid.
+ $8H_2O$. Sol. in warm H_2O or acetic acid.
 $Co_2(NH_2)_8(CrO_4)_2Cr_2O_7(H_2O)_2 + 8H_2O$.
Easily sol. in H_2O , from which it is precipitated by dil. HNO_3 + Aq. (Vortmann, B. 22. 2635.)

cobaltic nitrate, $\text{Co}_2(\text{NH}_3)_8(\text{NO}_3)_8$, $\cdot 5\text{H}_2\text{O}$.

H_2O ; precipitated by conc. HNO_3 + tmann.)

nitratocarbonate,
 $\text{NH}_3)_8(\text{NO}_3)_2(\text{CO}_3)_2 + \text{H}_2\text{O}$.

l. than other octamine carbonates.
n and Blasberg, B. 22. 2650.)

bonatotetramine cobaltic nitrate.

purpureochloride,
 $\text{NH}_3)_8\text{Cl}_6(\text{H}_2\text{O})_2$.

sol. in H_2O ; partly precipitated from
olution by conc. $\text{HCl} + \text{Aq}$. (Vort-
10. 1451.)

otetramine cobaltic chloride,
 $)_8(\text{OH}_2)\text{Cl}_2$, wick see. (Jörgen-
(2) 42. 211.)

purpureomercuric chloride,
 $\text{NH}_3)_8\text{Cl}_6(\text{H}_2\text{O})_2, 6\text{HgCl}_2$.

in cold, easily in hot H_2O . (Vort-

otetramine cobaltic mercuric chloro-
rgensen, J. pr. (2) 42. 211.)

purpureomercuric hydroxychloride,
 $\cdot \text{H}_{18}(\text{HgCl})_4(\text{HgOH})_4\text{Cl}_6$.

(Vortmann and Morgulis, B. 22.

$\cdot \text{H}_{18}(\text{HgOH})_8\text{Cl}_6$. (V. and M.)

$\cdot \text{H}_{18}(\text{HgOH})_8\text{Cl}_4(\text{OH})_2$. (V. and M.)

purpureomercuriodide, basic,
 $\cdot \text{H}_{18}(\text{HgOH})_8\text{I}_6$.

ann and Borsbach, B. 23. 2805.)

purpureochloroplatinate.

sol. in H_2O . (Vortmann.)

otetramine cobaltic chloroplatinate,
 $)_8(\text{OH}_2)\text{PtCl}_6 + 2\text{H}_2\text{O}$. (Jörgensen,
42. 215.)

roseochloride, $\text{Co}_2(\text{NH}_3)_8\text{Cl}_6(\text{H}_2\text{O})_2$,
 $\cdot 5\text{H}_2\text{O}$, or $4\text{H}_2\text{O}$.

H_2O . (Vortmann, B. 15. 1891.)

otetramine cobaltic chloride.

roseomercuric chloride,
 $\text{NH}_3)_8\text{Cl}_6(\text{H}_2\text{O})_2, 6\text{HgCl}_2 + 3\text{H}_2\text{O}$.

(Vortmann.)

roseomercuric hydroxychloride,
 $\cdot \text{H}_{18}(\text{HgCl})_6(\text{HgOH})_2\text{Cl}_6$.

ann and Morgulis, B. 22. 2647.)

$\cdot \text{H}_{18}(\text{HgOH})_8\text{Cl}_6$. (V. and M.)

$\cdot \text{H}_{18}(\text{HgOH})_8\text{Cl}_4(\text{OH})_2$. (V. and M.)

roseomercuric iodide,

$\cdot \text{H}_{21}(\text{HgI})_3\text{I}_6$.

sol. in HCl or HNO_3 . (Vortmann
ach, B. 23. 2806.)

$\text{Co}_2\text{N}_8\text{H}_{20}(\text{HgI})_4\text{I}_6$. Ppt. (V. and B.)

$\text{Co}_2\text{N}_8\text{H}_{20}(\text{HgI})_4\text{I}_4(\text{OH})_2$. Ppt. (V. and B.)

Octamine cobaltic sulphate,

$\text{Co}_2(\text{NH}_3)_8(\text{OH})_2(\text{SO}_4)_2 + 3\text{H}_2\text{O}$. (?).

Insol. in H_2O or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in
moderately conc. $\text{HCl} + \text{Aq}$. (Vortmann and
Blasberg, B. 22. 2653.)

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_4)_2 + 6\text{H}_2\text{O}$. Sol. in H_2O .
(Vortmann.)

+ $4\text{H}_2\text{O}$. Easily sol. in H_2O .

See **Roseotetramine cobaltic sulphate**.

sulphatocarbonate,

$\text{Co}_2(\text{NH}_3)_8\text{SO}_4(\text{CO}_3)_2 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Vortmann, B. 10. 1458.)

See **Carbonatotetramine cobaltic sulphate**.

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_4)_2\text{CO}_3 + 4\text{H}_2\text{O}$. Sol. in H_2O .
(Vortmann and Blasberg, B. 22. 2650.)

ammonium sulphite,

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_3\text{NH}_4)_2 + 10\text{H}_2\text{O}$.

See **Octamine cobaltisulphite, ammonium**.

Octamine cobaltisulphurous acid.

Ammonium octamine cobaltisulphite,

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_3\text{NH}_4)_2 + 10\text{H}_2\text{O}$.

Sol. in H_2O . (Vortmann and Magdeburg,
B. 22. 2632.)

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_3)_2(\text{SO}_3\text{NH}_4)_2 + 4\text{H}_2\text{O}$.

Ammonium barium — — — — —,

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_3)_6\text{Ba}_2(\text{NH}_4)_2 + 7\text{H}_2\text{O}$.

Ppt. (V. and M.)

Barium — — — — —, $\text{Co}_2(\text{NH}_3)_8(\text{SO}_3)_6\text{Ba}_2 +$
 $7\text{H}_2\text{O}$.

Ppt. (V. and M.)

Cobaltic — — — — —,

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_3)_6\text{Co}_2 + 36\text{H}_2\text{O}$, and
 $24\text{H}_2\text{O}$.

Luteocobaltic — — — — —,

$\text{Co}_2(\text{NH}_3)_8(\text{SO}_3)_6(\text{NH}_3)_{12}\text{Co}_2 + 8\text{H}_2\text{O}$.

Ppt. (V. and M.)

Octamine iridium chloride,

$\text{Ir}_2(\text{NH}_3)_8\text{Cl}_6$.

Very sol. in H_2O . (Palmaer, B. 22. 16.)

Octamine iridium chlorosulphate,

$\text{Ir}_2(\text{NH}_3)_8\text{Cl}_6\text{SO}_4 + 4\text{H}_2\text{O}$.

(Palmaer.)

Osmiamic acid, $\text{H}_2\text{N}_2\text{Os}_2\text{O}_6$, or

$\text{H}_2\text{N}_2\text{Os}_2\text{O}_6(?)$.

Known only in aqueous solution, which is
unstable.

Ammonium osmiamate.

Easily sol. in H_2O or alcohol. (Fritzsche and Struve, J. pr. 41. 97.)

Barium osmiamate, $BaN_2Os_2O_8$.

Moderately sol. in H_2O .

Lead osmiamate.

Ppt. Sol. in acids without decomp.

Lead osmiamate chloride.

Ppt.

Mercurous osmiamate.

Ppt.

Mercuric osmiamate.

Ppt.

Potassium osmiamate, $K_2N_2Os_2O_8$, or $K_2N_2Os_2O_6$.

Sl. sol. in cold, much more easily in hot H_2O . Sl. sol. in alcohol. Insol. in ether.

Silver osmiamate, $Ag_2N_2Os_2O_8$.

Extremely sl. sol. in H_2O or cold HNO_3 + Aq. Sol. in NH_4OH + Aq.

Sodium osmiamate.

Easily sol. in H_2O or alcohol.

Zinc osmiamate, $ZnN_2Os_2O_8$.

Decomp. by H_2O . Nearly insol. in NH_4OH + Aq.

Osmic acid, H_2OsO_4 .

Stable in H_2O containing alcohol. Sol. in HNO_3 or HCl + Aq. Not attacked by H_2SO_4 + Aq. (Moraht and Wischin, Z. anorg. 3. 153.)

100 g. H_2O dissolve 5.88 g. H_2OsO_4 at 15° . (Squire and Cains, Pharm. J. 1905, 74. 720.)

Attacked by liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 828.)

Barium osmate, $BaOsO_4 + H_2O$.

Insol. in H_2O . (Claus, Pogg. 65. 205.)

Calcium osmate, $CaOsO_4$.

Insol. in H_2O . (Freymy, J. pr. 33. 411.)

Lead osmate.

Insol. in H_2O . (Freymy.)

Potassium osmate, $K_2OsO_4 + 2H_2O$.

Sl. sol. in cold, much more sol. in hot H_2O , but is decomp. thereby. Sl. sol. in KNO_3 + Aq. Insol. in dil. or conc. alcohol and ether. Freymy, A. ch. (3) 12. 516.)

Insol. in conc. saline solutions. (Gibbs, Am. J. Sci. (2) 31. 70.)

Sodium osmate, Na_2OsO_4 .

Sol. in H_2O ; insol. in alcohol a (Freymy, l. c.)

Perosmic acid.

See Perosmic acid.

Osminitrous acid.**Ammonium osminitrite, $(NH_4)_2Os 2H_2O$.**

Sol. in H_2O . Decomp. when so warmed. (Wintrebert, C. R. 1905, 1

Barium osminitrite, $BaOs(NO_2)_2 + H_2O$; $+4H_2O$. (Wintrebert.)**Calcium osminitrite, $CaOs(NO_2)_2 + 4H_2O$. (Wintrebert.)****Magnesium osminitrite, $MgOs(NO_2)_2 + 4H_2O$. (Wintrebert.)****Potassium osminitrite, $K_2Os(NO_2)_2$. Very hygroscopic. Very sol. Decomp. by HCl , HBr and HI . (W. A. ch. 1903, (7) 23. 135.)****Silver osminitrite, $Ag_2Os(NO_2)_2 + 2H_2O$. Sl. sol. in H_2O with partial decomp. (Wintrebert, C. R. 1905, 140. 586.)****Sodium osminitrite, $Na_2Os(NO_2)_2 + 4H_2O$. Sol. in H_2O . (Wintrebert.)****Strontium osminitrite, $SrOs(NO_2)_2 + 4H_2O$. (Wintrebert.)****Zinc osminitrite, $ZnOs(NO_2)_2 + 4H_2O$. (Wintrebert.)****Osmyloxynitrous acid.****Ammonium osmyloxynitrite, $(NH_4)_2OsO_3(NO_2)_2$.**

Decomp. by boiling conc. HCl and + Aq. (Wintrebert, A. ch. 1903, (7)

Barium osmyloxynitrite, $BaOsO_3 + 4H_2O$. (Wintrebert.)**Potassium osmyloxynitrite, $K_2OsO_3 + 3H_2O$.**

Sl. sol. in cold H_2O . Aqueous so comp. slowly. Sol. with decomp. in + Aq. (Wintrebert.)

Silver osmyloxynitrite, $Ag_2OsO_3(NO_2)_2$. (Wintrebert.)

<p>osmyloxynitrite, $\text{SrOsO}_2(\text{NO}_2)_2 +$ (Wintrebert.)</p> <p>rous acid.</p> <p>osmylnitrite, $\text{K}_2\text{OsO}_2(\text{NO}_2)_4$. by H_2O and by excess of $\text{KOH} +$ nland, A. ch. 1903, (7) 28. 54.)</p> <p>churous acid.</p> <p>osmisulphite, $[\text{Os}(\text{H}_2\text{O})(\text{SO}_3)_5]\text{K}_5$ H_2O. Rosenheim, Z. anorg. 1899, 21. 144.)</p> <p>hydrogen osmisulphite, $[\text{Os}(\text{H}_2\text{O})(\text{SO}_3)_5]\text{K}_{11}\text{H}_2 + 5\text{H}_2\text{O}$. H_2O without decomp. (Rosenheim.) $[\text{Os}(\text{H}_2\text{O})(\text{SO}_3)_5]\text{K}_6\text{H}_2 + 2\text{H}_2\text{O}$. Sol. in H_2O with- out decomp. (Rosenheim.)</p> <p>osmisulphite, $[\text{Os}(\text{SO}_3)_5]\text{Na}_5 + 8\text{H}_2\text{O}$. Sol. in H_2O. $[\text{Os}(\text{SO}_3)_5]\text{Na}_4 + 4\text{H}_2\text{O}$. Ppt. $[\text{Os}(\text{SO}_3)_5]\text{Na}_3 + 3\text{H}_2\text{O}$. Ppt. (Rosen- heim.)</p> <p>osmium, Os.</p> <p>finely divided and not ignited to a temperature, Os is sol. in $\text{HNO}_3 +$ aqua regia. When ignited it is not at- tacked by any acid. in liquid NH_3. (Gore, Am. ch. J. 828)</p> <p>ammonium comps.</p> <p>osmiumamine comps., $\text{OsO}(\text{NH}_3)_2\text{X}$. osmiumdiamine comps., $\text{OsO}_2(\text{NH}_3)_4\text{X}_2$.</p> <p>bromide with MBr.</p> <p>osmiumosmate, M.</p> <p>dichloride, OsCl_2. Efflu- escent. Sol. in little, but decomp. by O, with pptn. of Os. Sol. in conc. chlorides + Aq with combination and decomp. (Berzelius.) in alcohol and ether. in H_2O. Insol. in HCl and H_2SO_4. in HNO_3 and aqua regia. Slowly sol. in alkali. Insol. in liquid Cl_2. Insol. in formaldehyde. (Ruff, Z. anorg. 455.)</p> <p>trichloride, OsCl_3. Efflu- scopic. Sol. in conc. H_2SO_4, HCl and HNO_3. Sol. in alkali and in NH_4OH. in liquid Cl_2. Easily sol. in alcohol. in ether. (Ruff, Z. anorg. 1910, 65.)</p> <p>oxide, OsO. Sol. in H_2O. (Moraht and Wis- anorg. 3. 153.)</p>	<p>Osmium tetrachloride, OsCl_4. Sol. in a little H_2O, but decomp. by further addition of that solvent. Sol. in conc. $\text{HCl} +$ Aq.</p> <p>Osmium trichloride with MCl. See Chlorosmite, M.</p> <p>Osmium tetrachloride with MCl. See Chlorosmate, M.</p> <p>Osmium sodium chloride, $\text{Na}_2\text{OsCl}_4 + 2\text{H}_2\text{O}$. Very sol. in H_2O and in alcohol. (Rosen- heim, Z. anorg. 1899, 21. 133.)</p> <p>Osmium tetrafluoride, OsF_4. Sol. in H_2O. (Ruff, B. 1913, 46. 948.)</p> <p>Osmium hexafluoride, OsF_6. Decomp. by H_2O and conc. H_2SO_4. Sol. in $\text{NaOH} + \text{Aq}$. (Ruff, B. 1913, 46. 945.)</p> <p>Osmium octofluoride, OsF_8. Sol. in H_2O, but is somewhat hydrolyzed. Sol. in conc. H_2SO_4 with decomp. Sol. in $\text{NaOH} + \text{Aq}$. (Ruff, B. 1913, 46. 944.)</p> <p>Osmium monohydroxide, $\text{OsO}, x\text{H}_2\text{O}$. Insol. in H_2O. Sl. sol. in $\text{KOH} + \text{Aq}$. Slowly but completely sol. in acids. (Ber- zelius.)</p> <p>Osmium dihydroxide, $\text{OsO}_2, \text{H}_2\text{O}$. Sol. in $\text{HCl} + \text{Aq}$ while still moist. Insol. in H_2SO_4 or $\text{HNO}_3 + \text{Aq}$. $+ 2\text{H}_2\text{O}$. Sol. in HCl, HNO_3, or $\text{H}_2\text{SO}_4 +$ Aq while still moist. (Claus and Jacoby.)</p> <p>Osmium sesquihydroxide, $\text{Os}_2\text{O}_3\text{H}_4$. Sol. in acids, and partly sol. in $\text{KOH} + \text{Aq}$. (Claus and Jacoby.)</p> <p>Osmium iodide, OsI_4. Extremely deliquescent. Sol. in H_2O or alcohol, but solution is unstable. (Moraht and Wischin, Z. anorg. 3. 153.)</p> <p>Osmium potassium nitrosochloride, $\text{K}_2\text{Os}(\text{NO})\text{Cl}_4$. Stable in aqueous solution. Only sl. at- tacked by hot HNO_3. (Wintrebert, A. ch. 1903, (7) 28. 132.)</p> <p>Osmium monoxide, OsO. Insol. in H_2O or acids. (Claus and Jacoby.)</p> <p>Osmium dioxide, OsO_2. Insol. in H_2O or acids.</p> <p>Osmium sesquioxide, Os_2O_3. Insol. in acids. (Claus and Jacoby.)</p>
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Osmium trioxide, "Osmic acid," OsO_3 .

See Osmic acid.

Osmium tetroxide, "Perosmic acid," OsO_4 .

Slowly but abundantly sol. in H_2O . Sol. in alcohol and ether with gradual decomposition. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$, the solution undergoing decomposition on heating.

Osmium oxide ammonia, $\text{OsO}_3 \cdot 2\text{NH}_3 + \text{H}_2\text{O}$.

See Oxyosmiumamine hydroxide.

Osmium oxysulphide, $\text{Os}_2\text{S}_7\text{O}_8 + 2\text{H}_2\text{O}$.

Unstable.

$\text{OsSO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O . (v. Meyer, J. pr. (2) 16. 77.)

$\text{Os}_2\text{O}_7\text{S}_2 + \text{H}_2\text{O}$. Decomp. and dissolved by HNO_3 , HCl , or $\text{H}_2\text{SO}_4 + \text{Aq}$. (Moraht and Wischin, Z. anorg. 3. 153.)

Osmium sulphide, Os_2S_3 (?).

(Berzelius.)

Min. *Laurite*. Insol. in all acids, even in aqua regia.

Osmium disulphide, OsS_2 .

Sl. sol. in H_2O ; not more sol. in alkali hydrates or carbonates + Aq . Insol. in alkalies after drying. (Fremy, A. ch. (3) 12. 521.)

Osmium tetrasulphide, $\text{OsS}_4 + x\text{H}_2\text{O}$.

Insol. in alkali sulphides, carbonates, or hydroxides + Aq . Sol. in cold dil. $\text{HNO}_3 + \text{Aq}$. (Claus.)

Osmocyanhydric acid, $\text{H}_4\text{Os}(\text{CN})_6$.

Easily sol. in H_2O and alcohol. Insol. in ether. (Martius, A. 117. 361.)

Barium osmocyanide, $\text{Ba}_2\text{Os}(\text{CN})_6 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O and dil. alcohol. (M.)

Barium potassium osmocyanide,

$\text{BaK}_2\text{Os}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Efflorescent. Sl. sol. in cold, easily in hot H_2O .

Ferric osmocyanide, $\text{Fe}_4[\text{Os}(\text{CN})_6]_3 + x\text{H}_2\text{O}$.

Insol. in H_2O .

Potassium osmocyanide, $\text{K}_4\text{Os}(\text{CN})_6 + 3\text{H}_2\text{O}$.

Moderately sol. in boiling, less in cold H_2O . Insol. in alcohol and ether.

Osmosyl ammonium comps.

See Oxyosmium amine comps.

Osmyl potassium bromide, $\text{K}_2\text{OsO}_3\text{Br}_4 + 2\text{H}_2\text{O}$.

Same properties as the chloride. (Wintrebert, A. ch. 1903, (7) 28. 94.)

Osmyl potassium chloride, $\text{K}_2\text{OsO}_3\text{Cl}_4$.

Very sol. in H_2O . Solution is stable in the presence of a small amt. of HCl comp. by hot conc. HCl .

+ $2\text{H}_2\text{O}$. As the anhydrous salt. (Wintrebert, A. ch. 1903, (7) 28. 86.)

Osmyl ditetramine comps.

See Oxyosmium diamine comps.

Oxamidosulphonic acid.

See Hydroxylamine monosulphonic acid.

Oximidosulphonic acid.

See Hydroxylamine disulphonic acid.

Oxyamidosulphonic acid.

See Hydroxylamine sulphonic acid.

Oxyammonium salts.

See Hydroxylamine salts.

Oxycobaltamines, acid comps.

(Maquenne, C. R. 96. 344.)

Are anhydrooxycobaltamine comps., see. (Vortmann, M. ch. 6. 404.)

Oxycobaltamine chloride,

$\text{Co}_2(\text{NH}_3)_{10}\text{O}\begin{smallmatrix} \text{OH} \\ \text{(OH)} \end{smallmatrix}\text{Cl}_4$.

(Vortmann, M. ch. 6. 404.)

$\text{Co}_2(\text{NH}_3)_{10}\text{O}_2\text{Cl}_4$, $\text{HCl} + 3\text{H}_2\text{O}$. Is anhydrooxycobaltamine chloride, which see.

— **chloronitrate hydrochloride,**

$\text{Co}_2(\text{NH}_3)_{10}(\text{OH})(\text{O.OH})(\text{NO}_2)_2\text{Cl}_4$, $4\text{HCl} + 3\text{H}_2\text{O}$.

Is anhydrooxycobaltamine chloronitrate, which see.

— **chlorosulphate,**

$\text{Co}_2(\text{NH}_3)_{10}\begin{smallmatrix} \text{OH} \\ \text{O(OH)} \end{smallmatrix}(\text{SO}_4)\text{Cl}_4$, 4HCl .

Easily decomp.

— **iodide,** $\text{Co}_2(\text{NH}_3)_{10}\begin{smallmatrix} \text{OH} \\ \text{O(OH)} \end{smallmatrix}\text{I}_4$.

Sl. sol. in H_2O . Decomp. by much H_2O . (Vortmann.)

— **nitrate,** $\text{Co}_2(\text{NH}_3)_{10}(\text{OH})(\text{O.OH})(\text{NO}_3)_2 + \text{H}_2\text{O}$.

Decomp. by H_2O .

$\text{Co}_2(\text{NH}_3)_{10}(\text{OH})(\text{O.OH})(\text{NO}_2)_2$, $\text{HNC} + 2\text{H}_2\text{O}$. Decomp. by H_2O .

— **nitratosulphate,**

$\text{Co}_2(\text{NH}_3)_{10}(\text{OH})(\text{O.OH})(\text{SO}_4)(\text{NO}_3)_2$, 4HNO_3 .

Decomp. at once by H_2O .

Itamine sulphate,
 $(\text{NH}_4)_{10}\text{O}_2(\text{SO}_4)_2, \text{H}_2\text{SO}_4 + \text{H}_2\text{O}.$
sl. sol. in H_2O with decomp.; more
sol. in acidified H_2O . Sol. in acids.
ane, C. R. 96. 344.)
 $\text{H}_2)_{10}\left(\begin{smallmatrix} \text{OH} \\ \text{O(OH)} \end{smallmatrix}\right)(\text{SO}_4)_2 + 3\text{H}_2\text{O}.$
 $\text{H}_2)_{10}\left(\begin{smallmatrix} \text{OH} \\ \text{O(OH)} \end{smallmatrix}\right)(\text{HSO}_4)_4.$ Decomp.
y by $\text{H}_2\text{O}.$
1, $\text{O}_2.$
s. H_2O absorb 4.6 vols. O gas at ord. temp.
(ham.)
27 pts. H_2O at ord. temp. (Pelouse and
s. H_2O dissolve 0.925 vol. O. (Gay-Lussac.)
 H_2O at t° and 760 mm. absorbs V vols.
gas, reduced to 0° and 760 mm.

V	t°	V	t°	V
04114	7	0.03465	14	0.03034
04007	8	0.03389	15	0.02989
03907	9	0.03317	16	0.02949
03810	10	0.03250	17	0.02914
03717	11	0.03189	18	0.02884
03628	12	0.03133	19	0.02858
03544	13	0.03082	20	0.02838

(Bunsen's Gasometry.)
icient of absorption of O by $\text{H}_2\text{O} =$
 $-0.0010899t + 0.000022563t^2.$ (Bun-
Pauli, A. 93. 21.)
icient of absorption of O in H_2O at
.041408; at $12.6^\circ = 0.036011.$ (Timofe-
phys. Ch. 6. 148.)
tion of O by $\text{H}_2\text{O}.$ $\beta_1 =$ "solubility,"
i. e., the amount of gas (reduced to 0° and
0 mm.) which is absorbed by 1 vol. of
liquid when the barometer indicates
0 mm. pressure; $\beta =$ coefficient of ab-
sorption, i. e., amount absorbed by the
liquid when the pressure of the gas itself
without the tension of the liquid amounts
760 mm.; $\beta_1 = \beta \frac{760-f}{760},$ when $f =$ va-
r tension of solvent at $t^\circ.$

β	β_1
0.04890	0.04860
4759	4728
4633	4601
4512	4479
4397	4362
4286	4250
4181	4142
4080	4040
3983	3941
3891	3847
3802	3756
3718	3670
3637	3587
3560	3507

Absorption of O by $\text{H}_2\text{O}.$ —Continued

t°	β	β_1
14	0.3486	0.3431
15	3415	3358
16	3347	3288
17	3283	3220
18	3220	3155
19	3161	3093
20	3102	3031
21	3044	2970
22	2988	2911
23	2934	2853
24	2881	2797
25	2831	2743
26	2783	2691
27	2736	2641
28	2691	2592
29	2649	2545
30	2608	2500
31	2572	2459
32	2537	2419
33	2503	2380
34	2471	2342
35	2440	2306
36	2410	2270
37	2382	2236
38	2355	2203
39	2330	2171
40	2306	2140
41	2280	2107
42	2256	2075
43	2232	2043
44	2209	2012
45	2187	1981
46	2166	1952
47	2145	1922
48	2126	1894
49	2108	1865
50	2090	1837
52	2057	1782
54	2026	1728
56	1998	1674
58	1971	1619
60	1946	1565
62	1921	1508
64	1897	1450
66	1874	1392
68	1853	1332
70	1833	1270
72	1815	1208
74	1799	1144
76	1785	1078
78	1772	1010
80	1761	0939
82	1752	0865
84	1743	0788
86	1736	0707
88	1729	0622
90	1723	0532
92	1717	0437
94	1712	0337
96	1708	0231
98	1704	0119
100	7001	0000

(Winkler, B. 24. 3600.)

Absorption of O by H₂O at t° and 760 mm.
β = coefficient of absorption.

t°	β	t°	β	t°	β
0	0.04961	23	0.03006	46	0.02163
1	4838	24	2956	47	2139
2	4720	25	2904	48	2115
3	4606	26	2855	49	2092
4	4496	27	2808	50	2070
5	4389	28	2762	51	2049
6	4286	29	2718	52	2029
7	4186	30	2676	53	2009
8	4089	31	2635	54	1990
9	3994	32	2596	55	1972
10	3903	33	2558	56	1955
11	3816	34	2521	57	1938
12	3732	35	2486	58	1922
13	3651	36	2452	59	1907
14	3573	37	2419	60	1893
15	3497	38	2387	65	1832
16	3425	39	2356	70	1787
17	3357	40	2326	75	1752
18	3292	41	2297	80	1726
19	3230	42	2269	85	1707
20	3171	43	2241	90	1693
21	3114	44	2214	95	1684
22	3059	45	2188	100	1679

(Bohr and Bock, W. Ann. (2) 44. 318.)

Coefficient of absorption of O by H₂O between 0° and 30° = 0.04890 — 0.0013413t + 0.0000283t² — 0.00000029534t³. (Winkler, l. c.)
Solubility in H₂O at 25° = 0.03080; at 15° = 0.03630. (Geffcken, Z. phys. Ch. 1904, 49. 269.)

Absorption of O₂ by distilled H₂O at t°.
a = ccm. of O₂ absorbed by 1 l. of H₂O at t° and 760 mm.

t°	a	t°	a	t°	a
0	49.24	17	33.21	34	25.19
1	47.94	18	32.58	35	24.85
2	46.65	19	32.01	36	24.52
3	45.45	20	31.44	37	24.20
4	44.31	21	30.91	38	23.89
5	43.21	22	30.38	39	23.59
6	42.15	23	29.86	40	23.30
7	41.15	24	29.38	41	23.02
8	40.19	25	28.90	42	22.75
9	39.28	26	28.42	43	22.49
10	38.37	27	27.94	44	22.24
11	37.51	28	27.51	45	22.00
12	36.75	29	27.08	46	21.77
13	35.98	30	26.65	47	21.55
14	35.26	31	26.27	48	21.34
15	34.55	32	25.90	49	21.14
16	33.88	33	25.54	50	20.95

(Fox, Trans. Faraday Soc. 1909, 5. 74.)

Solubility in H₂O at various pres

V = volume of the absorbing liquid
P = Hg-pressure in metres.
λ = coefficient of solubility.

V	t°	P
33.320 ccm.	23°	0.9595
		1.0941
		1.2883
		1.4976
		1.7638
		2.0638
		2.5011
		3.0402
		3.8675
		4.2504
		4.6301
		5.1360
32.003 ccm.	25.9°	5.6973
		6.1857
		6.7343
		7.3051
		7.7138
		8.1406
		0.8611
		0.9808
		1.0833
		1.2039
		1.4112
		1.6602
		2.3854
		2.6482
		2.8995
		3.2883
		3.9133
		4.2720
		4.6905
		5.0559
		5.6141
		6.0120
		6.5687
		7.1056
		7.4729
		8.1889

(Cassuto, Phys. Zeit. 1904, 5.

Solubility of O in H₂O at 25 (Findlay and Creighton, Bioch. J 294.)
Coefficient of absorption for H₂O at 15°; 0.03375 at 15.3°; 0.03330 (Müller, Z. phys. Ch. 1912, 81. 494

Solubility in H_2O at t° .Solubility of atmospheric O_2 in H_2O at t° .

lree	t°	lree
10.26	13	7.51
9.99	14	7.36
9.73	15	7.21
9.48	16	7.07
9.25	17	6.93
9.02	18	6.80
8.80	19	6.67
8.59	20	6.55
8.39	21	6.43
8.20	22	6.32
8.02	23	6.21
7.84	24	6.10
7.67	25	6.00

, Zeit. angew. Ch. 1913, 26. 714.)

Solubility of atmospheric O_2 in mixtures of H_2O with sea water diminishes regularly with the proportion of sea water present. (Soc. Chem. Ind. 1904, 23. 359.)

Volume of O_2 absorbed by 1 l. of sea water from a free dry atmosphere of 760 mm. pressure.

4°	8°	12°	16°	20°	24°	28°
9.26	8.40	7.68	7.08	6.57	6.14	5.75
8.85	8.04	7.36	6.80	6.33	5.91	5.53
8.45	7.68	7.04	6.52	6.07	5.67	5.31
8.04	7.33	6.74	6.24	5.82	5.44	5.08
7.64	6.97	6.43	5.96	5.56	5.20	4.86
7.23	6.62	6.11	5.69	5.31	4.95	4.62

Trans. Faraday Soc. 1909, 5. 77.)

Amount absorbed from the air, see also air, p. 1.

Absorption of O_2 by acids + Aq.
Content in gram-equivalents per litre of solution.

 HNO_3 + Aq.

S_{25°	S_{15°
0.03021	0.03478
0.03016	0.03490
0.02954	0.03354
0.02963	0.03365
0.02853	0.03175
...	0.03166

Absorption of O_2 by acids + Aq.—Continued. HCl + Aq.

M	S_{25°	S_{15°
0.578	0.02843	0.03431
0.579	0.02960	0.03410
1.170	0.02817	0.03217
1.176	0.02833	0.03109
1.736	0.02733	0.02969
1.982	0.02674	0.02988

 $\frac{H_2SO_4}{2}$ + Aq.

M	S_{25°	S_{15°
0.489	0.02887	0.03390
0.527	0.02875	0.03375
0.977	0.02757	0.03210
1.017	0.02745	0.03217
1.896	0.02545	0.02886
1.829	0.02577	0.02930
2.947	0.02285	0.02584
3.512	0.02198	0.02399
4.951	0.01918	0.02174
5.293	0.01918	0.02067

(Geffcken, Z. phys. Ch. 1904, 49. 269.)

Absorption of O_2 by H_2SO_4 + Aq at t° . α = coefficient of absorption.

Normality of the acid	t°	α
0	20.9	0.0310
4.9	20.9	0.0195
8.9	20.9	0.0155
10.7	21.2	0.0143
20.3	21.1	0.0119
24.8	21.5	0.0103
29.6	20.8	0.0117
34.3	20.9	0.0201
35.8	21.2	0.0275

(Bohr, Z. phys. Ch. 1910, 71. 49.)

Absorption of O_2 by $NaOH$ + Aq.

M = content in gram-equivalents per litre.
S = solubility.

M	S_{25°	S_{15°
0.559	0.02434	0.02777
0.601	0.02424	0.02784
1.033	0.02020	0.02291
1.059	0.01991	0.02262
2.077	0.01295	0.01479
2.089	0.01272	0.01458

Absorption of O ₂ by KOH + Aq.		
M	S 25°	S 15°
0.577	0.02447	0.02791
0.579	0.02435	0.02791
1.157	0.01920	0.02191
1.170	0.01914	0.02181

(Geffcken, Z. phys. Ch. 1904, 49. 270.)

NaCl + Aq with a chlorine content of 1,930 per 100,000 dissolved 82.9% of the amount of O₂ dissolved by distilled H₂O alone. (Clowes, J. Soc. Chem. Ind. 1904, 23. 359.)

Absorption of O₂ by salts + Aq.

M = content in gram-equivalents per litre.
S = solubility.

Absorption of O = by $\frac{K_2SO_4}{2}$ + Aq.

M	S 25°	S 15°
0.499	0.02528	0.02944
0.506	0.02530	0.02922
0.968	...	0.02395
0.970	0.02096	0.02377

Absorption of O₂ by NaCl + Aq.

M	S 25°	S 15°
0.530	0.02598	0.03045
0.535	0.02604	0.03052
1.020	0.02226	0.02601
1.034	0.02202	0.02557
1.880	...	0.01898
1.890	0.01663	0.01904
1.921	0.01654	0.01869

(Geffcken, Z. phys. Ch. 1904, 49. 270.)

Solubility of O₂ in NaCl + Aq.

Data indicate cc. O₂ dissolved per l. at 760 mm. and 0°.

t°	NaCl + Aq 1 g. mol. per l.	NaCl + Aq 2 g. mol. per l.	NaCl + Aq sat. at 20°
0	6.50	3.14	1.27
5	5.80	2.84	1.22
10	5.25	2.59	1.17
15	4.77	2.41	1.12
20	4.39	2.25	1.07
25	4.06	2.13	1.02
30	3.76	2.01	0.97

(Winkler, Z. anorg. 1911, 24. 342.)

Solubility of O₂ in KCN + Aq at 20°.

% KCN 1 10 20 30 50
Coeff. of abs. 0.029 0.018 0.013 0.008 0.003
(McLaurin, J. S. C. I. 1893, 63. 737.)

1 vol. alcohol absorbs 0.28397 vol. O₂ at temperatures between 0° and 24°. (B)

Absorption by alcohol (99.7%) at
β = coefficient of absorption;
β₁ = solubility. (See p. 635.)

t°	β	β ₁
0	0.23370	0.22
1	0.23296	0.22
2	0.23222	0.22
3	0.23149	0.22
4	0.23077	0.22
5	0.23005	0.22
6	0.22934	0.22
7	0.22863	0.22
8	0.22793	0.22
9	0.22724	0.22
10	0.22656	0.22
11	0.22588	0.22
12	0.22521	0.22
13	0.22455	0.22
14	0.22389	0.22
15	0.22324	0.22
16	0.22259	0.22
17	0.22195	0.22
18	0.22132	0.22
19	0.22069	0.22
20	0.22007	0.22
21	0.21946	0.22
22	0.21886	0.22
23	0.21826	0.22
24	0.21767	0.22

(Timofejew, Z. phys. Ch. 6. 1)

Solubility of O₂ in alcohol at 20° and

Wt. % alcohol	Vol. % abs. O ₂	Wt. % alcohol
0.0	2.98	33.33
9.09	2.78	50.0
16.67	2.63	66.67
23.08	2.52	80.0
28.57	2.49	

(Lubarsch, W. Ann. 1889, (2) 87

Solubility of O₂ in methyl alcohol

t°	l.	t°	
0	0.31864	25	0
5	0.30506	30	0
10	0.29005	40	0
15	0.27361	50	0
20	0.25574		

(Levi, Gazz. ch. it. 1901, 31. II,

Solubility of O₂ in ether at 0° = 0.4215. (Christoff, Z. phys. 79. 459.)

solubility of O_2 in acetone at t° .

L.	t°	L.
0.2997	25	0.2127
0.2835	30	0.1935
0.2667	40	0.1533
0.2493	50	0.1057
0.2313		

i, *Gazz. ch. it.* 1901, **31**, II, 513.)

absorption of O_2 by chloralhydrate + Aq.

mp. of the solution.

chloralhydrate in the solution.

coefficient of absorption at t° .

$\beta 15^\circ$ = coefficient of absorption at 15° .

$\beta 20^\circ$ = coefficient of absorption at 20° .

P	βt°	$\beta 15^\circ$
22.9	0.02759	0.02940
28.0	0.02690	0.02800
36.6	0.02590	0.02560
38.6	0.02402	0.02477
51.3	0.02439	0.02339
58.44	0.02350	0.02407
70.0	0.02659	0.02710
80.85	0.03200	0.03300
80.9	0.03140	0.03250
		$\beta 20^\circ$
16.9	0.02795	0.02795
32.0	0.02443	0.02495
52.9	0.02375	0.02325
61.08	0.02390	0.02410
65.5	0.02500	0.02580
71.4	0.02680	0.02730
78.0	0.03090	0.03280

ller, *Z. phys. Ch.* 1912, **81**, 499.)

orption of O_2 by glycerine + Aq.

mp. of the solution.

glycerine in the solution

coefficient of absorption at t°

$\beta 15^\circ$ = coefficient of absorption at 15° .

P	βt°	$\beta 15^\circ$
20.5	0.02904	0.02742
25.0	0.02654	0.02521
37.3	0.02038	0.02022
45.0	0.01800	0.01744
52.0	0.01623	0.01570
71.5	0.01010	0.00950
88.5	0.00903	0.00886

(Müller.)

Absorption of O_2 by glucose + Aq.

t° = temp. of the solution.

P = % glucose in the solution.

βt° = coefficient of absorption at t° .

$\beta 20^\circ$ = coefficient of absorption at 20° .

t°	P	βt°	$\beta 20^\circ$
21.2	10.84	0.02650	0.02690
21.5	20.7	0.02202	0.02250
19.9	33.8	0.01814	0.01815
20.5	51.9	0.01378	0.01390
21.7	58.84	0.01221	0.01250

(Müller.)

Absorption of O_2 by sucrose + Aq

t° = temp. of the solution.

P = % sucrose in the solution.

βt° = coefficient of absorption at t° .

$\beta 15^\circ$ = coefficient of absorption at 15° .

t°	P	βt°	$\beta 15^\circ$
15.3		0.03375	0.03400
16.2		0.03330	0.03397
18.0	12.1	0.02911	0.02969
15.6	24.38	0.02367	0.02396
16.6	28.44	0.02113	0.02181
15.6	42.96	0.01582	0.01600
16.2	49.25	0.01348	0.01380
17.2	50.0	0.01302	0.01359

(Müller.)

Abundantly absorbed by oil of turpentine. Oil of turpentine absorbs its own vol. O when exposed two weeks to the air, but does not give it off on boiling. (Brandes.)

Absorbed by other oils, but this is decomposition rather than absorption, as the oils are oxidized. (See Storer's Dict.)

100 vols. arterial blood dissolve 10-13 vols. O . (Magnus.)

Coefficient of absorption for petroleum = 0.202 at 20° ; 0.229 at 10° . (Gniewass and Walfisz, *Z. phys. Ch.* **1**, 70.)

The author examined the solubility of O_2 and N_2 at low temp. in alcohols, ethers, acetone, $CHCl_3$, petroleum, benzene and various inorganic liquids; at low temp. the solubility of the N_2 increases at the same rate as that of the O_2 . (Claude, *C. R.* 1900, **131**, 448.)

Oxydimercuri ammonium bromate,

$(NH_2OH)_2BrO_4$.

(Rammelsberg, *Pogg.* **55**, 82.)

— carbonate, $(NH_2OH)_2CO_3 + \frac{1}{2}H_2O$.

Insol. in H_2O . Decomp. by HCl + Aq. only when conc. Not decomp. by boiling KOH + Aq. Decomp. by KI or K_2S + Aq. (Hirzel.) + H_2O . As above. (Hirzel.)

Oxydimercuriammonium chloride,
 $(\text{NHg}_2\text{OH}_2)\text{Cl}$.

Is dimercuriammonium chloride, $\text{NHg}_2\text{Cl} + \text{H}_2\text{O}$, which see.

— **oxytrimercuriammonium chloride,**
 $(\text{NHg}_2\text{OH}_2)\text{Cl}$, $(\text{NHg}_2\text{O}_2\text{H}_2)\text{Cl}$ (?).

Insol. in H_2O . Easily sol. in dil. $\text{HCl} + \text{Aq}$. More difficultly sol. in very dil. H_2SO_4 or $\text{HNO}_3 + \text{Aq}$. Insol. in conc. H_2SO_4 . Sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$, or $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$. (Schmieder.)

— **chromate,** $(\text{NHg}_2\text{OH}_2)_2\text{CrO}_4$.

Not decomp. by $\text{KOH} + \text{Aq}$. (Hirzel, J. B. 1852. 421.)

— **mercuric chromate,** $(\text{NHg}_2\text{OH}_2)_2\text{CrO}_4$, 4HgO , 3CrO_3 .

Decomp. by HNO_3 without going into solution. Easily sol. in HCl . (Hirzel.)

Composition is $(\text{NHg}_2\text{OH}_2)_2\text{O}$, 2CrO_3 , $3[(\text{NH}_4)_2\text{O}$, $2\text{Cr}_2\text{O}_3] = (\text{NHg}_2\text{OH}_2)_2\text{Cr}_2\text{O}_7$, $3(\text{NH}_4)_2\text{Cr}_2\text{O}_7$. (Hensgen, R. t. c. 5. 187.)

Probably $(\text{NHg}_2)_2\text{Cr}_2\text{O}_7$, $3(\text{NH}_4)_2\text{Cr}_2\text{O}_7 + 2\text{H}_2\text{O}$.

— **fluoride, acid,** $(\text{NHg}_2\text{OH}_2)\text{F}$, HF .

(Finkener, Pogg. 110. 632.)

Probably NHg_2F , $\text{HF} + \text{H}_2\text{O}$.

— **hydroxide,** $(\text{NHg}_2\text{OH}_2)\text{OH} = \text{NHg}_2\text{OH} + \text{H}_2\text{O}$.

(Millon's base.) Sl. sol. in H_2O , especially if warm. Sol. in 13,000 pts. H_2O at 17° , and 1700 pts. at 80° . Insol. in alcohol or ether. (Gerresheim, A. 195. 373.)

+ H_2O . Insol. in H_2O or alcohol. Sol. in traces in $\text{NH}_4\text{OH} + \text{Aq}$. Not decomp. by cold $\text{KOH} + \text{Aq}$; sl. decomp. if hot. (Millon.)

— **ammonium iodate,** $(\text{NHg}_2\text{OH}_2)\text{IO}_3$, $2\text{NH}_4\text{IO}_3$.

Insol. in H_2O . (Millon, A. ch. (3) 18. 410.)

— **iodide,** $(\text{NHg}_2\text{OH}_2)\text{I}$.

Sol. in warm $\text{HCl} + \text{Aq}$. Not decomp. by boiling $\text{KOH} + \text{Aq}$. Sol. in warm $\text{KI} + \text{Aq}$. (Rammelsberg, Pogg. 48. 170.)

Correct formula is $\text{NHg}_2\text{I} + \text{H}_2\text{O}$. (Rammelsberg.)

— **nitrate,** $(\text{NHg}_2\text{OH}_2)\text{NO}_3$.

Insol. in H_2O ; not decomp. by boiling $\text{KOH} + \text{Aq}$. Sol. in cold $\text{HCl} + \text{Aq}$, from which it is precipitated by H_2O . Sl. sol. without decomp. in HNO_3 or $\text{H}_2\text{SO}_4 + \text{Aq}$. Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Soubeiran.)

Is dimercuriammonium nitrate, NHg_2NO_3 . (Pesci, Gazz. ch. it. 20. 485.)

— **ammonium nitrate,** $\text{NHg}_2(\text{OH}_2)\text{NO}_3$, $2\text{NH}_4\text{NO}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O . Kane, A. ch. 72. 242.)

Is dimercuriammonium ammonium nitrate, NHg_2NO_3 , $2\text{NH}_4\text{NO}_3 + 2\text{H}_2\text{O}$. (Pesci.)

Oxydimercuriammonium oxide,
 $(\text{NHg}_2\text{OH}_2)_2\text{O}$.

Insol. in H_2O or alcohol; not att. boiling conc. $\text{KOH} + \text{Aq}$. Sol. in hot + Aq , $\text{NH}_4\text{Cl} + \text{Aq}$, $(\text{NH}_4)_2\text{SO}_4$, $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$, $(\text{NH}_4)_2\text{C}_2\text{O}_4 + \text{Aq}$ lon, A. ch. (3) 18. 397.)

— **mercuric phosphate,** $\text{Hg}(\text{NHg}_2\text{OH}_2)_2\text{PO}_4$.

Insol. in H_2O . Slowly sol. in hot Aq ; not decomp. by boiling with H but by KI or $\text{K}_2\text{S} + \text{Aq}$. Sol. in H much hot $(\text{NH}_4)_2\text{HPO}_4 + \text{Aq}$. (Hirzel.)

— **mercuric sulphite,** $(\text{NHg}_2\text{OH}_2)_2\text{HgSO}_3$.

Insol. in H_2O . Sol. in much N Aq . Sol. in $\text{HCl} + \text{Aq}$ with decomp. Insol. in boiling $\text{KOH} + \text{Aq}$. (Hirzel.)

— **sulphate,** $(\text{NHg}_2\text{OH}_2)_2\text{SO}_4$.

Sol. in traces in H_2O . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Kane.)

Insol. in $\text{HNO}_3 + \text{Aq}$. (Hirzel.) Slowly sol. in boiling conc. H_2SO_4 . (Hirzel.)

Insol. in conc., easily sol. in dil. Aq . (Schmieder, J. pr. 75. 147.)

Moderately sol. in much $(\text{NH}_4)_2\text{SO}_4$ + Aq . Not decomp. by boiling $\text{KOH} + \text{Aq}$. (Hirzel.)

Easily decomp. by boiling with H_2O . (Schmieder.)

Does not exist. (Pesci.)

2NH_3 , 2HgO , SO_3 .

See Dimercuriammonium sulphate.

Oxytrimercuriammonium chloride,
 $(\text{NHg}_2\text{O}_2\text{H}_2)\text{Cl}$ (?).

Insol. in H_2O .

— **nitrate,** $(\text{NHg}_2\text{O}_2\text{H}_2)\text{NO}_3$.

Sol. in cold $\text{HCl} + \text{Aq}$, from which it is precipitated by $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in Aq without decomp. Not decomp. by boiling $\text{KOH} + \text{Aq}$. (Pagenstecher.)

Does not exist. (Pesci, Gazz. 485.)

Oxytrimercuridiammonium sulphate,
 2NH_3 , 3HgO , SO_3 .

See Trimercuriammonium sulphate.

Oxytrimercurioxydimercuriammonium sulphate,
 $\text{NH}_2\text{Hg}_2\text{O}_2$, $\text{NH}_2\text{Hg}_2\text{O}_2 > \text{SO}_4$.

Completely sol. in NH_4C $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. Sol. in dil. or conc. Aq , and very dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Insol. in H_2O . (Schmieder.)

Does not exist. (Pesci.)

curiammonium mercuric
(?), $2(\text{NH}_4\text{O}_2)\text{NO}_2$, HgNO_2 (?).
insol. in $\text{HNO}_3 + \text{Aq.}$ Sol. in
Aq. Slowly decomp. by boiling
Gradually sol. in hot conc.
l. (Hirzel.)
exist. (Pesci, Gazz. ch. it. 20.

phonic anhydride,

O_2
 NO_2 (?).

with decomp. (Weber, Pogg.

amine hydroxide (Osmono-
ine hydroxide),

$(\text{OH})_2$.

O_2 . Sl. sol. in acids. Sol. in
When moist, sol. in $\text{NH}_4\text{OH} +$

liamine chloride (Osmyltetra-
chloride), $\text{OsO}_2(\text{N}_2\text{H}_4\text{Cl})_2$.

cold, more easily in hot H_2O .
 $\text{Cl} + \text{Aq.}$ (Gibbs, Am. Ch. J.

**stinate, OsO}_2(\text{N}_2\text{H}_4\text{Cl})_2, PtCl_4 .
 O_2 . (Gibbs.)**

e, OsO}_2(\text{N}_2\text{H}_4\text{OH})_2.

in solution.

OsO}_2(\text{N}_2\text{H}_4\text{NO}_2)_2.

$\text{OsO}_2(\text{N}_2\text{H}_4)_2\text{SO}_4 + \text{H}_2\text{O}$.
Ch. J. 3. 233.)

retted hydrogen (?),

rier, and Goldschmidt has this
ding to Franke (J. pr. (2) 35.
p. slowly by H_2O or alkalies.
um salt, $\text{P}_4\text{H}(\text{OK})$, sol. in H_2O .

**ide, P}_4\text{H}(\text{OH}), HI .
 80° .**

**isulphuric acid, Pt}_2\text{O}_3,
 $\text{H}_2 + 11\frac{1}{2}\text{H}_2\text{O}$.**

(Blondel, A. ch. 1905, (8) 6.

**oxyplatissulphate, Pt}_2\text{O}_3, 3SO_3 ,
 H_2O .**

H_2O . (Blondel.)

**quioxypatissulphate, Pt}_2\text{O}_3,
 $\text{K}_2 + 2\text{H}_2\text{O}$.**

**Sodium sesquioxypatissulphate, Pt}_2\text{O}_3, 3SO_3 ,
 $\text{SO}_4\text{Na}_2 + 8\text{H}_2\text{O}$.**

Very sol. in H_2O . (Blondel.)

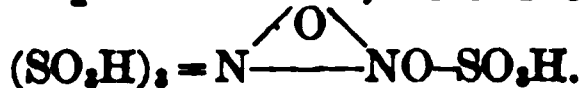
Oxysulphantimonic acid.

See Sulphoxyantimonic acid.

Oxysulpharsenic acid.

See Sulphoxyarsenic acid.

Oxysulphazotic acid, H}_4\text{S}_4\text{N}_2\text{O}_{14} =



Known only in its salts. (Claus, A. 158.
52, 194.)

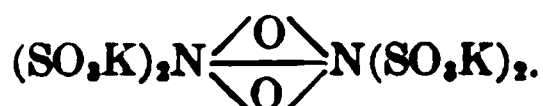
Has formula $(\text{SO}_3\text{H})_2\text{N} \begin{array}{c} \diagup \text{O} \diagdown \\ \text{---} \end{array} \text{N}(\text{SO}_3\text{H})_2$.

(Raschig, A. 241. 161.)

Potassium oxysulphazotate, NO(SO}_3\text{K})_2.

Insol. in alcohol. (Fremy, A. ch. (3) 15.
451.)

According to Raschig the formula is



Very sol. in water, with rapid decomposi-
tion. (Raschig.)

See also Peroxylaminesulphonate, potas-
sium.

Oxysulphotungstic acid.

See Sulphotungstic acid.

Oxysulphovanadic acid.

See Sulphoxyvanadic acid.

Ozone, O}_3.

Not appreciably sol. in H_2O . (Schönbein.)
Imparts its taste and properties to H_2O .
(Williamson.)

Later, Carius (B. 5. 520) found that 1000
vols. H_2O at $1-2.5^\circ$ absorb 5.11 vols. O_3 (red.
to 0° and 760 mm.). He also still later (A.
174. 1) found, by conducting the gas for 9-12
hours through H_2O , that 1000 vols. H_2O
absorb a maximum of 28.160 vols. O_3 . The
ozonized oxygen used contained 3.44 vols. O_3
in 100 vols. O_2 . Since gases are absorbed in
proportion to their partial pressure, which is
very small for the O_3 , the amount of absorp-
tion of water for the gas is very considerable.
Carius calculated the coefficient of absorption
at $+1^\circ$ to be 0.834.

Ozone is *not at all* absorbed by H_2O ; the
 H_2O through which ozone had been passed
gave no reactions for ozone. (Rammelsberg,
B. 6. 603.)

Schöne (B. 6. 1224) corroborates Carius,
and finds 8.81 vols. to 1000 vols. H_2O as a
maximum amount absorbed.

Sol. in H_2O . (Leeds, B. 12. 1831.)

H₂O takes up $\frac{1}{2}$ of its vol. of O₂ at 0° and 760 mm. pressure and $\frac{1}{2}$ of its vol. at 12°, or about 15 times that of oxygen at the same pressure and temp (Mailfert, C. R. 1894, 119. 951.)

Solubility in H₂O at t°.

Temp.	Wt. O ₂ dissolved in 1 l H ₂ O	Wt. O ₂ in gaseous mixture above the solution	Coefficient of solubility of O ₂
0	39.4 mgr.	61.5 mgr.	0.641
6	34.3	61	0.562
11.8	29.9	59.6	0.500
13	28	58.1	0.482
15	25.9	56.8	0.456
19	21	55.2	0.381
27	13.9	51.4	0.270
32	7.7	39.5	0.195
33	4.2	37.6	0.112
47	2.4	31.2	0.077
55	0.6	19.2	0.031
60	0.0	12.3	0.000

(Mailfert, C. R. 1894, 119. 952.)

Solubility of ozone in acidulated H₂O.

Temp.	Coefficient of solubility of ozone	Composition of solution
30°	0.240	1 l. H ₂ O + 0.7 cc. H ₂ SO ₄
33	0.224	
42.7	0.174	
49	0.156	1 l. H ₂ O + 0.9 cc. "
57	0.096	1 l. " + 0.3 cc. "

(Mailfert.)

0.00002 pt. by weight is sol. in 1 pt. by weight H₂O at ordinary temp. and pressure (Ladenburg, B. 1898, 31. 2510)

The solution of O₃ in H₂O cannot be brought into equilibrium, because when the gas is blown through the liquid, a portion is continually decomposed, although the concentration remains constant. (Ingles, Chem. Soc. 1903, 83. 1012)

About 10 mg. ozone are sol. in 1 l. H₂O at +2°; 1.5 mg. ozone are sol. at +28°. (Moufang, C. C. 1911, II. 1674.)

Solubility in 0.1 N H₂SO₄.

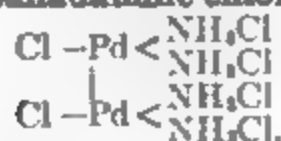
C solution: C gas = 0.23 at 20°; 0.44 at 0°. (Luther, Z Elektrochem 1905, 11. 833)

The absorption coefficient of the gas in 0.1 N H₂SO₄ solution is 0.487. (Rothmund, C. C. 1912, I. 1261.)

Sol. in H₂C₂O₄ + Aq. Jeremin, B. 11. 988.)

Completely absorbed by oil of turpentine and oil of cinnamon. (Soret, A. ch. (4) 17. 113)

Dipalladamine chloride, Cl₂Pd₂(NH₄)₂Cl₄ =



Sl. sol. in H₂O. (Dewille and Debray, C. R. 86. 296.)

Palladium, Pd.

Not attacked by H₂O. Sl. attad HC + Aq, but Pd sponge or filings are dissolved in warm HCl + Aq, with air. HNO₃ + Aq of 1.2 sp. gr. dissol slightly, but it is easily sol. in HNO₃ 1.35 sp. gr. (Rose.)

Easily sol. in aqua regia. Sl. sol. in but insol in dil. HI + Aq. Sol. in concing H₂SO₄. Sol. in boiling FeCl₃ + Aq in HBr + Aq with a little HNO₃.

Insol. in liquid NH₃. (Gore, Am 1898, 20. 828.)

Palladium ammonium compounds.

See—

Dipalladamine comp., Cl₂Pd₂(NH₄)₂

Palladodiamine " Pd(NH₄)₂Cl₂

Palladosamine " Pd(NH₄)₂Cl₂

Palladium dibromide.

Not known in pure state

Palladium bromide with MBr.

See Bromopalladite, M.

Palladium tetrabromide with MBr.

See Bromopalladate, M.

Palladous phosphorus bromide, Pd₂I

Properties as the corresponding (Strecker, B. 1909, 42. 1776)

Palladium subchloride, Pd₂Cl₃.

Deliquescent. Decomp. by H₂O, KI, or NH₄OH + Aq. (Kane)

Sol. in acetone. (Naumann, B. 1 4328.)

Palladium dichloride, PdCl₂.

Slowly but completely sol. in H₂O + 2H₂O. Not deliquescent wh slowly sol in H₂O. Much more so containing HCl.

Sol. in acetone. (Eidmann, C. C. 1014.)

Sol. in ethyl acetate. (Naumann, 37. 3601.)

Palladium dichloride with MCl.

See Chloropalladite, M.

Palladium tetrachloride with MCl.

See Chloropalladate, M.

Palladous phosphorus chloride, Pd₂Cl

Decomp. by H₂O into deliquescent PdCl₂. Decomp. by alcohol. (Fink 115. 176.)

Decomp. by H₂O. Sol. in CHCl₃. Insol. in ligroin and CCl₄. (Strecker 42. 1775)

PdCl₂, 2PCl₃. Sol. in C₆H₆, and by H₂O (Fink.)

chloride carbon monoxide, PdCl₂,
by heat. (Fink, C. R. 1898, 126.)

CO. Decomp. by H₂O. Sol. in
(.)

fluoride, PdF₂.

H₂O or HF + Aq. Sl. sol. while
H₂OH + Aq; insol. after drying,
-Aq. Insol. in boiling NaF or
(Berzelius.)

hydride, Pd₂H (?).

hydroxide, PdO, xH₂O (?).

. in acids or excess of alkali
and carbonates + Aq. Sol. in hot
(Rose.)
Na₂B₄O₇, and Na₂HPO₄ + Aq.

oxide, PdO₂, xH₂O.

. in acids. Sol. in conc. HCl +
decomp. With dil. HCl + Aq, Cl₂,
(Berzelius.)

hydroxide hydroxylamine,
O₂(OH)₂.

H₂O. Sol. in dil. HCl and in dil.
sel, A. 1907, 351. 445.)

dide, PdI₂.

H₂O. Can be detected as a brown
presence of 400,000 pts. H₂O.

HI + Aq. Easily sol. in KI + Aq.
[. ch. méd. 11. 57.)

il. HCl + Aq, but slightly sol. in
ns. (Fresenius.)

hot conc. HNO₃ + Aq. Sol. in
Cl₂ + Aq, Br₂ + Aq, I₂ + Aq, and
also in HCN, and MCN + Aq.

H₂SO₄, HCl, H₃PO₄, HNO₃, or
aq, or in the K, Na, or NH₄ salts
ds. Insol. in CuCl₂, ZnCl₂, or
+ Aq. Insol. in KBr + Aq ex-
cess of a free mineral acid, but not
Insol. in sugar or starch + Aq,
alcohol, ether, or oil of lemon.
l. in urine. Easily sol. in NH₄OH
when dil., with evolution of heat
position. (Kersten, A. 87. 28.)

cohol or ether.

ethyl acetate. (Naumann, B.
O.)

potassium iodide.

lladite, potassium.

boxide, Pd₂O.

by acids into palladious salt and
Phil. Trans. 1842, 1. 276.)

acids, even boiling aqua regia.
5. 220.)

Palladous oxide, PdO.

Slowly sol. in acids by boiling. (Wöhler,
A. 174. 160.)

Palladic oxide, PdO₂.

Very sl. attacked by acids.

Palladopalladic oxide, 4PdO, PdO₂.

Not attacked by aqua regia. (Schneider,
Pogg. 141. 528.)

**Palladous oxychloride, 3PdO, PdCl₂ +
4H₂O(?).**

Sol. in dil. acids. (Kane.)

**Palladous oxychloride ammonia, PdO, PdCl₂,
6NH₃(?).**

Sol. in HCl + Aq.

3PdO, PdCl₂, 2NH₃ + 3H₂O (?). Ppt.
(Kane.)

Palladium selenide, PdSe.

Insol. in HNO₃ and aqua regia. (Rössler,
A. 180. 240.)

Palladium subsulphide, Pd₂S.

Not attacked by acids except aqua regia,
which attacks slightly. (Schneider, Pogg.
141. 530.)

Palladium monosulphide, PdS.

Insol. in H₂O or (NH₄)₂S + Aq. Sol. in
HCl + Aq. Pptd. in presence of 10,000 pts.
H₂O. (Fellenberg, Pogg. 50. 65.)

Sol. in potassium thiocarbonate + Aq. (Ro-
senblatt, Z. anal. 26. 15.)

A sol. colloidal form was obtained in very
dilute solution. (Winnsinger, Bull. Soc. (2)
49. 452.)

Does not exist. (Kritschenko, Z. anorg. 4.
247.)

Palladium disulphide, PdS₂.

HNO₃ dissolves out part of the S. Easily
sol. in aqua regia without separation of S.
(Schneider.)

Palladium sulphide with M₂S.

See Sulphopalladate, M.

Palladodiamine bromide, Pd(N₂H₅Br)₂.

Easily sol. in H₂O.

— bromopalladite, Pd(N₂H₅Br)₂, PdBr₂.

Properties as the corresponding chloropal-
ladite.

— carbonate.

Sol. in H₂O.

— chloride, Pd(N₂H₅Cl)₂.

Easily sol. in H₂O.

Palladodiamine chloropalladite, $\text{Pd}(\text{N}_2\text{H}_4\text{Cl})_2$,
 PdCl_2 .

"Vauquelin's red salt." Insol. in cold H_2O .
(Fischer.)

Sol. in boiling H_2O with decomp. Sol. in
 HCl or $\text{HNO}_3 + \text{Aq}$.

— **fluoride**.,

Easily sol. in H_2O . (Müller.)

— **fluosilicate**.

Sl. sol. in cold, easily in warm H_2O . Insol.
in alcohol.

— **hydroxide**, $\text{Pd}(\text{N}_2\text{H}_4\text{OH})_2$.

Sol. in H_2O .

— **iodide**, $\text{Pd}(\text{N}_2\text{H}_4\text{I})_2$.

Sol. in H_2O .

— **nitrate**, $\text{Pd}(\text{N}_2\text{H}_4\text{NO}_3)_2$.

Easily sol. in H_2O , HNO_3 , or $\text{NH}_4\text{OH} + \text{Aq}$.
Insol. in alcohol.

— **palladous nitrite**, $\text{Pd}(\text{N}_2\text{H}_4\text{NO}_2)_2$,
 $\text{Pd}(\text{NO}_2)_2$.

Easily sol. in H_2O .

— **sulphate**, $\text{Pd}(\text{N}_2\text{H}_4)_2\text{SO}_4 + \text{H}_2\text{O}$.

Easily sol. in H_2O . Insol. in alcohol.

— **sulphite**, $\text{Pd}(\text{N}_2\text{H}_4)_2\text{SO}_3$.

Sl. sol. in H_2O .

Palladochloronitrous acid.

Potassium palladochloronitrite,
 $\text{Pd}(\text{NO}_2)_2\text{Cl}_2\text{K}_2$.

Sol. in 2 pts. hot, and 3 pts. cold H_2O .
(Vézes, C. R. 115. 111.)

Palladocyanhydric acid.

Ammonium palladocyanide,
 $(\text{NH}_4)_2\text{Pd}(\text{CN})_4$ (?).

Sol. in hot H_2O . (Rössler, Z. ch. 1866. 175.)

Barium —, $\text{BaPd}(\text{CN})_4 + 4\text{H}_2\text{O}$.

Not efflorescent. Sol. in H_2O .

Calcium —, $\text{CaPd}(\text{CN})_4 + 4\text{H}_2\text{O}$.

Sol. in H_2O .

Cupric —, $\text{CuPd}(\text{CN})_4$.

Ppt.

Lead —, $\text{PbPd}(\text{CN})_4$.

Ppt.

Magnesium —, $\text{MgPd}(\text{CN})_4$.

Very sol. in H_2O .

Magnesium — **platinocyanide**, $\text{MgPt}(\text{CN})_4 + 14\text{H}_2\text{O}$.

Extremely sol. in H_2O .

Potassium —, $\text{K}_2\text{Pd}(\text{CN})_4 + 3\text{H}_2\text{O}$.

Efflorescent. Sol. in H_2O .

+ H_2O . Not efflorescent.

Silver —, $\text{Ag}_2\text{Pd}(\text{CN})_4$.

Ppt.

Sodium —, $\text{Na}_2\text{Pd}(\text{CN})_4$.

Not efflorescent. Sol. in H_2O .

+ H_2O .

Palladoiodonitrous acid.

Potassium palladoiodonitrite,
 $\text{Pd}(\text{NO}_2)_2\text{I}_2\text{K}_2 + 3\text{H}_2\text{O}$.

Effloresces in the air.

Decomp. by H_2O and dil. acids.
heim, Z. anorg. 1900, 23. 28.)

Palladonitrous acid.

Potassium palladonitrite, $\text{K}_2\text{Pd}(\text{N}_2\text{H}_4\text{NO}_2)_2$.

Efflorescent; sol. in H_2O . (Lang,
415.)

Silver palladonitrite, $\text{Ag}_2\text{Pd}(\text{NO}_2)_4$.

Easily sol. in hot H_2O . (Lang.)

Sodium —, $\text{Na}_2\text{Pd}(\text{NO}_2)_4$.

(Fischer.)

Palladosamine bromide, $\text{Pd}(\text{NH}_4)_2\text{Br}_2$.

Insol. in cold, sl. sol. in hot H_2O .
sol. in $\text{HC}_2\text{H}_3\text{O}_2$, H_2SO_4 , KOH , NaOH ,
alkali carbonates + Aq . (Müller, A.

— **carbonate**, $\text{Pd}(\text{NH}_4)_2\text{CO}_3$.

Moderately sol. in H_2O .

— **chloride**, $\text{Pd}(\text{NH}_4\text{Cl})_2$.

Insol. in H_2O , but very gradually
by boiling therewith.

Sol. in warm HCl or $\text{HNO}_3 + \text{Aq}$.
cold $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in $\text{KOH} + \text{Aq}$.
out evolution of NH_3 .

+ $2\text{H}_2\text{O}$. Efflorescent. Insol.
(Baubigny, A. Suppl. 4. 253.)

— **cyanide**, $\text{Pd}(\text{NH}_4\text{CN})_2$.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

— **fluoride**.

Known only in solution.

— **hydroxide**, $\text{Pd}(\text{NH}_4\text{OH})_2$.

Easily sol. in H_2O . Slowly decomp.
boiling with H_2O . (Müller, A. 88.)

the iodide, $\text{Pd}(\text{NH}_4\text{I})_2$.
 H_2O . Sol. in boiling HNO_3 with
 I_2 . (Fehling, A. 39. 106.)

ly in solution, which decomp. on

$\text{Pd}(\text{NH}_4\text{NO}_2)_2$.
 y sol. in H_2O . (Lang.)

ous nitrite, $\text{Pd}(\text{NH}_4\text{NO}_2)_2$,
 I_2 .
 .. in cold, easily in hot H_2O .

e, $\text{Pd}(\text{NH}_4)_2\text{SO}_4$.
 y sol. in H_2O . (Müller.)

z, $\text{Pd}(\text{NH}_4)_2\text{SO}_4$.
 in H_2O . (Müller.)

chromium compounds.

ureochromium compounds.
 ureochromium compounds.
 reochromium compounds.
 omium compounds.
 mium compounds.

cobaltic compounds. .

pureocobaltic compounds,
 pureocobaltic compounds.
 pureocobaltic compounds.
 altic compounds,
 obaltic compounds.
 ltic compounds.
 urpureocobaltic compounds.
 altic compounds.

dicobaltic sulphite.
 cobaltic cobaltic sulphite.

iridium compounds.

pentamine, and Iridoaquopent-
 ounds.

rhodium compounds.

pureorhodium compounds.
 pureorhodium compounds.
 reorhodium compounds.
 pureorhodium compounds.
 lium compounds.
 dium compounds.

ic acid, $\text{H}_2\text{S}_2\text{O}_8$.

ly in aqueous solution.
 ition is decomp. by boiling, but
 by addition of acids.

Sp. gr. of aqueous solution of pentathionic
 acid at 22° :

Sp. gr.	1.233	1.320	1.474	1.506
% $\text{H}_2\text{S}_2\text{O}_8$	32.1	41.7	56	59.7

(Kessler, Pogg. 74. 279.)

Does not exist. (Spring, Bull. Acad. roy.
 Belg.)

Existence proven by Smith (Chem. Soc.
 43. 355.)

Barium pentathionate, $\text{BaS}_2\text{O}_8 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O . Aqueous solution is
 precipitated by alcohol.

Contains $3\text{H}_2\text{O}$. (Lewes, C. N. 43. 41.)

Barium pentathionate tetrathionate, BaS_2O_8 ,
 $\text{BaS}_4\text{O}_{12} + 6\text{N}_2\text{O}$.

Easily sol. in H_2O . Not precipitated from
 aqueous solution by two vols. alcohol. (Lud-
 wig, Arch. Pharm. (2) 51. 264.)

Cupric pentathionate, $\text{CuS}_2\text{O}_8 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O . (Debus, Chem. Soc.
 53. 360.)

Lead pentathionate, $\text{PbS}_2\text{O}_8 + 4\text{H}_2\text{O}$.

Ppt.

Potassium pentathionate, $\text{K}_2\text{S}_2\text{O}_8$.

Sol. in H_2O . (Rammelsberg, J. B. 1857.
 136.)

Solution decomposes very quickly when
 neutral, but is more stable in presence of salts
 or acids.

Sol. in about 2 pts. H_2O .

Insol. in alcohol. (Debus, Chem. Soc. 53.
 295.)

+ H_2O . (Shaw, Chem. Soc. 43. 351.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. (Debus, A. 244. 76.)

+ $2\text{H}_2\text{O}$. (Lewes, C. N. 43. 41.)

Perarsenic acid.

Sodium perarsenate, NaAsO_4 .

(Alvarez, C. N. 1906, 94. 270.)

Perboric acid.

Ammonium perborate, NH_4BO_3 .

(Constam and Bennett, Z. anorg. 1900, 25.
 265.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Stable in dry air. 100 pts. H_2O
 at 17.5° dissolve 1.55 g. anhydrous salt.

Decomp. in aqueous solution at ord. temp.

Decomp. by dil. and conc. H_2SO_4 and by
 HCl . (Melikoff, B. 1898, 31. 953.)

+ H_2O . (Bruhat, C. R. 1905, 140. 508.)

Much more sol. in H_2O than the Na salt.
 (Tanatar, Z. phys. Ch. 1898, 26. 133.)

+ $3\text{H}_2\text{O}$. (Melikoff, B. 1898, 31. 954.)

NH_4BO_3 , $\text{NH}_4\text{BO}_3 + \text{H}_2\text{O}$. (Petrenko, C.
 C. 1902, I. 1192.)

Barium perborate, $\text{Ba}(\text{BO}_3)_2 + 7\text{H}_2\text{O}$.

Difficultly sol. in H_2O . (Melikoff, B. 1898, 31. 954.)

Cæsium perborate, $\text{CsBO}_3 + \text{H}_2\text{O}$.

As NH_4^+ salt. (Christensen.)

Calcium perborate.

Sl. sol. in H_2O . Decomp. in water much more rapidly than the Ba salt. (Melikoff, B. 1898, 31. 954.)

Copper perborate.

Very unstable. Insol. in H_2O . (Melikoff, B. 1898, 31. 954.)

Nickel perborate.

Very unstable. Insol. in H_2O . (Melikoff.)

Potassium perborate, $2\text{KBO}_3 + \text{H}_2\text{O}$.

1.25 pts. are sol. in 100 pts. H_2O at 0° ; 2.5 pts., at 15° .

Insol. in alcohol and ether. (Girsewald, B. 1909, 42. 867.)

Potassium perborate hydrogen peroxide, $2\text{KBO}_3, \text{H}_2\text{O}_2$.

0.70 pt. is sol. in 100 pts. H_2O at 15° . (Girsewald, B. 1909, 42. 868.)

Potassium perdisborate, $\text{KB}_2\text{O}_5 + 2\text{H}_2\text{O}$.

Ppt. (Bruhat, C. R. 1905, 140. 508.)

Rubidium perborate, $\text{RbBO}_3 + \text{H}_2\text{O}$.

As Na salt. (Christensen.)

Sodium perborate, $\text{Na}_2\text{B}_4\text{O}_7 + 10\text{H}_2\text{O}$.

100 g. H_2O dissolve 4.2 g. at 11° ; 7.1 g. at 22° ; 13.8 g. at 32° . (Jaubert, C. R. 1904, 134. 796.)

+ $4\text{H}_2\text{O}$. Slowly decomp. in cold solution, rapidly when boiled. (Tanatar, Z. phys. Ch. 1898, 26. 132.)

Sol. in H_2O . 100 g. H_2O dissolve 1.17 g. Aq. solution decomp. on warming. (Melikoff, B. 1898, 31. 679.)

100 g. H_2O dissolve 2.55 g. at 15° ; 2.69 g. at 21° ; 2.85 g. at 26° ; 3.78 g. at 32° . (Jaubert and Lion, Rev. gén. Chim. 1905, (7) 8. 163.)

Uranyl perborate, UBO_4 .

(Bruhat, C. R. 1905, 140. 508.)

Perbromic acid, HBrO_4 .

Known only in aqueous solution, which can be concentrated to a thick liquid on water bath. Not decomp. by HCl , SO_2 , or H_2S . (Kämmerer, J. pr. 85. 452; 90. 190.)

Does not exist. (Muir, C. N. 33. 256; MacIvor, C. N. 33. 35.)

Barium perbromate, $\text{Ba}(\text{BrO}_4)_2$.

Very sl. sol. in boiling H_2O . (Kä J. pr. 90. 190.)

Does not exist. (Wolfram, A. 193.

Potassium perbromate, KBrO_4 .

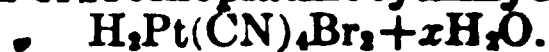
Less sol. in H_2O than KBrO_3 , but r than KClO_4 . (Kämmerer, J. pr. 90.

Does not exist. (Wolfram, A. 193

Silver perbromate, AgBrO_4 .

Sl. sol. in cold, more abundantly in (Kämmerer, J. pr. 90. 190.)

Does not exist. (Wolfram, A. 193

Perbromoplatinocyanhydric acid,

Deliquescent. Easily sol. in H_2O , and ether. (Holst, Bull. Soc. (2) 22

Aluminum perbromoplatinocyanide,

Deliquescent. Very sol. in H_2O .

Ammonium —, $(\text{NH}_4)_2\text{Pt}(\text{CN})_4\text{Br}$

Sol. in H_2O .

Barium —, $\text{BaPt}(\text{CN})_4\text{Br}_2 + 5\text{H}_2\text{O}$

Very sol. in H_2O or alcohol.

Cadmium —, $\text{CdPt}(\text{CN})_4\text{Br}_2 + x\text{H}$

Very sol. in H_2O .

Calcium —, $\text{CaPt}(\text{CN})_4\text{Br}_2 + 7\text{H}_2$

Sol. in H_2O .

Cobaltous —, $\text{CoPt}(\text{CN})_4\text{Br}_2 + 5\text{H}$

Sol. in H_2O . Sl. sol. in alcohol.

Glucinum —, $\text{GlPt}(\text{CN})_4\text{Br}_2$.

Deliquescent. Sol. in H_2O .

Ferrous —.

Very sl. sol. in H_2O .

Lead —, $\text{PbPt}(\text{CN})_4\text{Br}_2 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O .

Lithium —, $\text{Li}_2\text{Pt}(\text{CN})_4\text{Br}_2$.

Deliquescent. Sol. in H_2O .

Magnesium —, $\text{MgPt}(\text{CN})_4\text{Br}_2 +$

Sol. in H_2O .

Nickel —, $\text{NiPt}(\text{CN})_4\text{Br}_2 + x\text{H}_2\text{O}$

Sl. sol. in H_2O . Sol. in NH_4OH .

Potassium —, $\text{K}_2\text{Pt}(\text{CN})_4\text{Br}_2$.

Sol. in H_2O .

+ $2\text{H}_2\text{O}$. Efflorescent.

omoplatinocyanide,
 $\text{Pt}(\text{CN})_4$.
 olati, *Gazz. ch. it.* 1900, **30**, 588.)

γ , $\text{Na}_2\text{Pt}(\text{CN})_4\text{Br}_2$.
 ent. Sol. in H_2O .

δ , $\text{SrPt}(\text{CN})_4\text{Br}_2 + 7\text{H}_2\text{O}$.
 O.

$\text{InPt}(\text{CN})_4\text{Br}_2 + 5\text{H}_2\text{O}$.
 sol. in H_2O .

c acid.

percarbonate, $(\text{NH}_4)_2\text{CO}_4 +$

O with evolution of NH_3 . Insol.
 and ether. (Kasanesky, *C. C.*
 3.)

arbonate, BaCO_4 .

H_2O . (Merck, *C. C.* 1906, II

slowly in the air. Not rapidly
 H_2O . Rapidly decomp. by acids.
n. B. 1908, **41**, 280.)

ercarbonate, K_2CO_4 .

H_2O with decomp. Sl. sol. in al-
 iansen, *Z. Elektrochem.* 1897, **3**.

Sol in H_2O at 0° with only slight
 t is decomp. at ord. temp. Sl.
 ol. (Treadwell, *Ch. Z.* 1901, **25**.

ercarbonate, $\text{Rb}_2\text{CO}_4, 2\text{H}_2\text{O}_2 +$

ic; decomp. by H_2O ; pptd. by

$\text{H}_2\text{O}_2 + 2\text{H}_2\text{O}$. Hygroscopic; de-
 H_2O ; pptd. by alcohol.

$2\frac{1}{2}\text{H}_2\text{O}$. Hygroscopic; decomp.
 d. by alcohol. (Peltner, *B.* 1909,

Very deliquescent. (Constam
Z. Elektrochem. 1897, **3**, 144.)

arbonate, $\text{Na}_2\text{CO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$.

O with gradual decomp. (Tana-
32, 1544.)

rogen percarbonate, $4\text{Na}_2\text{CO}_4$.

erck, *Chem. Soc.* 1908, **94**, (2)

acid, HClO_4 .

with H_2O with a hissing sound
 n of much heat.

Solution in H_2O is very stable.

When dil. $\text{HClO}_4 + \text{Aq}$ is distilled, H_2O and
 HClO_4 distil off until a temp. of 203° is
 reached, when an acid of constant composi-
 tion containing 71.6–72.2% HClO_4 ($=\text{HClO}_4$
 $+2\text{H}_2\text{O}$) is obtained. Forms hydrate HClO_4
 $+ \text{H}_2\text{O}$, which is deliquescent, and dissolves
 in H_2O with evolution of much heat. HClO_4
 is very unstable, $\text{HClO}_4 + \text{H}_2\text{O}$ more stable,
 and $\text{HClO}_4 + 2\text{H}_2\text{O}$ is very stable. (Roscoe,
A. **121**, 346)

Sp. gr. of $\text{HClO}_4 + \text{Aq}$. at $15^\circ/4^\circ$.

Sp. gr.	% HClO_4	Sp. gr.	% HClO_4	Sp. gr.	% HClO_4
1.005	1.00	1.235	33.29	1.465	54.50
1.010	1.90	1.240	33.85	1.470	54.89
1.015	2.77	1.245	34.40	1.475	55.18
1.020	3.61	1.250	34.95	1.480	55.56
1.025	4.43	1.255	35.49	1.485	55.95
1.030	5.25	1.260	36.03	1.490	56.32
1.035	6.07	1.265	36.56	1.495	56.69
1.040	6.88	1.270	37.08	1.500	57.06
1.045	7.68	1.275	37.60	1.505	57.44
1.050	8.48	1.280	38.10	1.510	57.81
1.055	9.28	1.285	38.60	1.515	58.17
1.060	10.06	1.290	39.10	1.520	58.54
1.065	10.83	1.295	39.60	1.525	58.91
1.070	11.58	1.300	40.10	1.530	59.28
1.075	12.33	1.305	40.59	1.535	59.66
1.080	13.08	1.310	41.08	1.540	60.04
1.085	13.83	1.315	41.56	1.545	60.41
1.090	14.56	1.320	42.03	1.550	60.78
1.095	15.28	1.325	42.49	1.555	61.15
1.100	16.00	1.330	42.97	1.560	61.52
1.105	16.72	1.335	43.43	1.565	61.89
1.110	17.45	1.340	43.89	1.570	62.26
1.115	18.16	1.345	44.35	1.575	62.63
1.120	18.88	1.350	44.81	1.580	63.00
1.125	19.57	1.355	45.26	1.585	63.37
1.130	20.26	1.360	45.71	1.590	63.74
1.135	20.95	1.365	46.16	1.595	64.12
1.140	21.64	1.370	46.61	1.600	64.50
1.145	22.32	1.375	47.05	1.605	64.88
1.150	22.99	1.380	47.49	1.610	65.26
1.155	23.65	1.385	47.93	1.615	65.63
1.160	24.30	1.390	48.37	1.620	66.01
1.165	24.94	1.395	48.80	1.625	66.39
1.170	25.57	1.400	49.23	1.630	66.76
1.175	26.20	1.405	49.68	1.635	67.13
1.180	26.82	1.410	50.10	1.640	67.51
1.185	27.44	1.415	50.51	1.645	67.89
1.190	28.05	1.420	50.91	1.650	68.26
1.195	28.66	1.425	51.31	1.655	68.64
1.200	29.26	1.430	51.71	1.660	69.02
1.205	29.86	1.435	52.11	1.655	69.40
1.210	30.45	1.440	52.51	1.670	69.77
1.215	31.04	1.445	52.91	1.675	70.15
1.220	31.61	1.450	53.31		
1.225	32.18	1.455	53.71		
1.230	32.74	1.460	54.11		

(Emster, *Z. anorg.* 1907, **52**, 278.)

Sp. gr. of HClO ₄ +Aq.			
% HClO ₄	Sp. gr. at 15°/4°	Sp. gr. at 30°/4°	Sp. gr. at 50°/4°
11.14	1.0670	1.0507
35.63	1.2569	1.2451	1.2292
55.63	1.4807	1.4637	1.4421
69.81	1.6708	1.6284

(Emster, Z. anorg. 1907, 52. 279.)

Sp. gr. of HClO ₄ +Aq.				
Sp. gr.				% by wt. HClO ₄ in the liquid
corrected		uncorrected		
20°	50°	20°	50°	
1.7676	1.7098	1.7716	1.7312	100
1.7817	1.7259	1.7858	1.7475	98.62
1.8059	1.7531	1.8100	1.7751	94.67
...	1.7690	...	1.7912	90.80
...	1.7756	...	1.7979	84.81
...	1.7619	...	1.7840	81.07
1.7386	1.7023	1.7425	1.7237	75.59
1.6471	1.6110	1.6508	1.6311	68.42
1.5353	1.5007	1.5386	1.5194	60.38
1.4078	1.3779	1.4108	1.3949	50.51
1.2901	1.2649	1.2927	1.2804	39.73
1.1778	1.1574	1.1800	1.1715	27.07

(v. Wyk, Z. anorg. 1905, 48. 45.)

Bpt. of HClO₄+Aq. at atmospheric pressure.

% by wt. HClO ₄ in the liquid	% by wt. HClO ₄ in the vapor	Initial bpt.
72.4	72.4	203°
70.06	40.11	198.7
65.2	6.06	181.2
61.2	0.9	162.3
56.65	...	148.0
50.67	...	132.4
38.90	...	114.8
24.23	...	105.8
0.0	0.0	100

(v. Wyk, Z. anorg. 1905, 48. 33.)

Bpt. of HClO₄+Aq. at 18 mm. pressure.

% by wt. HClO ₄ in the liquid	Bpt.
100	16.0°
94.8	24.8
92.0	35
84.8	70
79.8	92
70.5	107

(v. Wyk, Z. anorg. 1905, 48. 36.)

+H₂O. Deliquescent. (Roscoe, A. 121. 346.)
+2H₂O. HClO₄+2H₂O has 1.65 sp. gr.

and boils at 200° (Serullas); has 1.1 sp. gr. and boils at 200° (Nativelle, J. 405).

Sol. in alcohol with decomp.; offensive.
+2½H₂O. (v. Wyck.)
+3H₂O. (v. Wyck.)
+3½H₂O. (v. Wyck.)

Perchlorates.

All perchlorates are sol. in H₂O, RbClO₄, and CsClO₄, somewhat dif. They are all deliquescent, and sol in excepting NH₄ClO₄, KClO₄, Pb(ClO₄)₂, Hg₂(ClO₄)₂. (Serullas, A. ch. (2) 48.

Aluminum perchlorate, Al(ClO₄)₃+6H₂O.
Very deliquescent. (Weinland, Z. 1913, 84. 370.)

Aluminum sodium perchlorate, AlNaClO₄+12H₂O.
Sl. hygroscopic. (Weinland, Z. and 84. 370.)
Sol. in acetone. (Naumann, B. 4328.)

Ammonium perchlorate, NH₄ClO₄.
Permanent. Sol. in 5 pts. H₂O; sol. in alcohol. (Mitscherlich, Pogg.

Solubility of NH₄ClO₄ in H₂O

t°	G. per l. solution	
0	115.63	1
20	208.45	1
40	305.77	1
60	390.50	1
80	481.86	1
100	570.06	1
107	591.15	1

(Carlson, Festsck. Stockholm. 191

100 g. H₂O dissolve 18.5 g. (Hofmann, Höbald and Quocq, A. 304.)

100 g. sat. solution in H₂O cont (17.35?) g. NH₄ClO₄ at 14.2°. (Cumming, Chem. Soc. 1915, 107. 36

Insol. in conc. HClO₄+Aq.
100 g. sat. solution in 98.8% eth contain 1.96 g. NH₄ClO₄ at 25.2 and Cumming.)
Sol. in acetone. Eidmann, C. C. 1014.)

Barium perchlorate, Ba(ClO₄)₂+4H₂O.
Deliquescent. Easily sol. in alcohol.

Solubility of $\text{Ba}(\text{ClO}_4)_2 + 3\text{H}_2\text{O}$ in H_2O at t° .

	G. per 100 g. H_2O	Sp. gr.
	206	1.782
	289	1.912
	358	2.009
	432	2.070
	497	2.114
	564	2.155
	645	2.195
	758	2.230

n, Festsck. Stockholm, 1911. 262.)

perchlorate, $(\text{BiO})\text{ClO}_4$.

$1\text{H}_2\text{O}$. Easily sol. in HCl or HNO_3 , easily in $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Muir, C. N.

perchlorate, $\text{Cd}(\text{ClO}_4)_2$.

deliquescent. Sol. in H_2O and alcohol. (Serullas, A. ch. 46. 305.)

l. (Salvadori, C. C. 1912, II. 414.)

l. (S.)

perchlorate ammonia, $\text{Cd}(\text{ClO}_4)_2$,

4NH_3 . (Salvadori, C. C. 1912,

perchlorate, CsClO_4 .

sol. in H_2O . (Retgers, Z. phys. Ch.

ity in H_2O . 100 g. H_2O dissolve at:

1° 33.7° 42° 50°
19 2.99 4.09 5.47 g. CsClO_4 ,

79° 84° 99°
79 16.51 28.57 g. CsClO_4 .

i, Acc. Sc. Med. Ferrara, 1911, 85. 150.)

Solubility in H_2O at t° .

	G. per 100 g. H_2O	Sp. gr.
	0.97	1.007
	2.05	1.010
	17.05	1.084

n, Festsck. Stockholm, 1911. 262.)

perchlorate, $\text{Ca}(\text{ClO}_4)_2$.

deliquescent. Very sol. in H_2O and alcohol. (Serullas, A. ch. 46. 304.)

perchlorate, $\text{Ce}(\text{ClO}_4)_3 + 8\text{H}_2\text{O}$.

deliquescent. (Jolin.)

perchlorate, $\text{Cr}(\text{ClO}_4)_3 + 6\text{H}_2\text{O}$.

hygroscopic. (Weinland, Z. anorg. 371.)

l. Can be cryst. from H_2O . (Wein-

Cobaltous perchlorate, $\text{Co}(\text{ClO}_4)_2 + 9\text{H}_2\text{O}$.Solubility in H_2O at t° .

t°	G. anhydrous salt in 100 ccm.	Sp. gr. of sat. solution at $t^\circ/4^\circ$
-30.7	83.14	...
-21.3	90.57	...
0	100.13	1.5639
+ 7.5	101.92	1.5658
18	103.80	1.5670
26	113.45	1.5811
45	115.10	1.5878

(Golblum and Terlikowsky, Bull. Soc. 1912, (4) 11. 146.)

+ $6\text{H}_2\text{O}$. (Salvadori, Gazz. ch. it. 1912, 42. (1) 458.)

Cobalt perchlorate ammonia, $\text{Co}(\text{ClO}_4)_2$, 6NH_3 .

$\text{Co}(\text{ClO}_4)_2$, 5NH_3 .

$\text{Co}(\text{ClO}_4)_2$, 4NH_3 , and $+2\text{H}_2\text{O}$.

$\text{Co}(\text{ClO}_4)_2$, 3NH_3 , and $+3\text{H}_2\text{O}$.

$\text{Co}(\text{ClO}_4)_2$, $3\text{NH}_3 + 2\text{H}_2\text{O}$.

(Salvadori, Gazz. ch. it. 1912, 42. (1) 458.)

Cupric perchlorate, basic, $\text{Cu}(\text{ClO}_4)_2$, $\text{Cu}(\text{OH})_2$.

Ppt. (Salvadori, C. C. 1912, II. 414.)

Cupric perchlorate, $\text{Cu}(\text{ClO}_4)_2$.

Deliquescent. Sol. in H_2O and alcohol. (Serullas, A. ch. 46. 306.)

+ $4\text{H}_2\text{O}$. (Salvadori, C. C. 1912, II. 414.)

Cupric perchlorate ammonia, $\text{Cu}(\text{ClO}_4)_2$, $4\text{NH}_3 + 2\text{H}_2\text{O}$.

Not deliquescent. Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Roscoe, A. 121. 346.)

$\text{Cu}(\text{ClO}_4)_2$, $\text{NH}_3 + \text{H}_2\text{O}$.

$\text{Cu}(\text{ClO}_4)_2$, $2\text{CuO} + 2\text{H}_2\text{O}$, NH_3 .

$\text{Cu}(\text{ClO}_4)_2$, $2\text{CuO} + 2\text{H}_2\text{O}$, 2NH_3 .

$\text{Cu}(\text{ClO}_4)_2$, $\text{Cu}(\text{OH})_2 + 2\text{H}_2\text{O}$, 6NH_3 .

$\text{Cu}(\text{ClO}_4)_2$, $\text{Cu}(\text{OH})_2 + 2\text{H}_2\text{O}$, 4NH_3 .

(Salvadori, C. C. 1912, II. 414.)

Didymium perchlorate, $\text{Di}(\text{ClO}_4)_2 + 9\text{H}_2\text{O}$.

Very deliquescent. Very sol. in H_2O and alcohol. (Cleve.)

Erbium perchlorate, $\text{Er}(\text{ClO}_4)_3 + 8\text{H}_2\text{O}$.

Very deliquescent.

Glucinum perchlorate, $\text{Gl}(\text{ClO}_4)_2 + 4\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O . (Atterberg.)

Hydrazine perchlorate, $(\text{N}_2\text{H}_4)(\text{HClO}_4)_2 + 3\text{H}_2\text{O}$.

1 l. of sat. solution in H_2O contains 417.2 g. at 18° , sp. gr. = 1.264; 669 g. at 35° , sp. gr. = 1.391. (Carlson, Festsck. Stockholm, 1911. 262.)

Indium perchlorate, $\text{In}(\text{ClO}_4)_3 + 8\text{H}_2\text{O}$.

Deliquescent. H_2O solution decomp. at 40° with separation of basic salt. Sol. in H_2O and easily forms sat. solutions. Sol. in abs. alcohol, but much less sol. in ether. (Mathers, J. Am. Chem. Soc. 1908, **30**. 212.)

Iodine perchlorate, $\text{I}(\text{ClO}_4)_3 + 2\text{H}_2\text{O}$.

Decomp. by H_2O . Indifferent toward organic solvents. (Fichter, Z. anorg. 1915, **91**. 135.)

Iron (ferrous) perchlorate, $\text{Fe}(\text{ClO}_4)_2$.

Tolerably permanent; sol. in H_2O . (Serullas, A. ch. **46**. 335.)

Iron (ferric) perchlorate, $\text{Fe}(\text{ClO}_4)_3$.

Sol. in H_2O . (Serullas.)

Iron (ferric) sodium perchlorate,
 $[\text{Fe}(\text{ClO}_4)_4]\text{Na} + 6\text{H}_2\text{O}$.

Hydroscopic. Can be cryst. from H_2O . (Weinland, Z. anorg. 1913, **84**. 366.)

Lanthanum perchlorate, $\text{La}(\text{ClO}_4)_3 + 9\text{H}_2\text{O}$.

Extremely deliquescent. Sol. in H_2O and absolute alcohol. (Cleve.)

Lead perchlorate, basic, 2PbO , $\text{Cl}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Decomp. by H_2O into an insol. more basic salt, and sol. $\text{Pb}(\text{ClO}_4)_2$. (Marignac.)

Lead perchlorate, $\text{Pb}(\text{ClO}_4)_2 + 3\text{H}_2\text{O}$.

Permanent; extremely easily sol. in H_2O . (Roscoe, A. **121**. 356.)

Sol. in about 1 pt. H_2O . (Serullas.)

Lithium perchlorate, LiClO_4 .

Deliquescent. Sol. in H_2O and alcohol. (Serullas.)

+ $3\text{H}_2\text{O}$. (Wyruboff, Zeit. Kryst. **10**. 626.)

Magnesium perchlorate, $\text{Mg}(\text{ClO}_4)_2$.

Deliquescent, and sol. in H_2O and alcohol. (Serullas.)

+ $6\text{H}_2\text{O}$. (Weinland, Z. anorg. 1913, **84**. 372.)

Manganous perchlorate, $\text{Mn}(\text{ClO}_4)_2$.

Very deliquescent. Sol. in H_2O and alcohol. (Serullas, A. ch. **46**. 335.)

+ $6\text{H}_2\text{O}$. Sol. in 0.342 pts. H_2O . (Salvadori, C. C. **1912**, II. 414.)

Manganous perchlorate ammonia, $\text{Mn}(\text{ClO}_4)_2$,
 $5\text{NH}_3 + \text{H}_2\text{O}$.

Sol. in HCl ; insol. in HNO_3 . (Salvadori, C. C. **1912**, II. 414.)

Mercurous perchlorate, $(\text{HgClO}_4)_2 + 4\text{H}_2\text{O}$.

Very sol. in H_2O . Gradually decomp. by

H_2O . Decomp. by alcohol. (Chika Chem. Soc. 1895, **67**. 1016.)

+ $6\text{H}_2\text{O}$. Very deliquescent. (Roscoe, **121**. 356.)

Permanent. (Serullas.)

Mercuric perchlorate, basic, HgO , $2\text{Hg}(\text{ClO}_4)_2$.

Anhydrous. Ppt. Insol. in either H_2O or HNO_3 . Decomp. and dissolved by a mixture of the two. (Chikashigé, Chem. Soc. **1895**, **67**. 824.)

+ $12\text{H}_2\text{O}$. Very sol. in H_2O . (Chikashigé, **1895**, **67**. 824.)

α -salt. Decomp. by H_2O . Sol. in H_2O . (Chikashigé, Chem. Soc. **1895**, **67**. 1016.)

β -salt. Insol. in H_2O ; insol. in HNO_3 . (Chikashigé, Chem. Soc. **1895**, **67**. 825.)

Mercuric perchlorate, $\text{Hg}(\text{ClO}_4)_2$.

Very deliquescent. Sol. in H_2O ; with decomp. in alcohol. (Serullas, **46**. 243.)

+ $6\text{H}_2\text{O}$. Very hygroscopic. Very sol. in H_2O . Slowly decomp. by H_2O , more by alcohol. (Chikashigé, Chem. Soc. **1895**, **67**. 1014.)

Mercuric perchlorate bromide, HgClO_4 , HgBrClO_4 .

Decomp. by H_2O . (Borelli, Gazz. **1908**, **38**. (2) 421.)

Mercuric perchlorate cyanide, $\text{Hg}(\text{ClO}_4)_2$, $\text{Hg}(\text{CN})_2$.

Very sol. in H_2O . Sol. in alcohol. (Borelli.)

Mercuric perchlorate iodide, $\text{Hg}(\text{ClO}_4)_2$, HgI_2 .

Deliquescent. Decomp. by H_2O . Sol. in much alcohol. Decomp. by HNO_3 . Completely sol. in KI or $\text{KCN} + \text{Aq}$. (Borelli.)

Mercuric perchlorate sulphocyanide, $\text{Hg}(\text{ClO}_4)_2$, $\text{Hg}(\text{SCN})_2$.

Insol. in H_2O and conc. acids. Sol. in regia. (Borelli.)

+ $6\text{H}_2\text{O}$. (Salvadori, C. C. **1912**, II. 414.)

Nickel perchlorate, $\text{Ni}(\text{ClO}_4)_2$.

Deliquescent; easily sol. in alcohol. Sol. in H_2O . (Groth, Pogg. **183**. 226.)

Solubility in H_2O at t° .

t°	G. anhydrous salt in 100 ccm.	Sp. gr. sat. sol.
—30.7	89.98	...
—21.3	92.48	...
0	104.55	1.57
+7.5	106.76	1.57
18	110.05	1.57
26	112.15	1.58
45	118.60	1.58

(Golblum and Terlikowsky, Bull. Soc. **1912**, (4) **11**. 147.)

K_2O . (Golblum and Terlikowsky.)
 K_2O . (Salvadori, C. C. 1912, II. 414.)
 K_2O . (Golblum and Terlikowsky.)

perchlorate, ammonia, $\text{Ni}(\text{ClO}_4)_2 \cdot \text{H}_2\text{O}$.
 (Salvadori.)

l perchlorate, $\text{NO.O.ClO}_4 + \text{H}_2\text{O}$.
 sl. hygroscopic; decomp. by H_2O .
 nn, B. 1909, 42. 2032.)

m perchlorate, $\text{Pt}_4\text{ClO}_4 + 15\text{H}_2\text{O}$.
 in H_2O . (Prost, Bull. Soc. (2) 46.)

um perchlorate, KClO_4 .

57.9 pts. H_2O at 21.3° (Longuine, A. 121.
 5 pts. H_2O at 15° (Serullas, A. ch. (2) 46. 297);
 H_2O at 10° ; in 55 pts. H_2O at 100° (Hutstein,
 1. 331.)

ility in H_2O .

KClO_4 dissolves in 142.9 pts. H_2O at
 solution has sp. gr. = 1.0005; in 52.5
 O at 25° , and solution has sp. gr. =
 in 15.5 pts. H_2O at 50° , and solution
 gr. = 1.0181; in 5.04 pts. H_2O at 100° ,
 ution has sp. gr. = 1.0660. (Muir, C.
 15.)

I_2O dissolves 78.07 millimols. KClO_4
 120.4 millimols. at 20° ; 179.9 milli-
 t 30° . (Noyes and Sammet, Z. phys.
 3, 43. 538.)

H_2O dissolves 0.1475 mol. KClO_4 at
 lothmund, Z. phys. Ch. 1909, 69. 539.)

Solubility in H_2O at t° .

G. KClO_4 in 100 g. H_2O	t°	G. KClO_4 in 100 g. H_2O
0.70	50	6.45
1.14	70	12.3
1.54	99	22.2
1.90

ri, Acc. Sci. Med. Ferrara, 1911, 85.
 150.)

	G. per 100 g. H_2O	Sp. gr.
0	0.79	1.007
0	1.80	1.011
0	4.81	1.022
0	8.71	1.033
0	14.78	1.053
0	20.98	1.067

on, Festsck. Stockholm, 1911. 262.)

H_2O dissolves 0.1481 equivalents
 at 25° . (Noyes and Boggs, J. Am.
 Soc. 1911, 33. 1652.)

. of sat. solution of KClO_4 in H_2O con-

tains 2.085 g. KClO_4 at 25.2° . (Thin and Cum-
 ming, Chem. Soc. 1915, 107. 361.)

KClO_4 is sol. in 22.6 pts. H_2O at ord. temp.,
 and 4.00 pts. at 100° ; in 29.6 pts. $\text{NH}_4\text{OH} +$
 Aq (conc.) at ord. temp.; in 30.4 pts. NH_4OH
 $+ \text{Aq}$ (1 vol. conc. + 3 vols. H_2O) at ord.
 temp.; in 22.4 pts. $\text{HNO}_3 + \text{Aq}$ (1 vol. conc. +
 5 vols. H_2O) at ord. temp., and 5.00 pts. at
 100° ; in 30.4 pts. $\text{HCl} + \text{Aq}$. (1 vol. conc. +
 4 vols. H_2O) at ord. temp.; 45.2 pts. $\text{HC}_2\text{H}_3\text{O}_2$
 $+ \text{Aq}$ (1 vol. commercial acid + 1 vol. H_2O)
 at ord. temp.; in 24.4 pts. $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$.
 (dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{dil. NH}_4\text{OH} + \text{Aq}$) at ord.
 temp., and 6.00 pts. at 100° ; in 25.6 pts.
 $\text{NH}_4\text{Cl} + \text{Aq}$ (1 pt. $\text{NH}_4\text{Cl} + 10$ pts. H_2O) at
 ord. temp., and 6.00 pts. at 100° ; in 16.0 pts.
 $\text{NH}_4\text{NO}_3 + \text{Aq}$ (1 pt. $\text{NH}_4\text{NO}_3 + 10$ pts. H_2O)
 at ord. temp., and 4.00 pts. at 100° ; in 25.6
 pts. $\text{NaC}_2\text{H}_3\text{O}_2 + \text{Aq}$ (conc. $\text{HC}_2\text{H}_3\text{O}_2 +$
 $\text{Na}_2\text{CO}_3 + 4$ vols. H_2O) at ord. temp., and
 7.00 pts. at 100° ; in 29.2 pts. $\text{Cu}(\text{C}_2\text{H}_3\text{O}_2)_2 +$
 Aq (Stolba, Z. anal. 2. 390) at ord. temp.,
 and 7.00 pts. at 100° ; in 27.2 pts. cane sugar
 (1 pt. + 10 pts. H_2O) at ord. temp.; in 36.8
 pts. grape sugar (1 pt. + 10 pts. H_2O) at ord.
 temp. (Approximate.) (Pearson, Zeit. Chem.
 1869. 662.)

Solubility of KClO_4 in HClO_4 at 25.2° .

Normality of HClO_4	% KClO_4
0.01	1.999
0.10	1.485
1.00	0.527

(Thin and Cumming, Chem. Soc. 1915, 107.
 361.)

Solubility in $\text{KCl} + \text{Aq}$ at 25° .

Concentration of KCl Equivalents per litre	Solubility of KClO_4 Equivalents per litre
0.04973	0.1282
0.09933	0.1123

(Noyes and Boggs, J. Am. Chem. Soc. 1911,
 33. 1652.)

Solubility in $\text{K}_2\text{SO}_4 + \text{Aq}$ at 25° .

Concentration of K_2SO_4 Equivalents per litre	Solubility of KClO_4 Equivalents per litre
0.04970	0.1315
0.09922	0.1181

(Noyes and Boggs.)

Very sl. sol. in abs. alcohol, and insol. if
 alcohol contains trace of an acetate. (Ros-
 coe.) Insol. in alcohol of 0.835 sp. gr. (Schlös-
 ing, C. R. 73. 1269.)

Sol. in 6400 pts. 97.2% alcohol; in 5000
 pts. 95.8% alcohol; in 2500-3000 pts. 90%
 alcohol; in 25,000 pts. alcohol-ether (2 pts.
 97% alcohol : 1 pt. ether). Practically insol.

in an alcoholic solution of HClO_4 . (Wenze' Z. angew. Ch. 1891. 691.)

Solubility of KClO_4 in ethyl alcohol+Aq at 25.2° .

Vol. % alcohol	G. KClO_4 sol. in 100 g. alcohol
51.2	0.754
93.5	0.051
98.8	0.019

(Thin and Cumming, Chem. Soc. 1915, 107. 361.)

Solubility in organic compds.+Aq. at 25° .

Solvent	Mol. KClO_4 sol. in 1 litre
0.5-N methyl alcohol	0.1402
" ethyl alcohol	0.1356
" propyl alcohol	0.1343
" tert. amyl alcohol	0.1279
" acetone	0.1451
" ether	0.1336
" glycol	0.1416
" glycerine	0.1404
" urea	0.1510
" ammonia	0.1474
" diethylamine	0.1342
" pyridine	0.1410
" urethane	0.1400
" formamide	0.1539
" acetamide	0.1447
" acetic acid	0.1462
" phenol	0.1362
" methylal	0.1400
" methyl acetate	0.1429

(Rothmund, Z. phys. Ch. 1909, 69. 539.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Potassium rubidium perchlorate, $\text{KRb}_2(\text{ClO}_4)_3$.

15.5 g. are contained in 1 l. solution sat. at 20° ; sp. gr. = 1.013. (Carlson.)

Rubidium perchlorate, RbClO_4 .

Sol. in 92.1 pts. H_2O at 21.3° . (Longuinine, A. 121. 123.)

1 pt. sol. in 92.1 pts. H_2O at 21° as compared with 1 pt. KClO_4 sol. in 57.9 pts. H_2O at 21° . (Erdmann, Arch. Pharm. 1894, 232. 23.)

Solubility in H_2O at t° .			
t°	G. RbClO_4 in 100 g. H_2O	t°	G. RbClO_4 in 100 g. H_2O
0	2.46	42.2	14.94
8	3.50	50	19.40
19.8	6.28	77	41.65
30	9.53	99	76.5

(Calzolari, Acc. Sci. Med. Ferrara, 1911, 85. 150.)

Solubility in H_2O at t° .

t°	G. in 100 g. H_2O	Sp. gr.
0	1.10	1.0
20	1.56	1.0
40	3.26	1.0
60	6.27	1.0
80	11.04	1.0
100	15.75	1.0

(Carlson, Festsk. Stockholm, 1911.

Scandium perchlorate.

(Crookes, Roy. Soc. Proc. 1908, 80.

Silver perchlorate, AgClO_4 .

Deliquescent. Sol. in H_2O and (Serullas, A. ch. 46. 307.)

Sodium perchlorate, NaClO_4 .

Deliquescent, and very sol. in alcohol. (Serullas.)

Not deliquescent. (Potilitsin, J. 1889, 1. 258.)

Solubility in H_2O at t° .

t°	G. in 1 l. of solution	Sp. gr.
15	1076	1.
50	1234	1.
143	1414	1.

(Carlson, Festsk. Stockholm, 1911

+ H_2O . Not deliquescent. (Pot

Strontium perchlorate, $\text{Sr}(\text{ClO}_4)_2$.

Very deliquescent. Sol. in H_2O cohol. (Serullas, A. ch. 46. 304.)

Terbium perchlorate.

Very sol. in H_2O and in alcohol. C. N. 1905, 92. 3.)

Thallous perchlorate, TlClO_4 .

1 pt. salt dissolves in 10 pts. H_2O and 0.6 pt. at 100° . (Roscoe, Chem 4. 504.)

Solubility in H_2O at t° .

t°	G. per 100 g. H_2O	Sp. gr.
0	6.00	1
10	8.04	1
30	19.72	1
50	39.62	1
70	65.32	1
80	81.49	1

(Carlson, Festsk. Stockholm, 1911

Sl. sol. in alcohol. (Roscoe.)

Thallium perchlorate, $\text{Tl}(\text{ClO}_4)_3 + 6\text{H}_2\text{O}$.
 Very hygroscopic, sol. in H_2O . Decomp. in moist air. (Gewecke, Z. anorg. 1912, 75. 1.)

Uranium perchlorate, $(\text{UO}_2)(\text{ClO}_4)_2 + 4\text{H}_2\text{O}$.
 (Salvadori, Ch. Z. 1912, 36. 513.)
 $+ 6\text{H}_2\text{O}$. (Salvadori.)

Yttrium perchlorate, $\text{Y}(\text{ClO}_4)_3 + 8\text{H}_2\text{O}$.
 Very deliquescent. Sol. in H_2O and alcohol. (Cleve.)

Zinc perchlorate, $\text{Zn}(\text{ClO}_4)_2$.
 Deliquescent. Sol. in H_2O and alcohol. (Serullas, A. ch. 48. 302.)
 $+ 4\text{H}_2\text{O}$, and $6\text{H}_2\text{O}$. (Salvadori, C. C. 1912, II. 414.)

Zinc perchlorate, ammonia, $\text{Zn}(\text{ClO}_4)_2, 4\text{NH}_3$.
 Ppt. (Salvadori, C. C. 1912, II. 414.)
 $\text{Zn}(\text{ClO}_4)_2, 6\text{NH}_3$. (Ephraim, B. 1915, 48. 43.)

Perchromic acid.

Sol. in ethyl acetate and valerate; in amyl chloride, formate, acetate, butyrate, and valerate. (All give blue solutions.)

Insol. in CS_2 , C_6H_6 , CHCl_3 , CCl_4 , $\text{C}_6\text{H}_5\text{NH}_2$, $\text{C}_6\text{H}_5\text{NO}_2$ and toluene. (Grosvenor, J. Am. Chem. Soc. 1895, 17. 41-43.)

$\text{H}_2\text{CrO}_5 + 2\text{H}_2\text{O}$. Decomp. above -30° . (Riesenfeld, B. 1914, 47. 552.)

Ammonium perchromate, $(\text{NH}_4)_2\text{CrO}_5$.

Very unstable. Sl. sol. in cold H_2O . Decomp. by conc. H_2SO_4 . Insol. in pure alcohol and pure ether. Decomp. by boiling alcohol containing more than 50% H_2O . (Wohlers, B. 1905, 38. 1888.)

$\text{CrO}_4, 3\text{NH}_3$. Sol. in 10% $\text{NH}_4\text{OH} + \text{Aq}$; sol. in H_2O with decomp., insol. in other solvents. (Wiede, B. 1897, 30. 2181.)

$\text{NH}_4\text{CrO}_5, \text{H}_2\text{O}_2$. Decomp. in the air. Sol. in ice cold H_2O , decomp. when warmed. Insol. in alcohol, ether, ligroin and CHCl_3 . (Wiede, B. 1898, 31. 518.)

Ammonium hydrogen perchromate, $\text{CrO}_2(\text{O.O.NH}_4)(\text{O.OH})$.

Sol. in H_2O with decomp. Difficultly sol. in cold abs. alcohol. (Hofmann, B. 1904, 37. 3406.)

Barium perchromate, BaCr_2O_8 .

(Byers and Reid, Am. Ch. J. 1904, 32. 513.)

Calcium perchromate, CaCr_2O_8 .

Very sol. in H_2O . (Mylius, B. 1900, 33. 3689; Byers and Reid, Am. Ch. J. 1904, 32. 513.)

Lithium perchromate, $\text{Li}_2\text{Cr}_2\text{O}_8$.

(Byers and Reid, Am. Ch. J. 1904, 32. 511.)

Magnesium perchromate, MgCr_2O_8 .

(Byers and Reid.)

Potassium perchromate, $\text{K}_2\text{Cr}_2\text{O}_8$.

Sl. sol. in cold H_2O . Decomp. by conc. H_2SO_4 . Insol. in pure alcohol and pure ether. Decomp. by boiling alcohol containing more than 50% H_2O . (Wohlers, B. 1905, 38. 1888.)

$+ x\text{H}_2\text{O}$. Sol. in H_2O at 0° without decomp. (Riesenfeld and Kutsch, B. 1908, 41. 3948.)

$\text{K}_2\text{Cr}_2\text{O}_8$. Sol. in H_2O . Decomp. in the air. (Byers and Reid, Am. Ch. J. 1904, 32. 505.)

$\text{KCrO}_5, \text{H}_2\text{O}_2$ or KH_2CrO_7 . Sol. in ice cold H_2O , decomp. when warmed; explosive. (Wiede, B. 1898, 31. 520.)

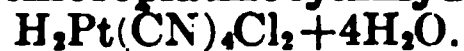
Sodium perchromate, $\text{Na}_2\text{Cr}_2\text{O}_8$.

Sl. sol. in cold H_2O . Decomp. by conc. H_2SO_4 . Insol. in pure alcohol and pure ether. Decomp. by boiling with alc. containing more than 50% H_2O . (Wohlers, B. 1905, 38. 1888.)

$\text{Na}_2\text{Cr}_2\text{O}_8 + 28\text{H}_2\text{O}$. Efflorescent. Sl. sol. in cold, easily in hot H_2O , with decomp. Not decomp. by $\text{NaOH} + \text{Aq}$. (Häussermann, J. pr. (2) 48. 70.)

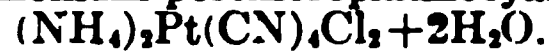
$\text{Na}_2\text{Cr}_2\text{O}_8$. (Byers and Reid, Am. Ch. J. 1904, 32. 511.)

Perchloroplatinocyanhydric acid,



Very sol. in H_2O and alcohol.

Ammonium perchloroplatinocyanide,



Sol. in H_2O .

Barium —, $\text{BaPt}(\text{CN})_4\text{Cl}_2 + 5\text{H}_2\text{O}$.

Very sol. in H_2O .

Calcium —, $\text{CaPt}(\text{CN})_4\text{Cl}_2$.

Sol. in H_2O .

Magnesium —, $\text{MgPt}(\text{CN})_4\text{Cl}_2 + x\text{H}_2\text{O}$.

Sol. in H_2O .

Manganous —, $\text{MnPt}(\text{CN})_4\text{Cl}_2 + 5\text{H}_2\text{O}$.

Sol. in H_2O and alcohol.

Potassium —, $\text{K}_2\text{Pt}(\text{CN})_4\text{Cl}_2 + 2\text{H}_2\text{O}$.

Very efflorescent, and sol. in H_2O and alcohol.

Percolumbic acid, $\text{HCbO}_4 + n\text{H}_2\text{O}$.

Insol. in H_2O . Sol. with decomp. in warm H_2SO_4 . (Melikoff, Z. anorg. 1899, 20. 341.)

Cæsium percolumbate, Cs_2CbO_8 .

Ppt. (E. F. Smith, J. Am. Chem. Soc. 1908, 30. 1658.)

Cæsium magnesium percolumbate,
 $\text{MgCsCbO}_8 + 8\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (E. F. Smith.)

Calcium potassium percolumbate,
 $\text{CaKCbO}_8 + 4\text{H}_2\text{O}$.

Difficultly sol. in H_2O . (E. F. Smith.)

Calcium sodium percolumbate, $\text{CaNaCbO}_8 + 4\text{H}_2\text{O}$.

Difficultly sol. in H_2O . (E. F. Smith.)

Magnesium potassium percolumbate,
 $\text{MgKCbO}_8 + 7\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (E. F. Smith.)

Magnesium rubidium percolumbate,
 $\text{MgRbCbO}_8 + 7\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (E. F. Smith.)

Magnesium sodium percolumbate,
 $\text{MgNaCbO}_8 + 8\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (E. F. Smith.)

Potassium percolumbate, K_2CbO_8 .

Sol. in H_2O . Ppt. from aq. solution by alcohol. (E. F. Smith.)

$\text{K}_4\text{Cb}_2\text{O}_{11} + 3\text{H}_2\text{O}$. Sol. with decomp. in H_2O . Ppt. by alcohol. Sol. in $\text{KOH} + \text{H}_2\text{O}$, + Aq. (Melikoff, Z. anorg. 1899, 20. 342.)

Rubidium percolumbate, Rb_2CbO_8 .

Sol. in H_2O . Insol. in alcohol. (E. F. Smith.)

Sodium percolumbate, Na_2CbO_8 .

Sol. in H_2O . Insol. in alcohol. (E. F. Smith.)

Perferricyanhydric acid.

Potassium perferricyanide, $\text{K}_3\text{Fe}(\text{CN})_6 + \text{H}_2\text{O}$ (?).

Very hygroscopic, and sol. in H_2O . Nearly insol. in absolute alcohol. Decomp. by hot H_2O . (Skraup, A. 189. 368.)

Periodic acid, H_5IO_6 .

Deliquescent in moist air; very sol. in H_2O . (Bengieser, A. 17. 254.)

Rather easily sol. in alcohol and ether. (Bengieser.)

Rather easily sol. in alcohol, less in ether. (Langtoch.)

Sl. sol. in alcohol, still less in ether. (Langlois, J. pr. 56. 36.)

Sp. gr. of $\text{H}_5\text{IO}_6 + \text{Aq}$.

$\text{H}_5\text{IO}_6 + 20\text{H}_2\text{O} = 1.4008$.

$\text{H}_5\text{IO}_6 + 40\text{H}_2\text{O} = 1.2165$.

$\text{H}_5\text{IO}_6 + 80\text{H}_2\text{O} = 1.1121$.

$\text{H}_5\text{IO}_6 + 160\text{H}_2\text{O} = 1.0570$.

$\text{H}_5\text{IO}_6 + 320\text{H}_2\text{O} = 1.0288$.

(Thomsen, B. 7. 71.)

Periodates.

Most periodates are insol. or sl. sol. all are insol. or very sl. sol. in aq. they all dissolve in dil. HNO_3 , + Aq gieser.)

Aluminum metaperiodate, $\text{Al}(\text{IO}_4)_3 +$

Stable in solution containing (Eakle, C. C. 1896. II, 649.)

Ammonium metaperiodate, NH_4IO_4 .

Sl. sol. in H_2O . Cryst. with $3\text{H}_2\text{O}$ (3. 316), $2\text{H}_2\text{O}$ (Langlois, A. ch. (3) :

Stable in solution containing free : (Eakle, Zeit. Kryst. 1896, 26. 258-8. 100 pts. H_2O dissolve 2.7 pts. N 16° ; sp. gr. of sat. solution at $16^\circ/4^\circ$ (Barker, Chem. Soc. 1908, 93. 17.)

Ammonium dimesoperiodate, $(\text{NH}_4)_2\text{I}_2\text{O}_8 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Rammelsberg, P 379.)

Stable in solution containing free : Two modifications. (Eakle, Zeit. 1896, 26. 558-88; C. C. 1896. II, 64

Ammonium lithium dimesoperiodate
 $(\text{NH}_4)_2\text{Li}_2\text{I}_2\text{O}_8 + 7\text{H}_2\text{O}$.

Sol. in H_2O . (Ihre.)

Ammonium magnesium mesoperiodate
 $\text{NH}_4\text{MgIO}_6 + 3\text{H}_2\text{O}$.

Precipitate. (Rammelsberg, P 510.)

Barium metaperiodate, $\text{Ba}(\text{IO}_4)_2$.

Known only in solution.

Barium dimesoperiodate, $\text{Ba}_2\text{I}_2\text{O}_8$.

Sl. sol. in H_2O ; easily sol. in dil. Aq. (Rammelsberg, Pogg. 134. 391. Cryst. also with $3\text{H}_2\text{O}$, $5\text{H}_2\text{O}$, and

Barium mesoperiodate, $\text{Ba}_2(\text{IO}_4)_2 + 6$
 (Ihre.)

Barium orthoperiodate, $\text{Ba}_4(\text{IO}_4)_4$.

Insol. in H_2O . Sol. in HNO_3 , + Aq melsberg.)

Barium dimesodiperiodate, $\text{Ba}_4\text{I}_4\text{O}_{16}$.

Precipitate. Sol. in dil. HCl (Rammelsberg, Pogg. 134. 395.)

Barium periodate tungstate.

See Tungstoperiodate, barium.

Cæsium metaperiodate, CsIO_4 .

Sl. sol. in cold H_2O ; readily sol. in (Wells, Am. Ch. J. 1901, 26. 279.)

2.15 pts. are sol. in 100 pts. H_2O . Sp. gr. of the sat. aq. solution at 1.0166 . (Barker, Chem. Soc. 1908,

iodate hydrogen fluoride, 2CsIO_4 , H_2O .

40–60% $\text{HF} + \text{Aq}$. Decomp. by prescent. (Weinland, Z. anorg. 13.)

metaperiodate, $\text{Cd}(\text{IO}_4)_2$.

Rammelsberg, Pogg. 134. 516.)

dimesoperiodate, $\text{Cd}_2\text{I}_2\text{O}_9 + 9\text{H}_2\text{O}$.

H_2O . (Rammelsberg.)

mesoperiodate, $\text{Cd}_3(\text{IO}_4)_2 + 5\text{H}_2\text{O}$.

(Kimmins, Chem. Soc. 55. 151.)

diperiodate, $\text{Cd}_4\text{I}_2\text{O}_{11} + 3\text{H}_2\text{O}$.

H_2O . (Rammelsberg.)

eriodate, $\text{Cd}_{10}\text{I}_6\text{O}_{31} + 15\text{H}_2\text{O}$.

H_2O . (Rammelsberg.)

ta-periodate, $\text{Ca}(\text{IO}_4)_2$.

$\text{IO}_4 + \text{Aq}$ and acids. (Rammelsberg, 134. 405.)

dimesoperiodate, $\text{Ca}_2\text{I}_2\text{O}_9 + 7\text{H}_2\text{O}$, I_2O_7 .

H_2O . (Rammelsberg.)

(Langlois.)

hoperiodate, $\text{Ca}_3(\text{IO}_4)_2$.

H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$. (Rammelsberg, 44. 577.)

eriodate, 7CoO , $2\text{I}_2\text{O}_7 + 18\text{H}_2\text{O}$.

by HCl , and sol. on warming. Completely sol. in NHO_3 . (Lautsch, 19.)

It be obtained by Rammelsberg.

soperiodate, $\text{Cu}_2\text{I}_2\text{O}_9 + 6\text{H}_2\text{O}$.

by H_2O without dissolving. (Rammelsberg.)

periodate, Cu_2HIO_6 .

in $\text{HNO}_3 + \text{Aq}$. (Kimmins, Chem. Soc. 55. 149.)

riodate, $\text{Cu}_4\text{I}_2\text{O}_{11} + \text{H}_2\text{O}$.

H_2O ; sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Rammelsberg, 137. 313.)

odate, 5CuO , $\text{I}_2\text{O}_5 + 5\text{H}_2\text{O}$.

sol. in H_2O . (Rammelsberg, B. 1. 132.)

peroidate, $\text{Di}_2\text{O}_2(\text{IO}_4)_2$.

H_2O . Ppt. (Cleve, Bull. Soc. (2) 1911, 99. 1326.)

Erbium periodate.

Sol. in H_2O . (Höglund.)

Glucinum periodate, $\text{Gl}_2(\text{IO}_4)_2 + 11\text{H}_2\text{O}$.

Decomp. by H_2O without dissolving. Easily sol. in $\text{HNO}_3 + \text{Aq}$.

+ $13\text{H}_2\text{O}$. Nearly insol. in H_2O . (Atterberg, B. 7. 474.)

Iron (ferrous) orthoperiodate, $\text{Fe}_2(\text{IO}_4)_2$.

(Kimmins, Chem. Soc. 55. 150.)

FeH_2IO_6 . (Kimmins.)

Iron (ferric) periodate, $2\text{Fe}_2\text{O}_3$, $\text{I}_2\text{O}_7 + 21\text{H}_2\text{O}$.

Ppt. (Rammelsberg.)

Iron (ferric) dimesoperiodate, FeHI_2O_9 .

Insol. in dil. $\text{HNO}_3 + \text{Aq}$. (Kimmins, Chem. Soc. 55. 149.)

Iron (ferric) metaperiodate, $\text{Fe}(\text{IO}_4)_3$.

(Kimmins.)

Lanthanum periodate, $\text{La}(\text{IO}_4)_3 + 2\text{H}_2\text{O}$.

Precipitate. (Cleve.)

Lead metaperiodate, $\text{Pb}(\text{IO}_4)_2$.

Sol. in $\text{HNO}_3 + \text{Aq}$. (Kimmins.)

Lead orthoperiodate, $\text{Pb}_2\text{H}_4(\text{IO}_4)_2$.

Sol. in $\text{HNO}_3 + \text{Aq}$. (Kimmins, Chem. Soc. 55. 149.)

Lead mesoperiodate, $\text{Pb}_3(\text{IO}_4)_2 + 2\text{H}_2\text{O}$.

Insol. in H_2O or excess of periodic acid + Aq . Decomp. by dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Bengieser, A. 17. 254.)

Lithium metaperiodate, LiIO_4 .

Difficultly sol. in H_2O . (Rammelsberg, B. 1. 132.)

Somewhat deliquescent.

+ H_2O ; sol. in H_2O . (Barker, Chem. Soc. 1911, 99. 1326.)

Lithium dimesoperiodate, $\text{Li}_4\text{I}_2\text{O}_9 + 3\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (Rammelsberg, Pogg. 134. 387.)

Lithium orthoperiodate, Li_5IO_6 .

H_2O dissolves out a slight amount of LiI . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Rammelsberg, Pogg. 137. 313.)

Magnesium metaperiodate, $\text{Mg}(\text{IO}_4)_2 + 10\text{H}_2\text{O}$.

Easily sol. in H_2O . (Rammelsberg.)

Magnesium diperiodate, $\text{Mg}_4\text{I}_2\text{O}_{11} + 6\text{H}_2\text{O}$, or $9\text{H}_2\text{O}$.

Sl. efflorescent. Insol. in H_2O . (Rammelsberg.)

Magnesium dimesoperiodate, $\text{Mg}_2\text{I}_2\text{O}_8 + 3\text{H}_2\text{O}$.

(Rammelsberg, Pogg. 134. 499.)
+15 H_2O . Insol. in H_2O . Sol. in periodic acid + Aq. (Langlois.)

Manganic periodate.

See Manganiperiodic acid.

Mercurous diperiodate, $5\text{Hg}_2\text{O}$, I_2O_7 , or $4\text{Hg}_2\text{O}$, $\text{I}_2\text{O}_7 = \text{Hg}_4\text{I}_2\text{O}_{11}$.

Insol. in H_2O . Easily sol. in HNO_3 + Aq and in HCl + Aq. (Lautsch, J. pr. 100. 86.)

Mercuric orthoperiodate, $\text{Hg}_2(\text{IO}_6)_2$.

Insol. in H_2O . Easily sol. in HCl . Sl. sol. in HNO_3 . (Lautsch.)

Mercuric potassium periodate, 10HgO , $5\text{K}_2\text{O}$, $6\text{I}_2\text{O}_7$.

Insol. in H_2O . Difficultly sol. in warm HNO_3 without decomp. (Rammelsberg, Pogg. 134. 526.)

Nickel dimesoperiodate, $\text{Ni}_2\text{I}_2\text{O}_8$.

(Kimmins, Chem. Soc. 55. 151.)

Nickel mesoperiodate, $\text{Ni}_2(\text{IO}_6)_2$.

(Kimmins.)

Nickel periodate, 7NiO , $4\text{I}_2\text{O}_7 + 63\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in H_2IO_6 + Aq. (Rammelsberg, Pogg. 134. 514.)

Potassium metaperiodate, KIO_4 .

Sl. sol. in H_2O . Sol. in 290 pts. cold H_2O . (Rammelsberg, Pogg. 134. 320.)

Almost insol. in KOH + Aq.
0.66 pt. is sol. in 100 pts. H_2O at 13° . Sp. gr. of the sat. sol. at $13^\circ/4^\circ = 1.0051$. (Barker, Chem. Soc. 1908, 93. 16.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Potassium mesoperiodate, $\text{K}_2\text{IO}_6 + 4\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O . (Ihre.)

Potassium dimesoperiodate, $\text{K}_4\text{I}_2\text{O}_8 + 9\text{H}_2\text{O}$.

Sol. in 9.7 pts. cold H_2O . (Rammelsberg, Pogg. 134. 320.)

Sol. in KOH + Aq.
+3 H_2O .

Potassium hydrogen dimesoperiodate, $\text{K}_2\text{HI}_2\text{O}_8$.

Less sol. in H_2O than KIO_4 . (Kimmins, Chem. Soc. 51. 356.)

Potassium manganic periodate.

See Manganiperiodate, potassium.

Potassium zinc periodate, K_2O , 4ZnO , $2\text{I}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Ppt. (Rammelsberg, Pogg. 134. 368.)

Potassium periodate tungstate.

See Tungstoperiodate, potassium.

Rubidium periodate, RbIO_4 .

0.65 pt. is sol. in 100 pts. H_2O at 1 gr. of the sat. aq. solution at $13^\circ/4^\circ$ = (Barker, Chem. Soc. 1908, 93. 16.)

Samarium periodate, $\text{Sm}(\text{IO}_6) + 4\text{H}_2\text{O}$
Precipitate. (Cleve.)

Silver metaperiodate, AgIO_4 .

Decomp. by cold H_2O into $\text{Ag}_2\text{I}_2\text{O}_8$ and by warm H_2O into $\text{Ag}_2\text{I}_2\text{O}_8 + \text{H}_2\text{O}$ (Mermüller and Magnus, Pogg. 28. 51 + H_2O . Insol. ppt. (Kimmins.)

Silver mesoperiodate, Ag_2IO_6 .

(Fernlunds, J. pr. 100. 99.)
 Ag_2HIO_6 . Insol. ppt. (Kimmins Soc. 51. 358.)
Ppt. by dil. HNO_3 ; sol. in HNO_3 . heim, A. 1899, 308. 57.)

Silver dimesoperiodate, $\text{Ag}_2\text{I}_2\text{O}_8 + 3\text{H}_2\text{O}$.

Insol. ppt. (Kimmins.)
Decomp. by boiling H_2O into (Rammelsberg.)

Silver orthoperiodate, Ag_2IO_6 .

Sol. in HNO_3 or NH_4OH + Aq. (Rammelsberg, Pogg. 134. 386.)

Sol. in excess NH_4OH + Aq; ppt. HNO_3 . (Rosenheim, A. 1899, 308. 5)

$\text{Ag}_2\text{H}_2\text{IO}_6$. Insol. ppt. (Kimmins Soc. 51. 358.)

$\text{Ag}_2\text{H}_2\text{IO}_6$. As above. (Kimmins.)
Sol. in dil. HNO_3 . (Rosenheim, 308. 53.)

Silver diperiodate, $\text{Ag}_2\text{I}_2\text{O}_{11}$.

Sl. sol. in HNO_3 + Aq; insol. in NH_4OH + Aq. (Lautsch, J. pr. 100. 75.)

Silver dimesodiperiodate, $\text{Ag}_{10}\text{I}_2\text{O}_{18}$.

HNO_3 + Aq dissolves out Ag_2O . 1 NH_4OH + Aq. (Lautsch.)

Sodium metaperiodate, NaIO_4 .

Easily sol. in H_2O .
+2 H_2O . (Langlois.)
+3 H_2O . Efflorescent; sol. in 12 p at ord. temp. (Rammelsberg, J. pr. 11)

Sodium dimesoperiodate, $\text{Na}_4\text{I}_2\text{O}_8 + 3\text{H}_2\text{O}$.

Scarcely sol. in cold, sl. sol. in h (Magnus and Ammermüller, Pogg. 28)

Very sol. in dil. HNO_3 + Aq. (Langlois)

Sol. in $\text{HC}_2\text{H}_3\text{O}_2$ + Aq with (Bengieser, A. 17. 254.)

Insol. in methyl acetate. (Naumann, 1909, 42. 3790.)
+4 H_2O .

periodate, $\text{Na}_2\text{IO}_6 + \frac{1}{4}\text{H}_2\text{O}$.
O. (Ihre.)
 $\text{Na}_2\text{H}_2\text{IO}_6$. Less sol. in H_2O than
 $\text{H}_2\text{O} (= \text{Na}_2\text{H}_2\text{IO}_6)$. (Kimmins,
51. 357.)

periodate, Na_2IO_6 .
Correct composition for
 I_2O_7 . (Kimmins.)
Correct composition for Na_2IO_6 ,
mmms.)

etaperiodate, $\text{Sr}(\text{IO}_4)_2 + 6\text{H}_2\text{O}$.
O.

mesoperiodate, $\text{Sr}_2\text{I}_2\text{O}_9$.
by H_2O .

esoperiodate, $\text{Sr}_2(\text{IO}_4)_2$.
e.

thoperiodate, $\text{Sr}_2(\text{IO}_4)_2$.
berg, Pogg. 44. 577.)

date, $3\text{Tl}_2\text{O}_3, \text{I}_2\text{O}_7 + 30\text{H}_2\text{O}$.
 H_2O . Decomp. by alkalis.
rg, B. 3. 361.)

iodate.
e. Sol. in $\text{HNO}_3 + \text{Aq}$.

iodate.
e, which quickly decomposes.

eriodate, $\text{YbIO}_6 + 2\text{H}_2\text{O}$.
ic. (Cleve, Z. anorg. 1902, 32.)

odate, $\text{Y}_2(\text{IO}_4)_2 + 8\text{H}_2\text{O}$.
tly sol. (Cleve.)
 $\text{O}_7 + 6\text{H}_2\text{O}$. Precipitate. (Cleve.)

eriodate, $\text{Zn}_2\text{I}_2\text{O}_9 + 6\text{H}_2\text{O}$.
berg, Pogg. 134. 513.)

te, $3\text{ZnO}, 2\text{I}_2\text{O}_7 + 7\text{H}_2\text{O}$.
)

late, $\text{Zn}_4\text{I}_2\text{O}_{11} + \text{H}_2\text{O}$.
. in H_2O , sl. acid with HNO_3 .
. ch. (3) 34. 257.)

kiperiodate, $\text{Zn}_5\text{I}_4\text{O}_{19} + 14\text{H}_2\text{O} (?)$.
berg.)

Zinc periodate, $9\text{ZnO}, 2\text{I}_2\text{O}_7 + 12\text{H}_2\text{O}$.
(Rammelsberg.)

Periodoplatinocyanhydric acid.

**Barium periodoplatinocyanide, $\text{BaPt}(\text{CN})_4\text{I}_2$
 $+ x\text{H}_2\text{O}$.**
Easily sol. in H_2O or alcohol. (Holst, Bull.
Soc. (2) 22. 347.)

**Potassium periodoplatinocyanide,
 $\text{K}_2\text{Pt}(\text{CN})_4\text{I}_2$.**
Permanent. Easily sol. in H_2O or alcohol.

Permanganic acid, HMnO_4 .
Known only in solution, which decomposes
by evaporation or warming.

Permanganates.
All permanganates are sol. in H_2O , except-
ing AgMnO_4 , which is sl. sol.

Ammonium permanganate, NH_4MnO_4 .
Sol. in 12.6 pts. H_2O at 15° . (Aschoff.)
Sol. in H_2O with decomp. (Christensen,
Z. anorg. 1900, 24. 206.)

Barium permanganate, $\text{Ba}(\text{MnO}_4)_2$.
Sol. in H_2O .

Cadmium permanganate, $\text{Cd}(\text{MnO}_4)_2 + 8\text{H}_2\text{O}$.
Stable. (Klobb, Bull. Soc. 1894, (3) 11
607.)

**Cadmium permanganate ammonia,
 $\text{Cd}(\text{MnO}_4)_2, 4\text{NH}_3$.**
Sol. in H_2O with decomp. (Klobb, Bull.
Soc. (3) 3. 510.)

Cæsium permanganate, CsMnO_4 .
Sl. sol. in cold, somewhat more easily sol.
in hot H_2O . (Muthmann, B. 1893, 26. 1018.)
Solubility in H_2O .
100 ccm. of the sat. solution contain at:

1°	19°	59°
0.097	0.23	1.25 g. CsMnO_4 .

(Patterson, J. Am. Chem. Soc. 1906, 28.
1735.)

Calcium permanganate, $\text{Ca}(\text{MnO}_4)_2 + 5\text{H}_2\text{O}$.
Deliquescent.

Cupric permanganate.
Deliquescent.

Cupric permanganate ammonia, $\text{Cu}(\text{MnO}_4)_2 \cdot 4\text{NH}_3$.

Sol. in H_2O with slow decomp. (Klobb, Bull. Soc. (3) 3. 509.)

Didymium permanganate, $\text{Di}(\text{MnO}_4)_2 + 21\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Frerichs and Smith, A. 191. 331.)

Has not been prepared. (Cleve, B. 11. 912.)

Lanthanum permanganate, $\text{La}(\text{MnO}_4)_3 + 21\text{H}_2\text{O}$.

Ppt. (Frerichs and Smith, A. 191. 331.)

Has not been prepared. (Cleve, B. 11. 910.)

Lead permanganate.

Sol. in $\text{HNO}_3 + \text{Aq}$. (Forchhammer.)

Lithium permanganate, $\text{LiMnO}_4 + 3\text{H}_2\text{O}$.

Sol. in 1.4 pts. H_2O at 16° . (Aschoff.)

Magnesium permanganate, $\text{Mg}(\text{MnO}_4)_2$.

Insol. in CHCl_3 , CCl_4 , C_6H_6 , toluene, nitrobenzene, ligroin, ether and CS_2 . Sol. in methyl alcohol, acetone, pyridine, and readily sol. in glacial acetic acid. Only pyridine and glacial acetic acid are sufficiently stable toward the salt to be of any practical use for oxidation purposes. (Michael and Garner, Am. Ch. J. 1906, 35. 288.)

+6 H_2O . Easily deliquescent.

Nickel permanganate ammonia, $\text{Ni}(\text{MnO}_4)_2 \cdot 4\text{NH}_3$.

Sol. in H_2O with decomp. (Klobb, Bull. Soc. (3) 3. 509.)

Potassium permanganate, KMnO_4 .

Sol. in 16 pts. H_2O at 15° . (Mitscherlich.)

Solubility in 100 pts. H_2O at t° .

t°	Pts. KMnO_4
0	2.83
9.8	4.31
19.8	6.34
24.8	7.59
29.8	9.03
34.8	10.67
40.0	12.56
45.0	14.58
50.0	16.89
55.0	19.33
65.0	25.03

(Baxter, J. Am. Chem. Soc. 1906, 28. 1343.)

Solubility in H_2O at t° .

p = pts. KMnO_4 sol. in 100 pts. H_2O

t°	p	t°	p	t°	p
0	2.76	10	6.26	38	11.74
1	2.90	20	6.48	39	12.12
2	3.06	21	6.70	40	12.51
3	3.22	22	6.94	41	12.91
4	3.38	23	7.18	42	13.31
5	3.54	24	7.42	43	13.72
6	3.70	25	7.68	44	14.14
7	3.86	26	7.94	45	14.56
8	4.04	27	8.20	46	15.00
9	4.22	28	8.48	47	15.44
10	4.40	29	8.77	48	15.88
11	4.58	30	9.07	49	16.32
12	4.78	31	9.37	50	16.77
13	4.98	32	9.69	51	17.23
14	5.18	33	10.01	52	17.71
15	5.38	34	10.34	53	18.21
16	5.60	35	10.68	54	18.71
17	5.82	36	11.02	55	19.23
18	6.04	37	11.38	56	19.75

(Worden, J. Soc. Chem. Ind. 190)

Solubility in H_2O .

100 cc. of the sat. solution con

0° 15° 15.3° 30°
2.84 5.22 5.30 8.69 g. l

Sp. gr. of sat. solution at
(Patterson, J. Am. Chem. Soc. 1735.)

1 l. sat. $\text{KMnO}_4 + \text{Aq}$ contains :

0° 10° 20° 30° 40°
0.176 0.278 0.411 0.573 0.792 mo

53° 63° 70° 75°
1.154 1.429 1.812 2.047 mo

(Sackur, Z. Elektrochem. 1912,

Solubility of KMnO_4 in H

Grams KMnO_4 sol. in 100 grams H_2O	t°
0.58	—
1.01	—
2.02	—
2.91	—
4.22	+10
5.20	+12
7.53	+22
11.61	+46
16.75	+56

(Voerman, C. C. 1906, I. 1

Sol. in conc. H_2SO_4 . Deliquescent
 HCl , but does not dissolve. (Gor
Slowly sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$. (Cl
Edwards.)

PERMANGANATE AMMONIA, SILVER

solubility in KOH + Aq at t° expressed in mol. per l. of the sat. solution.

H ₂ O	1-n KOH	2-n KOH	4-n KOH	6-n KOH	8-n KOH	10-n KOH
0.176	0.050	0.031	0.027	0.023	0.017	0.012
0.278	0.112	0.068	0.048	0.042	0.028	0.016
0.411	0.179	0.119	0.079	19° 0.074	0.032	0.029
0.573	32° 0.316	32° 0.213	32° 0.149	0.114	32° 0.062	0.040
0.792	0.439	0.306	0.211	0.161	0.084	0.052
3° 1.154	50° 0.638	0.462	0.304	0.219	0.111	...
1.429	61° 0.904	60° 0.639	0.427	0.291	61° 0.143	0.071
1.812	1.172	0.869	0.572	0.390	0.188	0.082
2.047	0.651	0.089
....	1.513	1.190	...	0.500	0.231	...
....	1.655	1.352	83° 0.803	85° 0.572
....	0.649	0.297	...

(Sackur, Z. Elektrochem. 1912, 18. 723.)

Solubility in salts + Aq. at t°.

Solvent	t°	Mol. KMnO ₄ in 1 l. of sat. solution
0.1-n $\frac{K_2CO_3}{2}$	0	0.1462
	25	0.4375
	40	0.7380
1-n $\frac{K_2CO_3}{2}$	0	0.0629
	25	0.2589
	40	0.5007
2-n $\frac{K_2CO_3}{2}$	0	0.0446
	40	0.3519
4-n $\frac{K_2CO_3}{2}$	0	0.0270
	25	0.0930
6-n $\frac{K_2CO_3}{2}$	0	0.0156
0.1-n KCl	0	0.1395
	25	0.4315
	40	0.7380
0.5-n KCl	0	0.0760
	25	0.3060
	40	0.5840
1-n KCl	0	0.0532
	25	0.220
	40	0.444
2-n KCl	0	0.0379
	25	0.1432
	40	0.288

Sackur, Z. Elektrochem. 1912, 18. 723.)

ery sol. in liquid NH₃. (Moissan, A. ch. (7) 6. 428; Franklin, Am. Ch. J. 1898, 29.)

decomp. immediately by alcohol. Sol. in me. (Eidmann, C. C. 1899. II, 1014; mann. B. 1904, 37. 4328.)

Solubility in acetone + Aq. at 13°.

A = ccm. acetone in 100 ccm. acetone + Aq.
1/5 KMnO₄ = millimols KMnO₄ in 100 ccm. of the solution.

A	1/5 KMnO ₄
0	148.5
10	162.2
20	177.3
30	208.2
40	257.4
50	289.7
60	316.8
70	328.0
80	312.5
90	227.0
100	67.6

(Herz and Knoch, Z. anorg. 1904, 41. 317.)

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3795.)

Sol. in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Rubidium permanganate, RbMnO₄.

Solubility in H₂O lies between K and Cs salts. (Muthmann, B. 1893, 26. 1018.)

Solubility in H₂O.

100 ccm. of the sat. solution contain at:

2° 19° 60°
0.46 1.06 4.68 g. RbMnO₄.

(Patterson, J. Am. Chem. Soc. 1906, 28 1735.)

Silver permanganate, Ag₂MnO₄.

Sol. in 109 pts. cold H₂O and much less h H₂O. Decomp. by boiling. (Mitscherlic Pogg. 25. 301.)

Silver permanganate ammonia.

Sl. sol. in cold, more easily in hot F (Klobb, C. R. 103. 384.)

Sodium permanganate, $\text{NaMnO}_4 + 3\text{H}_2\text{O}$.

Deliquescent. Extremely sol. in H_2O .

Moderately sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Strontium permanganate, $\text{Sr}(\text{MnO}_4)_2 + 4\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . (Fromherz.)

Thallous permanganate, TlMnO_4 .

Sol. in H_2O with decomp. (R. Meyer, Z. anorg. 1899, 22. 188.)

Zinc permanganate, $\text{Zn}(\text{MnO}_4)_2 + 6\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O . (Martenson, J. B. 1873. 274.)

Zinc permanganate ammonia,

$\text{Zn}(\text{MnO}_4)_2, 4\text{NH}_3$.

Sol. in H_2O with decomp. (Klobb, Bull. Soc. (3) 3. 509.)

Permanganomolybdic acid, MnO_2 , $12\text{MoO}_3 + 10\text{H}_2\text{O}$.

Sol. in H_2O . Decomp. by alkalis. Sol. in alcohol. (Péchar, C. R. 1897, 125. 31.)

Ammonium permanganomolybdate,

$2(\text{NH}_4)_2\text{O}, \text{MnO}_2, 7\text{MoO}_3 + 5\text{H}_2\text{O}$.

(Friedheim and Samelson, Z. anorg. 1900, 24. 73.)

$3(\text{NH}_4)_2\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 6\text{H}_2\text{O}$. (Friedheim and Allemann, Mit. d. Nat. Ges. Bern. 1904. 23.)

$+ 7\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 70.)

$4(\text{NH}_4)_2\text{O}, \text{MnO}_2, 11\text{MoO}_3 + 7\text{H}_2\text{O}$. (Friedheim and Samelson.)

$3(\text{NH}_4)_2\text{O}, \text{MnO}_2, 12\text{MoO}_3 + 5\text{H}_2\text{O}$. Sl. sol. in cold H_2O . Decomp. by alkalis. Insol. in alcohol. (Péchar, C. R. 1897, 125. 30.)

Ammonium manganous permanganomolybdate, $3[(\text{NH}_4)_2, \text{Mn}]\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 6\text{H}_2\text{O}$ and $+ 7\text{H}_2\text{O}$.

(Friedheim and Allemann, Mitt. d. Nat. Ges. Bern. 1904. 23.)

$3[(\text{NH}_4)_2, \text{Mn}]\text{O}, \text{MnO}_2, 10\text{MoO}_3 + 10\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 94.)

$4[(\text{NH}_4)_2, \text{Mn}]\text{O}, \text{MnO}_2, 10\text{MoO}_3 + 6\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 75.)

$4[(\text{NH}_4)_2, \text{Mn}]\text{O}, \text{MnO}_2, 11\text{MoO}_3 + 8\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 72.)

Ammonium manganous potassium permanganomolybdate, $2(\text{NH}_4)_2\text{O}, \text{MnO}, \text{K}_2\text{O}, \text{MnO}_2, 10\text{MoO}_3 + 5\text{H}_2\text{O}$.

Very sl. sol. in cold, easily sol. in H_2O at $70-80^\circ$. (Rosenheim, Z. anorg. 1898, 16. 79.)

$3[(\text{NH}_4)_2, \text{K}, \text{Mn}]\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 7\text{H}_2\text{O}$. (Friedheim and Allemann, Mitt. d. Nat. Ges. Bern. 1904. 23.)

$4[(\text{NH}_4)_2, \text{K}, \text{Mn}]\text{O}, \text{MnO}_2, 10\text{MoO}_3 + 5\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 97.)

$3[(\text{NH}_4)_2, \text{K}, \text{Mn}]\text{O}, \text{MnO}_2, 10\text{MoO}_3 + 6\text{H}_2\text{O}$, and $+ 10\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 92.)

Ammonium potassium permanganate, $3[(\text{NH}_4)_2, \text{K}, \text{Mn}]\text{O}, \text{MnO}_2, 4\text{H}_2\text{O}$.

(Friedheim and Samelson.)

Barium permanganomolybdate, $3\text{BaO}, 9\text{MoO}_3 + 12\text{H}_2\text{O}$.

Ppt. (Hall, J. Am. Chem. Soc. 1900, 22. 700.)

Manganous potassium permanganate, $2\text{K}_2\text{O}, \text{MnO}, \text{MnO}_2, 8\text{H}_2\text{O}$.

True formula for $5\text{K}_2\text{O}, \text{Mn}_2\text{O}_3, 12\text{H}_2\text{O}$ of Struve. (Friedheim and Samelson, Z. anorg. 1900, 24. 86.)

$3[\text{K}_2, \text{Mn}]\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 6\text{H}_2\text{O}$. (Friedheim and Allemann, Mitt. d. Nat. Ges. Bern. 1904. 23.)

$2.6 \text{K}_2\text{O}, 0.4 \text{MnO}, \text{MnO}_2, 9\text{MoO}_3$. Ppt. (Hall, J. Am. Chem. Soc. 1900, 22. 700.)

$4[\text{K}_2, \text{Mn}]\text{O}, \text{MnO}_2, 11\text{MoO}_3 + 7\text{H}_2\text{O}$. (Friedheim and Samelson, Z. anorg. 1900, 24. 80.)

Manganous potassium sodium permanganomolybdate, $3[\text{K}_2, \text{Na}, \text{Mn}]\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 4\text{H}_2\text{O}$.

(Friedheim and Allemann, Mitt. d. Nat. Ges. Bern. 1904. 48.)

Manganous sodium permanganomolybdate, $3[\text{Na}_2, \text{Mn}]\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 4\text{H}_2\text{O}$.

(Friedheim and Allemann.)

Potassium permanganomolybdate, $\text{MnO}_2, 8\text{MoO}_3 + 3\text{H}_2\text{O}$.

Much less sol. in H_2O than sodium permanganomolybdate. (Friedheim and Samelson, Z. anorg. 1900, 24. 78.)

$+ 5\text{H}_2\text{O}$. Nearly insol. in cold H_2O . (Rosenheim and Itzig, Z. anorg. 1898, 16. 79.)

$3\text{K}_2\text{O}, \text{MnO}_2, 9\text{MoO}_3 + 5\text{H}_2\text{O}$. and Samelson, Z. anorg. 1900, 24. 75.)

$+ 6\text{H}_2\text{O}$. (Hall, J. Am. Chem. Soc. 1900, 22. 700.)

$3\text{K}_2\text{O}, \text{MnO}_2, 12\text{MoO}_3 + 4\text{H}_2\text{O}$. insol. in cold H_2O . Decomp. Insol. in alcohol. (Péchar, C. R. 1897, 125. 31.)

Silver permanganomolybdate, $3\text{Ag}_2\text{O}, 9\text{MoO}_3 + 6\text{H}_2\text{O}$.

Ppt. (Hall, J. Am. Chem. Soc. 1900, 22. 700.)

manganomolybdate, $3\text{Na}_2\text{O}$, $12\text{MoO}_3 + 13\text{H}_2\text{O}$.

ent. Very sol. in H_2O . Decomp. Insol. in alcohol. (Péchar, C. R.

notungstic acid.

manganous permanganotung-
 $4(\text{NH}_4)_2\text{O}$, MnO , MnO_2 , $12\text{WO}_3 +$

sol. in H_2O . Can be cryst. there-
ggers and Smith, J. Am. Chem.
26. 1475.)

manganotungstate, $3\text{Na}_2\text{O}$, MnO_2 ,
 $+18\text{H}_2\text{O}$.

easily sol. in hot H_2O . Solution
a long boiling with separation of
peroxide. (Just, B. 1903, 36.

dic acid, Mo_2O_7 , $5\text{H}_2\text{O} =$
 $4 + 2\text{H}_2\text{O}$.

in H_2O , and not decomp. by boil-
ard, A. ch. (6) 28. 550.)

$+1\frac{1}{2}\text{H}_2\text{O}$. "Ozo-molybdic acid."
sl. sol. in H_2O after being dried in
ol. in fairly conc. H_2SO_4 . (Muth-
898, 31. 1838.)

, H_2O_2 . Sl. sol. in cold, more eas-
not H_2O , but does not separate on
ol. in dil. acids, also in H_3PO_4 .
, Ch. Z. 1891, 15. 957.)

permolybdate, $\text{NH}_4\text{MoO}_4 +$

. in H_2O ; sl. sol. in alcohol, but
tracts it from H_2O , forming a very
sat. solution, which is pptd. by a
 NH_4MoO_4 , and only a sl. amount
solution. (Péchar, d.)

5MoO_3 , $2\text{MoO}_4 + 6\text{H}_2\text{O}$. (Muth-
898, 31. 1837.)

$7\text{MoO}_4 + 12\text{H}_2\text{O}$. Ppt. (Muth-
norg. 1898, 17. 76.)

$5\text{MoO}_4 + 6\text{H}_2\text{O}$. Ppt. (Muth-

nickel permolybdate ammonia,
 $2\text{Ni}(\text{MoO}_4)_2$, 2NH_3 .

by H_2O . Sol. in dil. NH_4OH .
nem. Soc. 1904, 85. 674.)

molybdate, $\text{Ba}(\text{MoO}_4)_2 + 2\text{H}_2\text{O}$.

, A. ch. 1893, (6) 28. 537.)

9MoO_3 , $2\text{H}_2\text{O}_2 + 13\text{H}_2\text{O}$. (Baer-
rt. 1885.)

rmolybdate, Cs_2O , $4\text{MoO}_4 + 6\text{H}_2\text{O}$.
 H_2O . (Muthmann, B. 1898, 31.

MoO_3 , $3\text{MoO}_4 + 4\text{H}_2\text{O}$. Ppt.
n.)

Copper permolybdate, $\text{Cu}(\text{MoO}_4)_2 + \text{H}_2\text{O}$.

Insol. in H_2O ; easily sol. in acids. Sol. in
 $\text{NH}_4\text{OH} + \text{Aq}$ with decomp. (Péchar, d.)

Magnesium permolybdate, $\text{Mg}(\text{MoO}_4)_2 +$
 $10\text{H}_2\text{O}$.

Very sol. in H_2O ; sl. sol. in alcohol. (Péch-
ard.)

Mercurous permolybdate.

Insol. in H_2O or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Péchar, d.)

Potassium permolybdate, $\text{KMoO}_4 + 2\text{H}_2\text{O}$.

Sl. sol. in cold, more in hot H_2O . Sl. sol.
in alcohol. (Péchar, d.)

K_2O , 2MoO_3 , $\text{MoO}_4 + 3\text{H}_2\text{O}$. Ppt. (Muth-
mann, Z. anorg. 1898, 17. 77.)

K_2O_2 , MoO_4 , H_2O_2 . Decomp. by H_2O .
(Melikoff and Pissarjewsky, B. 1898, 31.
2449.)

$\text{K}_2\text{MoO}_4 + 3\text{H}_2\text{O}$. Nearly insol. in cold,
easily sol. in hot H_2O . (Mazzuchelli and
Zangrilli, Gazz. ch. it. 1910, 40. (2) 56.)

Rubidium permolybdates.

"Rubidium ozo-molybdate."

$3\text{Rb}_2\text{O}$, $10\text{MoO}_4 + 14\text{H}_2\text{O}$. Ppt.

Rb_2O , 2MoO_3 , $\text{MoO}_4 + 3\text{H}_2\text{O}$. May be re-
cryst. from $\text{H}_2\text{O}_2 + \text{Aq}$.

$3\text{Rb}_2\text{O}$, 5MoO_3 , $2\text{MoO}_4 + 6\text{H}_2\text{O}$. Ppt.

Rb_2O , 3MoO_3 , $\text{MoO}_4 + 4\text{H}_2\text{O}$. Ppt. (Muth-
mann, B. 1898, 31. 1839-41.)

Silver permolybdate, AgMoO_4 .

(Péchar, d.)

Sodium permolybdate, $\text{NaMoO}_4 + 3\text{H}_2\text{O}$.

Very sol. in H_2O ; insol. in alcohol, but
behaves similarly to K salt. (Péchar, d.)

Thallous permolybdate.

Insol. in H_2O . (Péchar, A. ch. 1893, (6)
28. 559.)

Pernitric acid, NO_3 .

See Nitrogen hexoxide.

Silver pernitrate, basic, $3\text{Ag}_2\text{O}_3$, AgNO_3 .

Decomp. H_2O . (Mulder, R. t. c. 1898, 17.
142.)

Perosmic acid.

Potassium perosmate (?).

Sol. in H_2O , but very easily decomp.

Peroxynitric acid.

Silver peroxynitrate.

Analysis of the black compound formed,
under certain circumstances, in a silver volt-
ameter when an aqueous solution of AgNO_3

is electrolyzed, points to the composition $3\text{Ag}_2\text{O}$, 5O , AgNO_3 , perhaps $2\text{Ag}_2\text{O}_4$, AgNO_3 , or $3\text{Ag}_2\text{O}_3$, AgNO_3 . (Mulder, Chem. Soc. 1896, 70. (2) 561.)

Peroxyaminesulphonic acid.

Potassium peroxyaminesulphonate, $\text{N}_2\text{O}_4(\text{SO}_3\text{K})_4$.

Very unstable in H_2O . Very sl. sol. in cold H_2O . More stable in $\text{N}/10$ $\text{KOH} + \text{Aq}$.

100 pts. $\text{N}/10$ $\text{KOH} + \text{Aq}$ dissolve 0.62 pt. of the salt at 3° ; 6.6 pts. at 29° . (Haga, Chem. Soc. 1904, 85. 86.)

Perstannic acid, $\text{H}_2\text{Sn}_2\text{O}_7$.

Known in colloidal state, sol. in H_2O . (Spring, Bull. Soc. (2) 51. 180.)

Potassium perstannate, $\text{KSnO}_4 + 2\text{H}_2\text{O}$.

Sol. in H_2O . Insol. in alcohol. (Tanatar, B. 1905, 38. 1185.)

Sodium perstannate, $\text{NaSnO}_4 + 2\text{H}_2\text{O}$.

Difficultly sol. in H_2O with decomp. (Tanatar.)

Persulphuric acid, $\text{H}_2\text{S}_2\text{O}_8$.

See Sulphur heptoxide.

$\text{H}_2\text{S}_2\text{O}_8$.

Sp. gr. of $\text{H}_2\text{S}_2\text{O}_8 + \text{Aq}$.

Sp. gr. $14^\circ/14^\circ$	% $\text{H}_2\text{S}_2\text{O}_8$	g. $\text{H}_2\text{S}_2\text{O}_8$ per l.
1.042	7.2	75
1.096	15.4	169
1.154	23.6	272
1.246	35.2	438

(Elbs and Schönherr, Z. Elektrochem. 1896, 2. 245.)

Ammonium persulphate, $(\text{NH}_4)_2\text{S}_2\text{O}_8$.

Very sol. in H_2O . 100 pts. H_2O at 0° dissolve 58.2 pts. $(\text{NH}_4)_2\text{S}_2\text{O}_8$. (Marshall, Chem. Soc. 59. 771.)

Solubility in H_2O equals 58% at 8° . (Moreau, C. C. 1901, II. 56.)

100 pts. H_2O dissolve 65 pts. at ord. temp. (Elbs, J. pr. 1893, (2) 48. 185.)

Ammonium lead persulphate, $(\text{NH}_4)_2\text{Pb}(\text{SO}_4)_2$.

Decomp. by H_2O . Almost insol. in cold H_2SO_4 of sp. gr. = 1.7. Sl. sol. in H_2SO_4 (sp. gr. = 1.7) at 50° . Sol. in fuming H_2SO_4 and in cold conc. HCl . Sol. in acetic acid, in Na acetate + Aq acidified with acetic acid and in excess of cold 20% $\text{NaOH} + \text{Aq}$. (Elbs, Z. Elektrochem. 1900, 7. 346.)

Ammonium mercurous persulphate ammonia, $(\text{NH}_4)_2\text{HgS}_2\text{O}_8 \cdot 2\text{NH}_3$.

Decomp. by H_2O . Insol. in dil. hot or cold H_2SO_4 or HNO_3 . Sol. (Tarugi, Gazz. ch. it. 1903, 33. (1) 11.)

Barium persulphate, $\text{BaS}_2\text{O}_8 + 4\text{H}_2\text{O}$.

Very sol. in H_2O . 100 pts. H_2O solve 39.1 pts. BaS_2O_8 , or 52.2 pts. $\text{BaS}_2\text{O}_8 + 4\text{H}_2\text{O}$. Sol. in absolute alcohol with $\text{BaS}_2\text{O}_8 + \text{H}_2\text{O}$. Insol. in alcohol. (M.)

Cadmium persulphate ammonia, 6NH_3 .

Sol. in H_2O . (Barbieri, Z. anorg. 350.)

Cæsium persulphate, $\text{Cs}_2\text{S}_2\text{O}_8$.

Sol. in H_2O . 8.71–8.98 pts. are 1 pt. H_2O at 23° . (E. F. Smith, J. A. Soc. 1899, 21. 935.)

Calcium persulphate.

Very sol. in H_2O . (Marshall, J. S. Ind. 1897, 16. 396.)

Copper persulphate ammonia, CuS_2O_8 .

Sol. in H_2O . (Barbieri, Z. anorg. 351.)

Lead persulphate, PbS_2O_8 .

Decomp. by H_2O . Sl. sol. in H_2SO_4 pyrosulphuric acid. Sol. in cold H_2SO_4 . Insol. or sol. with decomp. in all ord. (Elbs, Z. Elektrochem. 1900, 7. 345.)

Solubility of $\text{Pb}(\text{SO}_4)_2$ in $\text{H}_2\text{SO}_4 + \text{Aq}$
v = moles of H_2SO_4 per mole
c = millimols $\text{Pb}(\text{SO}_4)_2$ in 1 l.

v	c	v
0.304	0.00	0.558
0.348	1.8	0.600
0.387	3.0	0.917
0.407	3.9	1.11
0.435	5.3	1.54
0.477	14.4	2.08
0.515	23.3	2.13

(Dolezalek and Finckli, Z. anorg. 321.)

+ $3\text{H}_2\text{O}$. Deliquescent. Very sol (Marshall.)

Lead potassium persulphate, $\text{K}_2\text{Pb}(\text{SO}_4)_2$.

Decomp. by H_2O . Almost insol. in cold H_2SO_4 of sp. gr. = 1.7. Sl. sol. in H_2SO_4 (sp. gr. = 1.7) at 50° . Sol. in fuming H_2SO_4 and in cold conc. HCl , excess of cold 20% Na acetate + Aq acidified with acetic acid, and in Na acetate + Aq with acetic acid. (Elbs, Z. Elektrochem. 1900, 7. 346.)

sulphate ammonia, $\text{NiS}_2\text{O}_8 \cdot 6\text{NH}_3$.
e in the air. Sol. in H_2O with de-
arbieri, Z. anorg. 1911, 71. 351.)

persulphate, $\text{K}_2\text{S}_2\text{O}_8$.
 H_2O at 0° dissolve 1.77 pts. $\text{K}_2\text{S}_2\text{O}_8$;
in hot H_2O with very sl. decomp.
1 H_2O than any other persulphate.
)

persulphate, $\text{Rb}_2\text{S}_2\text{O}_8$.
 I_2O . 3.32–3.49 pts. are sol. in 100
at 22.5° . (E. F. Smith, J. Am.
. 1899, 21. 934.)

ulphate, basic, $5\text{Ag}_2\text{O}_3$, $2\text{Ag}_2\text{SO}_7$.
by H_2O and acids. (Mulder, C. C.
.)

rsulphate, $\text{Na}_2\text{S}_2\text{O}_8$.
in H_2O . (Löwenherz.)

persulphate.
in H_2O . (Marshall, J. Soc. Chem.
16. 396.)

ersulphate, $\text{Tl}_2\text{S}_2\text{O}_8$.
in H_2O . (Smith, J. Am. Chem.
21. 936.)

phate ammonia, $\text{ZnS}_2\text{O}_8 \cdot 4\text{NH}_3$.
 O . (Barbieri, Z. anorg. 1911, 71.)

molybdic acid.
alphomolybdic acid.

c acid, $\text{HTaO}_4 + n\text{H}_2\text{O}$.
Melikoff, Z. anorg. 1899, 20. 345.)

rtantalate, Cs_2TaO_8 .
E. F. Smith, J. Am. Chem. Soc.
1867.)

assium pertantalate, $\text{CaKTaO}_8 +$
).
cold H_2O ; decomp. by hot H_2O .
Z. anorg. 1899, 20. 347.)

dium pertantalate, $\text{CaNaTaO}_8 +$
).
sol. in H_2O . (E. F. Smith, J.
Soc. 1908, 30. 1668.)

potassium pertantalate,
 $\text{aO}_8 + 7\text{H}_2\text{O}$.
t sol. in H_2O . (E. F. Smith.)

rubidium pertantalate,
 $\text{TaO}_8 + 9\text{H}_2\text{O}$.
t sol. in H_2O . (E. F. Smith.)

Magnesium sodium pertantalate, MgNaTaO_8
 $+ 8\text{H}_2\text{O}$.
Somewhat sol. in H_2O . (E. F. Smith.)

Potassium pertantalate, $\text{K}_2\text{TaO}_8 + \frac{1}{2}\text{H}_2\text{O}$.
Sol. in H_2O with decomp.; sol. in $\text{KOH} +$
 $\text{H}_2\text{O}_2 + \text{Aq}$; pptd. by alcohol. (Melikoff, Z.
anorg. 1899, 20. 346.)

Rubidium pertantalate, Rb_2TaO_8 .
Somewhat sol. in H_2O . (E. F. Smith.)

Sodium pertantalate, $\text{Na}_2\text{TaO}_8 + \text{H}_2\text{O}$.
Pptd. by alcohol. Sl. sol. in H_2O , decomp.
on heating with H_2O . (Melikoff, Z. anorg.
1899, 20. 348.)
 $\text{NaTaO}_4 + \text{NaOTaO}_4 + 13\text{H}_2\text{O}$. Sol. in
 $\text{H}_2\text{O}_2 + \text{Aq}$; pptd. by alcohol. (Melikoff, Z.
anorg. 1899, 20. 349.)

Pertitanic acid.

Ammonium pertitanate, $(\text{NH}_4)_2\text{O}_2$, $\text{TiO}_2 +$
 H_2O_2 .
Fairly stable; decomp. rapidly in aq. solu-
tion. (Melikoff, B. 1898, 31. 955.)

Barium pertitanate, BaO_2 , $\text{TiO}_2 + 5\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Melikoff and Pissarjew-
sky, Z. anorg. 1898, 18. 59.)

Potassium pertitanate, K_2O_2 , TiO_2 , $\text{K}_2\text{O}_4 +$
 $10\text{H}_2\text{O}$.
Stable at zero; deliquesces and decomp. at
ordinary temp. (Melikoff, B. 1898, 31. 680.)

Sodium pertitanate, Na_2O_2 , $\text{TiO}_2 + 3\text{H}_2\text{O}$.
Sol. in H_2O . Pptd. in alcohol. (Melikoff,
B. 1898, 31. 955.)
 $4\text{Na}_2\text{O}_2$, $\text{Ti}_2\text{O}_7 + 10\text{H}_2\text{O}$. Decomp. by H_2O .
(Melikoff.)

Pertungstic acid.

Barium pertungstate, BaO , 2WO_3 , $\text{O} + 6\text{H}_2\text{O}$.
Insol. in H_2O . Decomp. by acids. (Kell-
ner, Dissert. 1909.)

Cæsium pertungstate, $3\text{Cs}_2\text{O}$, 12WO_3 , $2\text{O} +$
 $12\text{H}_2\text{O}$.
Sl. sol. in cold, easily sol. in warm H_2O .
(Kellner.)
 $5\text{Cs}_2\text{O}$, 12WO_3 , $24\text{O} + 11\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Kellner.)

Calcium pertungstate, 3CaO , 6WO_3 , $8\text{O} +$
 $8\text{H}_2\text{O}$.
Sol. in H_2O . (Kellner.)

Lithium pertungstate, Li_2O , 2WO_3 , $2\text{O} +$
 $6\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Kellner.)
 $3\text{Li}_2\text{O}$, 4WO_3 , $\text{O} + 9\text{H}_2\text{O}$. Sol. in H_2O .
(Kellner.)

Magnesium pertungstate, 2MgO , 4WO_3 , $6\text{O} + 9\text{H}_2\text{O}$.

Easily sol. in H_2O . (Kellner.)

Potassium pertungstate, K_2O_4 , $\text{WO}_4 + \text{H}_2\text{O}$.

Sol. in H_2O with decomp.; explodes in the air at 80° . (Melikoff, B. 1898, 31. 634.)

K_2O , 2WO_3 , $4\text{O} + 4\text{H}_2\text{O}$. Sol. in H_2O from which it is ppt. by alcohol and ether. (Kellner.)

$7\text{K}_2\text{O}$, 10WO_3 , $5\text{O} + 22\text{H}_2\text{O}$. Very sl. sol. in H_2O . (Kellner.)

Rubidium pertungstate, $2\text{Rb}_2\text{O}$, 4WO_3 , $\text{O} + 3\text{H}_2\text{O}$.

Sol. in H_2O with slow decomp. (Kellner.)

$5\text{Rb}_2\text{O}$, 12WO_3 , $3\text{O} + 12\text{H}_2\text{O}$. Insol. in H_2O . (Kellner.)

Sodium pertungstate, $\text{NaWO}_4 + \text{H}_2\text{O}$.

Very sol. in H_2O . (Péchar, C. R. 112. 1060.)

$+2\text{H}_2\text{O}$. Sol. in H_2O but easily decomp. (Kellner.)

$\text{Na}_2\text{W}_2\text{O}_7 + 6\text{H}_2\text{O}$. Sol. in H_2O and can be cryst. therefrom. (Pissarjewsky, Z. anorg. 1900, 24. 113.)

Na_2O_2 , $\text{WO}_4 + \text{H}_2\text{O}_2$, $(\text{Na}_2\text{O}_2)\text{WO}_4 + 7\text{H}_2\text{O}$. Decomp. in the air. Sol. in H_2O with decomp. (Melikoff, B. 1898, 31. 633.)

Na_2O_2 , WO_4 , H_2O_2 . Very unstable. Decomp. in the air and by H_2O . (Melikoff.)

Strontium pertungstate, SrO , 2WO_3 , $\text{O} + 6\text{H}_2\text{O}$.

(Kellner.)

Peruranic acid, $\text{UO}_4 \cdot x\text{H}_2\text{O}$ (?).

Known only in its salts.

Ammonium peruranate, $(\text{NH}_4)_2\text{O}_2$, $(\text{UO}_4)_2 + 8\text{H}_2\text{O}$.

Sol. in H_2O ; decomp. by acids and by $\text{Al}(\text{OH})_3$ in aq. solution. (Melikoff, B. 1897, 30. 2904.)

Ammonium uranyl peruranate, $(\text{NH}_4)_2(\text{UO}_2)\text{UO}_4 + 8\text{H}_2\text{O}$ (?).

Easily sol. in H_2O . (Fairley, Chem. Soc. (2) 31. 134.)

Barium peruranate, BaUO_4 .

As K salt. (de Coninck, C. C. 1909, I. 1970.)

$(\text{BaO}_2)_2\text{UO}_4 + 8\text{H}_2\text{O}$. Decomp. by H_2SO_4 and H_2CO_3 . (Melikoff, B. 1897, 30. 2905.)

$\text{BaO}_2(\text{UO}_4)_2 + 9\text{H}_2\text{O}$. Ppt. (Melikoff.)

Calcium peruranate, CaUO_4 .

As K salt. (de Coninck.)

Calcium peruranate, $(\text{CaO}_2)_2\text{UO}_4 + 10\text{H}_2\text{O}$.

Ppt. (Melikoff, B. 1897, 30. 2906.)

Copper peruranate, $(\text{CuO}_2)_2\text{UO}_4$.

Ppt. (Melikoff.)

Lead peruranate, $(\text{PbO})_2\text{UO}_4$, PbO .

Ppt. (Melikoff.)

Lithium peruranate, $(\text{Li}_2\text{O}_2)(\text{UO}_4)_2$.

Sol. in H_2O ; decomp. by acids $\text{Al}(\text{OH})_3$ in aq. solution; very (Melikoff.)

Nickel peruranate, $(\text{NiO})_2\text{UO}_4$.

Ppt. (Melikoff.)

Potassium peruranate, $\text{K}_4\text{UO}_8 + 10\text{H}_2\text{O}$.

Unstable. (Fairley.)

K_2UO_6 . (de Coninck, C. R. 1769.)

$+3\text{H}_2\text{O}$. Decomp. by H_2O , HCl , HNO_3 . (Aloy, Bull. Soc. 1903, (3)

Silver peruranate, $\text{Ag}_2\text{U}_2\text{O}_{11}$ (?).

(Guyard, Bull. Soc. (2) 1. 95.)

Does not exist. (Alibegoff, A. 23)

Sodium peruranate, $\text{Na}_4\text{UO}_8 + 5\text{H}_2\text{O}$.

Sol. in H_2O . Sl. sol. in alcohol.

Na_2UO_6 . As K salt. (de Coninck, 1909, I. 1970.)

$+5\text{H}_2\text{O}$. Decomp. by H_2O (Aloy, Bull. Soc. 1903, (3) 29. 293.)

$(\text{Na}_2\text{O}_2)_2\text{UO}_4 + 8\text{H}_2\text{O}$. Sol. in comp. by dil. HCl , H_2SO_4 , and by in aq. solution. (Melikoff, B. 1897,

Sodium uranyl peruranate, $\text{Na}_2(\text{UO}_2)_2\text{UO}_4 + 6\text{H}_2\text{O}$ (?).

Sol. in H_2O . (Fairley.)

Pervanadic acid, HVO_4 (?).

Sol. in H_2O . (Pissarjewsky, C. C. 565.)

Ammonium pervanadate, NH_4VO_4 .

Sol. in $\text{H}_2\text{O}_2 + \text{Aq}$; insol. in alcohol. Z. anorg. 1898, 16. 294.)

$(\text{NH}_4)_2\text{VO}_6 + 2\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O in alcohol. (Melikoff, B. 1909, 42.)

$(\text{NH}_4)_4\text{V}_2\text{O}_{11}$. Sol. in $\text{H}_2\text{O}_2 + \text{Aq}$; aq. solution by alcohol. (Melikoff, 1899, 19. 406.)

Barium pervanadate, $\text{Ba}(\text{VO}_4)_2$.

Sl. sol. in $\text{H}_2\text{O}_2 + \text{Aq}$ free from insol. in alcohol. (Scheuer, Z. anorg. 16. 288.)

Cadmium pervanadate, $\text{Cd}(\text{VO}_4)_2$.

Sl. sol. in $\text{H}_2\text{O}_2 + \text{Aq}$; insol. in alcohol. (Scheuer.)

Calcium pervanadate, $\text{Ca}(\text{VO}_4)_2$.

Sol. in $\text{H}_2\text{O}_2 + \text{Aq}$; insol. in alcohol. (Scheuer.)

vanadate, $\text{Pb}(\text{VO}_4)_2$.
 in $\text{H}_2\text{O}_2 + \text{Aq}$ free from H_2SO_4 ; insol.
 (Scheuer.)

ervanadate, LiVO_4 .
 $\text{H}_2\text{O}_2 + \text{Aq}$; insol. in alcohol.

pervanadate, KVO_4 .
 $\text{H}_2\text{O}_2 + \text{Aq}$ acidified with H_2SO_4 ;
 alcohol. (Scheuer.)
 $+2\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O ; insol. in
 (Melikoff, B. 1909, 42. 2293.)
 O_4 , $2\text{KVO}_4 + 2\text{H}_2\text{O}$. 0.855 g. is
 1 g. H_2O at 19° ; sl. sol. in $\text{KOH} +$
 stable in the air. (Melikoff and
 sky, Z. anorg. 1899, 19. 408.)
 $+2\text{H}_2\text{O}$. (Melikoff and Pissar-
 anorg. 1899, 19. 411.)
 $+3\frac{1}{2}\text{H}_2\text{O}$. Moderately sol. in
 slow decomp. (Melikoff and Pissar-
 anorg. 1899, 19. 410.)

vanadate, AgVO_4 .
 in $\text{H}_2\text{O}_2 + \text{Aq}$ free from H_2SO_4 ; sol.
 (Scheuer.)

ervanadate, NaVO_4 .
 H_2O_2 acidified with H_2SO_4 ; insol.
 (Scheuer.)

pervanadate, $\text{Sr}(\text{VO}_4)_2$.
 in $\text{H}_2\text{O}_2 + \text{Aq}$ free from H_2SO_4 ;
 alcohol. (Scheuer.)

um, Ph (?).
 ontaine, C. R. 87. 559.)
 s of terbium and yttrium. (Roscoe,
 74.)

e, CoCl_2 .
bonyl chloride.

me, PN_2H (?).
 in H_2O . Insol. in dil. $\text{HNO}_3 + \text{Aq}$;
 decomp. by conc. HNO_3 . (Rose,
 308.)
 in conc. HNO_3 . (Pauli, A. 123. 236.)
 H_2SO_4 with decomp. (Rose.)
 in dil., but decomp. by conc. KOH
 $+ \text{Aq}$.
 in alcohol or ether.
 la is perhaps $\text{P}_3\text{N}_3\text{H}_4$. (Salzmann,
 .)

mic acid, $\text{PO} \begin{smallmatrix} \diagup \text{NH} \\ \diagdown \text{OH} \end{smallmatrix}$

)
 not exist, but was impure *pyrophos-*
c acid. (Gladstone.) Also Mente
 245).

**Pyrophosphamic acid, $\text{P}_2\text{NH}_4\text{O}_6 =$
 $\text{P}_2\text{O}_5(\text{OH})_2\text{NH}_2$.**

Deliquescent in moist air; easily sol. in
 H_2O or alcohol; sl. sol. in ether. (Gladstone,
 Chem. Soc. 3. 152.)

Correct composition is imidodiphosphoric
 acid, $\text{P}_2\text{NH}_4\text{O}_6 = \text{HO}-\text{PO} < \begin{smallmatrix} \text{O} \\ \text{NH} \end{smallmatrix} > \text{PO}-\text{OH}$.
 (Mente, A. 248. 232.)

Barium pyrophosphamate, $\text{Ba}_2(\text{P}_2\text{NH}_4\text{O}_6)_2$.

Sol. in HCl or $\text{HNO}_3 + \text{Aq}$, not in $\text{HC}_2\text{H}_3\text{O}_2$
 $+ \text{Aq}$. (Gladstone and Holmes, Chem. Soc.
 (2) 2. 233.)

Cupric —, $\text{Cu}_2(\text{P}_2\text{NH}_4\text{O}_6)_2 + 2\text{H}_2\text{O}$.

Ppt. Decomp. by cold $\text{KOH} + \text{Aq}$. (Glad-
 stone, Chem. Soc. 3. 135.)

Ferric —, $\text{Fe}_2(\text{P}_2\text{NH}_4\text{O}_6)_2 + 2\text{H}_2\text{O}$.

Insol. in dil. acids. Sol. in conc. H_2SO_4 ,
 and decomp. by warming. Easily sol. in
 $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$.
 (Gladstone, Chem. Soc. 3. 142.)

Lead —, $\text{Pb}_2(\text{P}_2\text{NH}_4\text{O}_6)_2 + 4\text{H}_2\text{O}$.

Insol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Potassium —, $\text{K}_2\text{P}_2\text{NH}_4\text{O}_6$.

Deliquescent. Sol. in H_2O . Insol. in
 alcohol. (Gladstone, A. 76. 85.)

Silver —, $\text{Ag}_2\text{P}_2\text{NH}_4\text{O}_6 + 5\text{H}_2\text{O}$.

Ppt.

Zinc —, $\text{Zn}_2(\text{P}_2\text{NH}_4\text{O}_6)_2$.

(Gladstone and Holmes, Chem. Soc. (2) 2.
 225.)

Phosphamide, PON .

See Phosphoryl nitride.

$\text{PN}_2\text{H}_3\text{O}$.

See Phosphoryl imidoamide.

Triphosphamide, PON_3H_4 .

See Phosphoryl triamide.

Trimetaphosphimic acid, $\text{P}_3\text{N}_3\text{H}_4\text{O}_6$.

Sol. in H_2O ; aq. solution does not coagulate
 albumen. (Stokes, Am. Ch. J. 1895, 17. 275.)

Ammonium trimetaphosphimate,

$(\text{NH}_4)_3\text{P}_3\text{N}_3\text{O}_6\text{H}_3$.

Sol. in H_2O , insol. in alcohol; unstable.
 (Stokes, Am. Ch. J. 1896, 18. 643.)

**Barium trimetaphosphimate, $\text{Ba}_3(\text{P}_3\text{N}_3\text{O}_6\text{H}_3)_2$
 $+ 4\text{H}_2\text{O}$.**

$+ 6\text{H}_2\text{O}$. Sl. sol. in H_2O . Easily sol. in
 $\text{NH}_4\text{Cl} + \text{Aq}$ and in $\text{NaCl} + \text{Aq}$. (Stokes.)

Barium sodium trimetaphosphimate,
 $P_3N_3O_6H_3NaBa + 1\frac{1}{2}H_2O$.

Almost insol. in H_2O ; easily sol. in $NH_4Cl + Aq$ and in $NaCl + Aq$. (Stokes.)

Magnesium trimetaphosphimate,
 $(P_3N_3O_6H_3)_2Mg_3 (?)$.

Sol. in H_2O ; insol. in alcohol; aq. solution decomp. on boiling. (Stokes.)

Potassium trimetaphosphimate, $K_3P_3N_3O_6H_3$.
 Sol. in H_2O ; insol. in alcohol. (Stokes.)

Silver trimetaphosphimate, $Ag_3P_3N_3O_6H_3$.

Ppt.; sol. in $NH_4OH + Aq$; insol. in H_2O ; sl. sol. in $HNO_3 + Aq$. (Stokes.)

α **Sodium trimetaphosphimate,** $P_3N_3O_6H_3Na_3 + 4H_2O$.

18.3 pts. are sol. in 100 pts. H_2O at 20° ; very sol. in hot H_2O ; decomp. by alkali on long boiling. (Stokes.)

β **Sodium trimetaphosphimate,** $P_3N_3O_6H_3Na_3 + H_2O$.

Sol. in H_2O ; insol. in alcohol. (Stokes.)

Tetrametaphosphimic acid,
 $P_4N_4O_8H_8 + 2H_2O$.

Very sl. sol. in H_2O ; decomposes the sol. salts of HCl , H_2SO_4 and HNO_3 . (Stokes, Am. Ch. J. 1895, 17. 290.)

100 pts. H_2O at 20° dissolve 0.64 pt. crystallized acid. Somewhat more sol. in boiling H_2O . (Stokes.)

100 pts. 10% $HNO_3 + Aq$ at 20° dissolve 0.26 pt. of crystallized acid. (Stokes.)

Not decomp. by boiling alkalies + Aq . (Stokes, Am. Ch. J. 1896, 18. 785.)

Insol. in alcohol. (Stokes, Am. Ch. J. 1896, 18. 784.)

Ammonium tetrametaphosphimate,
 $P_4N_4O_8H_8(NH_4)_2$.

Only sl. sol. in boiling H_2O ; sol. in excess of hot 5% HNO_3 . (Stokes.)

$P_4N_4O_8H_8(NH_4)_4 + 4H_2O$. Readily sol. in H_2O ; sl. sol. in $NH_4OH + Aq$. (Stokes.)

Barium tetrametaphosphimate, $P_4N_4O_8H_8Ba_2 + 2H_2O$.

Ppt.; insol. in H_2O . (Stokes.)

Potassium tetrametaphosphimate,
 $P_4N_4O_8H_8K_2$.

Sl. sol. in boiling H_2O ; sol. in cold dil. $KOH + Aq$. (Stokes.)

$P_4N_4O_8H_8K_4 (?)$. Very sol. in H_2O . (Stokes.)

Silver tetrametaphosphimate, $P_4N_4O_8H_8Ag_4$.

Ppt.; insol. in H_2O ; sl. sol. in $HNO_3 + Aq$. (Stokes.)

$P_4N_4O_8Ag_3$. Ppt.; sol. in NH_4NO_3 (Stokes.)

Sodium tetrametaphosphimate, $P_4N_4O_8 + 2\frac{1}{2} (?) H_2O$.

Sl. sol. in cold H_2O . Easily sol. in h. Ppt. from aqueous solution by excess kali. (Stokes.)

Pentametaphosphimic acid,

$PO(OH) \begin{matrix} \diagup NH.PO(OH)NH.PO(OH) \\ \diagdown NH.PO(OH)NH.PO(OH) \end{matrix}$
 $= H_{10}P_5N_5O_{10}$.

Sol. in H_2O ; pptd. by alcohol. (Stokes, Ch. J. 1898, 20. 748.)

Magnesium pentametaphosphimate,
 $(P_5N_5O_{10}H_5)_2Mg$.

Ppt. (Stokes.)

$P_5N_5O_{10}H_5Mg_2 + 5H_2O$. Ppt., insol. in alcohol; almost insol. in H_2O ; sl. sol. in acetic acid. (Stokes.)

Silver pentametaphosphimate, $P_5N_5O_{10}H_5$.

Ppt., sol. in cold $KOH + Aq$ with (Stokes.)

Sodium pentametaphosphimate,
 $P_5N_5O_{10}H_5Na_3 + 2H_2O$.

Sol. in H_2O ; insol. in alcohol. (Stokes, Am. Ch. J. 1896, 18. 784.)
 $P_5N_5O_{10}H_5Na_4 + 2H_2O$. Sol. in 80% acid; pptd. by alcohol. (Stokes.)

Hexametaphosphimic acid.

Silver hexametaphosphimate, $P_6N_6O_{12}H_6$.

Ppt.; decomp. by cold $KOH + Aq$. (Stokes, Am. Ch. J. 1898, 20. 757.)

Sodium hexametaphosphimate, $P_6N_6O_{12}H_6 + 2H_2O$.

Sol. in H_2O ; pptd. by alcohol. (Stokes.)

Phosphine.

See Hydrogen phosphide.

Pyrophosphodiamic acid,

$P_2N_2H_4O_4 = P_2O_2(OH)_2(NH_2)_2$.

Deliquescent. Easily sol. in H_2O , or ether. Sol. in cold conc. H_2SO_4 ; decomp. (Gladstone, Chem. Soc. 2.)

Correct composition is dimidodiphosphic acid, $P_2N_2H_4O_4 + H_2O = HO-PO=PO-OH$. (Mente.)

Aluminum pyrophosphodiamate.

Precipitate. Sol. in $NH_4OH + Aq$ in acids. (Gladstone, A. 76. 82.)

Ammonium —, $P_2O_2(ONH_2)_2(NH_4)_2$.

Very deliquescent in moist air. Sol. (Schiff, A. 103. 168.)

Barium pyrophosphodiamate,
 $\text{BaP}_2\text{O}_6(\text{NH}_2)_2$.

Precipitate. Sl. sol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Gladstone.)

Calcium —, $\text{CaP}_2\text{O}_6(\text{NH}_2)_2$.

Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Sol. in $\text{NH}_4\text{Cl} + \text{Aq.}$ and acids. (Gladstone and Holmes.)

Lead —.

Ppt. Decomp. by H_2O .

Magnesium —.

Ppt. (Gladstone and Holmes.)

Silver —, $\text{Ag}_2\text{P}_2\text{O}_6(\text{NH}_2)_2$.

Sl. sol. in H_2O . Sol. in $\text{HNO}_3 + \text{Aq.}$ (Gladstone and Holmes.)

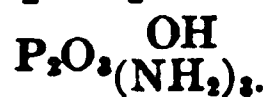
Strontium —.

Sol. in acids and $\text{NH}_4\text{Cl} + \text{Aq.}$ Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Gladstone and Holmes, Chem. Soc. (2) 4. 295.)

Zinc —, $\text{ZnP}_2\text{O}_6(\text{NH}_2)_2$.

Ppt. (Gladstone and Holmes.)

Pyrophosphotriamic acid, $\text{P}_2\text{N}_2\text{H}_7\text{O}_4 =$



Decomp. by boiling H_2O or HCl . Sol. in conc. H_2SO_4 upon heating. (Gladstone and Holmes.)

Correct formula is $\text{HO}-\text{PO} < \begin{matrix} \text{NH} \\ \text{NH} \end{matrix} > \text{PO}-\text{NH}_2 = \text{diimidodiphosphomonamic acid}$. (Mente, A. 248. 241.)

Ammonium pyrophosphotriamate,



Insol. in H_2O . (Gladstone and Holmes.)

Barium —, $\text{BaP}_2\text{N}_2\text{H}_5\text{O}_4$.

$\text{BaH}_2(\text{P}_2\text{N}_2\text{H}_5\text{O}_4)_2$. Decomp. by $\text{HCl} + \text{Aq.}$ (Gladstone, Chem. Soc. 4. 6.)

Cobaltous —, $\text{CoP}_2\text{N}_2\text{H}_5\text{O}_4$.

Slowly decomp. by dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ not by $\text{HCl} + \text{Aq.}$ (Gladstone and Holmes, Chem. Soc. (2) 4. 1.)

Cupric —, $\text{CuP}_2\text{N}_2\text{H}_5\text{O}_4$.

Insol. in H_2O or $\text{NH}_4\text{OH} + \text{Aq.}$ (Gladstone and Holmes, Chem. Soc. (2) 4. 1.)

Ferrous —, $\text{FeH}_2(\text{P}_2\text{N}_2\text{H}_5\text{O}_4)_2$.

Insol. in dil. acids. (Gladstone, Chem. Soc. (2) 4. 1.)

Lead —, $\text{H}_2\text{Pb}_2(\text{P}_2\text{N}_2\text{H}_5\text{O}_4)_2$.

Ppt. (Gladstone and Holmes, Chem. Soc. (2) 4. 1.)

$\text{H}_4\text{Pb}_2(\text{P}_2\text{N}_2\text{H}_5\text{O}_4)_2$. Ppt. (G. and H.)

$\text{H}_4\text{Pb}(\text{P}_2\text{N}_2\text{H}_5\text{O}_4)_2$. (G. and H.)

Mercuric pyrophosphotriamate,



Insol. in H_2O or dil. HCl or $\text{HNO}_3 + \text{Aq.}$ (Gladstone and Holmes, Chem. Soc. (2) 4. 1.)

Platinum —, $\text{Pt}_2\text{P}_2\text{N}_2\text{H}_5\text{O}_4$.

Decomp. by H_2O when freshly pptd. (G. and H.)

Potassium —, $\text{KP}_2\text{N}_2\text{H}_5\text{O}_4$.

Almost insol. in H_2O . (Gladstone, Chem. Soc. 4. 10.)

Silver —, $\text{Ag}_2\text{P}_2\text{N}_2\text{H}_5\text{O}_4$.

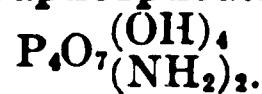
Ppt. Sl. attacked by $\text{HC}_2\text{H}_3\text{O}_2$; decomp. by HNO_3 or $\text{NH}_4\text{OH} + \text{Aq}$ into—

$\text{AgH}_2\text{P}_2\text{N}_2\text{H}_5\text{O}_4$. Insol. in H_2O . Decomp. by HCl . (Gladstone, Chem. Soc. (2) 4. 1.)

Zinc —.

Insol. in H_2O . (Gladstone and Holmes.)

Tetraphosphodiamic acid, $\text{P}_4\text{N}_2\text{H}_5\text{O}_{11} =$



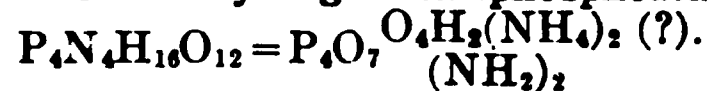
Known only as NH_4 salt.

Ammonium tetraphosphodiamate,



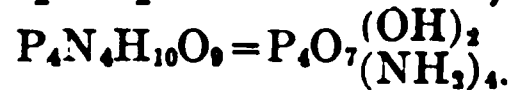
Very deliquescent, and sol. in H_2O . (Gladstone.)

Ammonium dihydrogen tetraphosphodiamate,



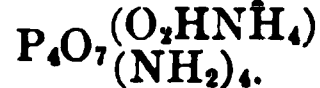
Insol. in cold, easily sol. in hot H_2O and dil. acids. (Gladstone.)

Tetraphosphotetramic acid,



Sol. in H_2O . Insol. in alcohol. (Gladstone.)

Ammonium tetraphosphotetramate,



Sol. in H_2O , and precipitated from solution by alcohol. (Gladstone.)

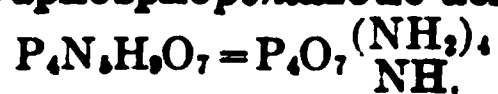
Silver —, $\text{Ag}_2\text{P}_4\text{N}_4\text{H}_4\text{O}_9$.

Ppt.

$\text{Ag}_2\text{H}_4\text{P}_4\text{N}_4\text{H}_4\text{O}_9$. Ppt.

**Ammonium phosphoarseniovanadico-
vanadiotungstate.**

See **Arseniophosphovanadico-
vanadiotungstate, ammonium.**

Tetraphosphopentazotic acid,

Insol. in H_2O . Decomp. gradually by boiling with H_2O . (Gladstone.)

Ammoniotetraphosphopentazotic acid (?),

Decomp. by H_2O . (Gladstone.)

Cupric tetraphosphopentazotate.

(Gladstone, Chem. Soc. (2) 6. 261.)

Lead —.

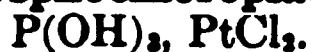
(Gladstone, Chem. Soc. (2) 6. 261.)

Potassium —, $KOP_4N_5H_5O_6$.

Insol. in H_2O . (Gladstone, Chem. Soc. (2) 6. 268.)

Phosphoboric acid, H_3BO_3 , $H_3PO_4 = BPO_4 + 3H_2O$.

Not decomp. by boiling H_2O or conc. acids. Sol. in boiling solution of caustic alkalies. (Vogel, N. Repert. Pharm. 18. 611.)

Phosphochloroplatinous acid,

See Chloroplatinophosphoric acid.

Phosphochromic acid.**Ammonium phosphochromate, $3(NH_4)_2O$, P_2O_5 , $8CrO_3 + H_2O$.**

Sol. in H_2O with decomp. (Friedheim, Z. anorg. 1894, 6. 284.)

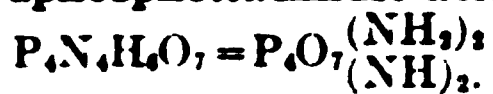
Potassium phosphochromate, $2K_2O$, P_2O_5 , $4CrO_3 + H_2O$.

Sol. in H_2O but cannot be cryst. therefrom without decomp. Can be cryst. without decomp. from H_2O containing phosphoric acid. (Friedheim.)

$3K_2O$, P_2O_5 , $8CrO_3$. Sol. in H_2O but cannot be cryst. therefrom without decomp. (Blondel, C. R. 1894, 118. 194.)

Phosphohypophosphotungstic acid.**Potassium sodium phosphohypophosphotungstate, $9K_2O$, Na_2O , $4P_2O_5$, $2PO_2H_2$, $26WO_3 + 23H_2O$.**

Precipitate. Easily sol. in hot H_2O . (Gibbs, Am. Ch. J. 7. 313.)

Tetraphosphotetrimidic acid,

Known only in its salts. (Gladstone.)

Silver tetraphosphotetrimidate.

Ppt. (Gladstone.)

Phosphiodic acid, P_2O_5 , $18I_2O_5 +$

Decomp. by H_2O . (Chrétien, A. (7) 15. 389.)

Ammonium phosphiodate, $4(NH_4)$, $18I_2O_5 + 12H_2O$.

Sol. in H_2O . Sl. sol. in conc. H_2 . (Chrétien.)

Lithium phosphiodate, $3Li_2O$, P_2O_5 , $11H_2O$.

Sol. in H_2O . Sl. sol. in conc. H_2 . (Chrétien.)

Potassium phosphiodate, $4K_2O$, $18I_2O_5 + 5H_2O$.

Decomp. by a small amt. of H_2O large amt. Sl. sol. in conc. H_2 . (Chrétien.)

Sodium phosphiodate, $6Na_2O$, $P_2O_5 + 5H_2O$.

Sol. in H_2O . Sl. sol. in conc. H_2 . (Chrétien.)

Phosphoiridic acid.

See Chlorophosphoiridic acid.

Phospholuteotungstic acid, H_3PV

See under Phosphotungstic acid.

Phosphomolybdic acid, P_2O_5 , $18I_2O_5 + xH_2O$.

"Phospholuteomolybdic acid."

Deliquescent. Sol. in H_2O in all proportions. (Kehrmann, Z. anorg. 1894, 3H₂O, P_2O_5 , $20MoO_3 + 21H_2O$. in H_2O . Sol. in ether. By evapn H_2O solution crystals with $44H_2O$ a strong solution in conc. HNO_3 + $19H_2O$, are obtained; also crystals and $48H_2O$ are known. (Debray, 704.)

According to Rammelsberg (B. formula is $3H_2O$, P_2O_5 , $22MoO_3$.

According to Gibbs (Am. Ch. J. formula is $3H_2O$, P_2O_5 , $24MoO_3 + 5H_2O$.

Finkener (B. 11. 1638) gives the formula $3H_2O$, P_2O_5 , $24MoO_3 + 58H_2O$, + $29H_2O$.

P_2O_5 , $20MoO_3 + 52H_2O$. Sol. in H_2O with evolution of heat, and subsequent addition into two layers, the upper contains pure ether, and lower of a solution of ether. Sp. gr. of lower layer, when 13°, is 1.3. On warming lower layer separates out and forms an upper layer which redissolves on cooling and shaking. The lower layer is insol. in H_2O and miscible with alcohol.

ether thus dissolve 80.6 pts. acid
pts. at 8.1°; 96.7 pts. at 19.3°;
at 27.4°; 107.9 pts. at 32.9°. (Par-
R. 104. 688.)

$\text{MoO}_3 + 57\text{H}_2\text{O}$, and $+58\text{H}_2\text{O}$.
ert. 1906.)

$\text{MoO}_3 + 61\text{H}_2\text{O}$. (Pohl.)

$\text{MoO}_3 + 61\text{H}_2\text{O}$. (Miolati, C. C.
39.)

. (Pohl.)

pentamolybdic acid,

$10\text{O}_3 = 3\text{H}_2\text{O}$, P_2O_5 , 5MoO_3 .

own in free state.

phosphomolybdate, $(\text{NH}_4)_2\text{O}$,
 $2\text{MoO}_3 + 2\text{H}_2\text{O}$.

rier, Dissert. 1894.)

O , P_2O_5 , $4\text{MoO}_3 + 5\text{H}_2\text{O}$. (Fried-
urg. 1894, 6. 33.)

(Perlberger, Dissert. 1904.)

O_4 , $11\text{MoO}_3 + 6\text{H}_2\text{O}$.

is $(\text{NH}_4)_2\text{PO}_4$, $10\text{MoO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$,
o the older authorities.

sol. in H_2O or aqueous acid solu-
ily sol. in ammonia, and alkalies +
berg and Struve, J. pr. 44. 291.)

not completely insol. in a mixture
 $\text{MoO}_3 + \text{Aq}$, and dil. $\text{HNO}_3 + \text{Aq}$
insol. in a dil. nitric acid solution
in nitrate. (Richters, Z anal. 10.

is increased even in presence of
molybdate and free HNO_3 by
onium, and other chlorides, tar-
or large quantities of ammonium
nitrate. Not precipitated in pres-
ence of H_3PO_4 . (Fresenius, Z. anal.

1,000 pts. H_2O at 16°; in 6600 pts.
ning 1 vol. % HNO_3 ; in 550 pts.
f 1.12 sp. gr.; in 620 pts. alcohol
gr.; in 190 pts. $\text{HNO}_3 + \text{Aq}$ (sp.
50°; in 5 pts. conc. H_2SO_4 at 100°;
 $\text{LiOH} + \text{Aq}$ of 0.95 sp. gr. (Eggertz,
96.)

1,186 pts. H_2O , 38,117 pts. dil.
id 13,513 pts. strong alcohol
nalyst, 1879. 23.)

g to Sonnenschein, the solubility is
y much H_2O or alcohol, alkaline
carbonates, ortho-, pyro-, and
ates; sodium borate, hyposul-
sulphate, acetate, arsenate, and
tassium sodium tartrate, ammo-
ze, orthophosphoric acid, and sul-
. It is not increased by ammo-
xide or sulphate, potassium sul-
tartrate, acid oxalate, nitrate, or
lide, chloride, or bromide, sodium
nitrate; nitric, hydrochloric, boric,
alic, and dilute sulphuric acids.
in, J. pr. 53. 342.)

not H_2O . Sol. in cold caustic
ali carbonates, and phosphates,

NH_4Cl , and $(\text{NH}_4)_2\text{C}_2\text{O}_4 + \text{Aq}$; al. sol. in
 $(\text{NH}_4)_2\text{SO}_4$, KNO_3 , and $\text{KCl} + \text{Aq}$; very al. sol.
in $\text{NH}_4\text{NO}_3 + \text{Aq}$. Sol. in K_2SO_4 , Na_2SO_4 ,
 NaCl , MgCl_2 , H_2SO_4 , HCl , and conc. or dil.
 $\text{HNO}_3 + \text{Aq}$.

Presence of $(\text{NH}_4)_2\text{MoO}_4$ totally changes
the effect of acid liquids; insol. in dil. HNO_3 or
 $\text{H}_2\text{SO}_4 + \text{Aq}$ containing $(\text{NH}_4)_2\text{MoO}_4$, but
somewhat sol. in $\text{HCl} + \text{Aq}$, even in presence
of that salt. Tartaric acid and similar organic
substances totally prevent the precipitation
of this salt. (Eggertz in Fresenius' Quant.
anal.)

$5(\text{NH}_4)_2\text{O}$, 48MoO_3 , $2\text{P}_2\text{O}_5 + 17\text{H}_2\text{O} =$
 $3(\text{NH}_4)_2\text{O}$, 24MoO_3 , $\text{P}_2\text{O}_5 + 2(\text{NH}_4)_2\text{O}$, H_2O ,
 24MoO_3 , $\text{P}_2\text{O}_5 + 16\text{H}_2\text{O}$. Formula of above
salt according to Gibbs.

$3(\text{NH}_4)_2\text{O}$, 22MoO_3 , $\text{P}_2\text{O}_5 + 9\text{H}_2\text{O}$, or
 $12\text{H}_2\text{O}$.

$8(\text{NH}_4)_2\text{O}$, H_2O , 60MoO_3 , $3\text{P}_2\text{O}_5 + 11\text{H}_2\text{O}$.
Sl. sol. in H_2O .

$3(\text{NH}_4)_2\text{O}$, 16MoO_3 , $\text{P}_2\text{O}_5 + 14\text{H}_2\text{O}$. Insol.
in cold, sol. with decomp. in hot H_2O . Sol.
in $\text{NH}_4\text{OH} + \text{Aq}$. (Gibbs, Am. Ch. J. 3. 317.)

$5(\text{NH}_4)_2\text{O}$, P_2O_5 , 16MoO_3 . (Meschoirer,
Dissert. 1894.)

$3(\text{NH}_4)_2\text{O}$, P_2O_5 , $18\text{MoO}_3 + 14\text{H}_2\text{O}$. Sol.
in H_2O . The aqueous solution is stable at
ordinary temp. for several days, but when
warmed ordinary ammonium phosphomolyb-
date separates. (Kehrmann, Z. anorg. 1894,
7. 414.)

$3(\text{NH}_4)_2\text{O}$, P_2O_5 , $28\text{MoO}_3 + 8\text{H}_2\text{O}$. 100 g.
 H_2O dissolve 0.0238 g. at 15°. 1 pt. is sol. at
15° in 4206 pts. H_2O ; 7300 pts. 5% $\text{NH}_4\text{NO}_3 +$
 Aq ; 4930 pts. 1% $\text{HNO}_3 + \text{Aq}$. (de Lucchi,
Rass. Min. 1910, 32. 21.)

$9(\text{NH}_4)_2\text{O}$, $2\text{P}_2\text{O}_5$, $28\text{MoO}_3 + 8\text{H}_2\text{O}$. (Mes-
choirer, Dissert. 1894.)

Ammonium diphosphopentamolybdate,

$2(\text{NH}_4)_2\text{PO}_4$, $5\text{MoO}_3 + 7\text{H}_2\text{O} = 3(\text{NH}_4)_2\text{O}$,
 5MoO_3 , $\text{P}_2\text{O}_5 + 7\text{H}_2\text{O}$.

Easily sol. in hot, less in cold H_2O . (Zenk-
ner, J. pr. 58. 256.)

$5(\text{NH}_4)_2\text{O}$, H_2O , 10MoO_3 , $2\text{P}_2\text{O}_5 + 6\text{H}_2\text{O}$
 $= 3(\text{NH}_4)_2\text{O}$, 5MoO_3 , $\text{P}_2\text{O}_5 + 2(\text{NH}_4)_2\text{O}$, H_2O ,
 5MoO_3 , $\text{P}_2\text{O}_5 + 6\text{H}_2\text{O}$. Sol. in H_2O . (Gibbs,
Am. Ch. J. 1895, 17. 87.)

$+8\text{H}_2\text{O}$. (Perlberger.)

$+18\text{H}_2\text{O}$. (Mazzuchelli and Zangrilli,
Gazz. ch. it 1910, 40. (2) 55.)

$5(\text{NH}_4)_2\text{O}$, P_2O_5 , $10\text{MoO}_3 + 13\text{H}_2\text{O}$, and
 $+14\text{H}_2\text{O}$. (Perlberger, Dissert. 1904.)

Ammonium barium phosphomolybdate,

$3(\text{NH}_4)_2\text{O}$, 30BaO , P_2O_5 , 30MoO_3 .

Insol. precipitate. (Seligsohn, J. pr. 67.
478.)

Ammonium cadmium phosphomolybdate,

$5(\text{NH}_4)_2\text{O}$, CdO , P_2O_5 , $6\text{MoO}_3 + 8\text{H}_2\text{O}$.

(Perlberger, Dissert. 1904.)

$3(\text{NH}_4)_2\text{O}$, 2CdO , $2\text{P}_2\text{O}_5$, $9\text{MoO}_3 + 14\text{H}_2\text{O}$.
(Perlberger.)

Ammonium cobaltous phosphomolybdate,
 $(\text{NH}_4)_2\text{O}, 2\text{CoO}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 10\text{H}_2\text{O}.$

Decomp. by cold H_2O . Sol. in acids and hot H_2O . (Arnfeld, Dissert. 1898.)

$4(\text{NH}_4)_2\text{O}, \text{CoO}, 2\text{P}_2\text{O}_5, 10\text{MoO}_3 + 12\text{H}_2\text{O}.$
 Sl. sol. in cold, easily sol. in hot H_2O . (Arnfeld.)

Ammonium manganous phosphomolybdate,
 $(\text{NH}_4)_2\text{O}, 2\text{MnO}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 20\text{H}_2\text{O}.$

Decomp. by H_2O , but dissolves clear on heating. (Arnfeld.)

$4(\text{NH}_4)_2\text{O}, \text{MnO}, 2\text{P}_2\text{O}_5, 10\text{MoO}_3 + 13\text{H}_2\text{O}.$
 (Arnfeld.)

$5(\text{NH}_4)_2\text{O}, 10\text{MnO}, 2\text{P}_2\text{O}_5, 20\text{MoO}_3 + 10\text{H}_2\text{O}.$ Very sl. sol. in H_2O . (Gibbs, Am. Ch. J. 1895, 17. 87.)

Ammonium nickel phosphomolybdate,
 $(\text{NH}_4)_2\text{O}, 2\text{NiO}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 10\text{H}_2\text{O}.$

Decomp. by cold, but sol. in hot H_2O . (Arnfeld.)

$(\text{NH}_4)_2\text{O}, \text{NiO}, 2\text{P}_2\text{O}_5, 10\text{MoO}_3 + 12\text{H}_2\text{O}.$
 Very sol. in H_2O . (Arnfeld.)

Ammonium potassium phosphomolybdate,
 $6(\text{NH}_4)_2\text{O}, 15\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 60\text{MoO}_3 + 12\text{H}_2\text{O}.$

Sol. in H_2O . Insol. in alcohol. (Seligsohn, J. pr. 67. 477.)

Ammonium sodium phosphomolybdate,
 $6(\text{NH}_4)_2\text{O}, 15\text{Na}_2\text{O}, 2\text{P}_2\text{O}_5, 60\text{MoO}_3 + 18\text{H}_2\text{O}.$

Sol. in much boiling H_2O . Insol. in alcohol. (Seligsohn, J. pr. 67. 474.)

Barium phosphomolybdate, $3\text{BaO}, \text{P}_2\text{O}_5, 24\text{MoO}_3 + x\text{H}_2\text{O}.$

Moderately sol. in cold, very easily sol. in hot H_2O . Decomp. in aqueous solution at ordinary temp. on standing. (Kehrmann, Z. anorg. 1894, 7. 414.)

Cæsium phosphomolybdate, $3\text{Cs}_2\text{O}, \text{P}_2\text{O}_5, 6\text{MoO}_3 + 8\text{H}_2\text{O}.$

Ppt. (Ephraim, Z. anorg. 1910, 65. 240.)
 $2\text{Cs}_2\text{O}, \text{P}_2\text{O}_5, 14\text{MoO}_3 + 3\text{H}_2\text{O}.$ Difficultly sol. in H_2O . (Ephraim.)

$3\text{Cs}_2\text{O}, \text{P}_2\text{O}_5, 21\text{MoO}_3 + 4\text{H}_2\text{O} (?)$. (Ephraim.)

Calcium potassium phosphomolybdate, $2\text{CaO}, 3\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 10\text{MoO}_3 + 22\text{H}_2\text{O}.$

(Friedheim, Z. anorg. 1893, 4. 293.)

Cobaltous phosphomolybdate, $2\text{CoO}, \text{P}_2\text{O}_5, 4\text{MoO}_3 + x\text{H}_2\text{O}.$

(Arnfeld, Dissert. 1898.)

$3\text{CoO}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 16\frac{1}{2}\text{H}_2\text{O}$, and $+17\frac{1}{2}\text{H}_2\text{O}.$ Extremely sol. in H_2O . (Arnfeld.)

$3\text{CoO}, \text{P}_2\text{O}_5, 18\text{MoO}_3 + 38\text{H}_2\text{O}.$ Sol. in H_2O . (Arnfeld.)

$3\text{CoO}, \text{P}_2\text{O}_5, 24\text{MoO}_3 + 58\text{H}_2\text{O}$, and $+60\text{H}_2\text{O}.$ Sol. in H_2O . (Arnfeld.)

Cobaltous potassium phosphomolybdate,
 $\text{K}_2\text{O}, 2\text{CoO}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 15\text{H}_2\text{O}.$

(Arnfeld.)

$4\text{K}_2\text{O}, \text{CoO}, 2\text{P}_2\text{O}_5, 10\text{MoO}_3 + 12\text{H}_2\text{O}.$
 sol. in cold, easily sol. in hot H_2O . (Arnfeld.)

Croceocobaltic phosphomolybdate, $2\text{K}_2\text{O}, [\text{Co}(\text{NH}_3)_4(\text{NO}_2)_2]_2\text{O}, 2\text{H}_2\text{O}, 21\text{H}_2\text{O}.$

Sl. sol. in cold, easily in hot H_2O . Am. Ch. J. 3. 317.)

Gold phosphomolybdate ammonia, $1\text{K}_2\text{O}, 7\text{P}_2\text{O}_5, 3\text{MoO}_3, 24\text{NH}_3 + 21\text{H}_2\text{O}.$

Insol. in H_2O . (Gibbs, Am. Ch. J. 17. 172.)

Gold sodium phosphomolybdate and ammonia,
 $5\text{Au}_2\text{O}_3, \text{Na}_2\text{O}, \text{P}_2\text{O}_5, 11\text{MoO}_3, 1\text{H}_2\text{O}.$

Sol. in dil. HCl . Almost insol. in NH_4OAc . (Gibbs.)

Lead phosphomolybdate, $23\text{PbMoO}_4, 2\text{PbPO}_4 + 7\text{H}_2\text{O}.$

Sol. in 500,000 pts. H_2O . Insol. in H_2O + Aq. Easily sol. in KOH , NaOH , or HCl + Aq; somewhat less sol. in $\text{HC}_2\text{H}_3\text{O}_2$. (Beuf, Bull. Soc. (3) 3. 852.)

Lithium phosphomolybdate, $12\text{Li}_2\text{O}, 5\text{MoO}_3 + 18\text{H}_2\text{O}.$

Partially sol. in H_2O . (Ephraim, Z. 1909, 64. 233.)

$3\text{Li}_2\text{O}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 16\text{H}_2\text{O}.$ Ppt. (Ephraim, Z. anorg. 1910, 65. 233-6.)

$3\text{Li}_2\text{O}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 17\text{H}_2\text{O}.$ Ppt. $5\text{Li}_2\text{O}, 2\text{P}_2\text{O}_5, 8\text{MoO}_3 + 28\text{H}_2\text{O}.$ Easily sol. in H_2O . (E.)

$3\text{Li}_2\text{O}, \text{P}_2\text{O}_5, 12\text{MoO}_3 + 18\text{H}_2\text{O}.$ Ppt. $3\text{Li}_2\text{O}, \text{P}_2\text{O}_5, 18\text{MoO}_3 + 27\text{H}_2\text{O}.$ Sol. in H_2O . (E.)

Manganous phosphomolybdate, $3\text{MnO}, 5\text{MoO}_3 + 20\text{H}_2\text{O}.$

Very sol. in H_2O . Sl. sol. in alcohol. (Arnfeld.)

$3\text{MnO}, \text{P}_2\text{O}_5, 18\text{MoO}_3 + 38\text{H}_2\text{O}.$ Sol. in H_2O . (Arnfeld.)

$3\text{MnO}, \text{P}_2\text{O}_5, 24\text{MoO}_3 + 58\text{H}_2\text{O}.$ Sol. in H_2O . Easily sol. in H_2O . (Arnfeld.)

Manganous potassium phosphomolybdate,
 $2\text{MnO}, 3\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 10\text{MoO}_3 + 3\text{H}_2\text{O}.$

Sol. in hot H_2O . (Arnfeld.)

Manganous sodium phosphomolybdate,
 $7\text{MnO}, 9\text{Na}_2\text{O}, 2\text{P}_2\text{O}_5, 22\text{MoO}_3 + 3\text{H}_2\text{O}.$

Nearly insol. in cold H_2O . Sol. in H_2O but decomp. thereby. (Gibbs, Am. Ch. J. 1895, 17. 85.)

Nickel phosphomolybdate, $2\text{NiO}, \text{P}_2\text{O}_5, 4\text{MoO}_3 + x\text{H}_2\text{O}.$

Ppt. (Arnfeld.)

$3\text{NiO}, \text{P}_2\text{O}_5, 5\text{MoO}_3 + 20\text{H}_2\text{O}.$ Decomp. in H_2O . (Arnfeld.)

P_2O_5 , $18MoO_3 + 34H_2O$. Sol. in nfeld.)
 P_2O_5 , $24MoO_3 + 58H_2O$, and $+60H_2O$.
 it. Sol. in H_2O . (Arnfeld.)

potassium phosphomolybdate, K_2O ,
 P_2O_5 , $5MoO_3 + 13H_2O$.

by cold H_2O , but goes into solution on boiling. (Arnfeld.)

SiO_2 , $2P_2O_5$, $10MoO_3 + 12H_2O$. Sol. (Arnfeld.)

potassium phosphomolybdate, K_3PO_4 ,
 $O_3 + 1\frac{1}{2}H_2O = 3K_2O$, P_2O_5 , $22MoO_3$,
 P_2O_5 .

in H_2O . Easily sol. in alkalies. (and Struve.)

I_2O dissolve 0.007 g. at 30° ; 100 g. $O_3 + Aq$ dissolve 0.204 g. at 30° .
 ill. 90, Bur. of Chem., U. S. Dept. (1905.)

According to older authorities the formula
 $10MoO_3 + 1\frac{1}{2}H_2O$.

(Rammelsberg.)

I_2O , $24MoO_3$, $P_2O_5 + 3H_2O$. Sl. sol. O_3 .

H_2O , $44MoO_3$, $2P_2O_5 + 21H_2O$.
 m. Ch. J. 3. 317.)

P_2O_5 , $18MoO_3 + 11H_2O$, and $+15H_2O$

Sl. sol. in cold, very easily sol. O_3 . Can be cryst. from hot H_2O .
 n, Z. anorg. 1894, 7. 416.)

P_2O_5 , $17MoO_3 + 12H_2O$. Moder-
 in H_2O . (Elias, Dissert. 1906.)

P_2O_5 , $17MoO_3 + xH_2O$. Aqueous
 decomp. rapidly in the cold. (Kehr-
 anorg. 1894, 7. 423.)

$2H_2O$, $9MoO_3$, $P_2O_5 + 18H_2O$.

I_2O , $10MoO_3$, $P_2O_5 + 19H_2O$. Eas-
 H_2O . (Rammelsberg, B. 10. 1776.)

$5MoO_3$, P_2O_5 . Insol. in H_2O . Sol.
 Aq. (Rammelsberg.)

P_2O_5 , $2MoO_3 + 13H_2O$. Very sol. in
 Friedheim, Z. anorg. 4. 287.)

P_2O_5 , $4MoO_3 + 8H_2O$. Sol. in H_2O .
 n.)

diphosphopentamolybdate, $3K_2O$,
 $5MoO_3 + 7H_2O$.

H_2O ; precipitated by HNO_3 or HCl
 enkner, J. pr. 58. 261.)

P_2O_5 , $5MoO_3 + 6H_2O$. (Friedheim.)

diphosphopentamolybdate nitrate,
 O_3 , $5MoO_3$, $6KNO_3 + 9H_2O$.

, C. R. 66. 706.)

phosphomolybdates:

P_2O_5 , $22MoO_3 + 12H_2O$; $3Rb_2O$,
 $6MoO_3 + 12H_2O$; $6Rb_2O$, P_2O_5 , $18MoO_3$,
 $5Rb_2O$, $2P_2O_5$, $9MoO_3 + 13H_2O$;

P_2O_5 , $10MoO_3 + 15H_2O$. (Ephraim,
 1910, 65. 237-9.)

Silver phosphomolybdate, $7Ag_2O$, P_2O_5 ,
 $20MoO_3 + 24H_2O$.

Ppt. Sol. in dil. $HNO_3 + Aq$, forming—
 $2Ag_2O$, P_2O_5 , $20MoO_3 + 7H_2O$. Sl. sol. in
 H_2O . (Rammelsberg.)

Formula of first salt is—

$7Ag_2O$, $22MoO_3$, $P_2O_5 + 14H_2O$. Sol. in hot
 H_2O , but solution is quickly decomp. (Gibbs,
 Am. Ch. J. 3. 317.)

$7Ag_2O$, P_2O_5 , $24MoO_3$. Ppt. (Miolati, J.
 pr. 1908 (2) 77. 451.)

Silver diphosphopentamolybdate,
 $Ag_3Mo_5P_2O_{23} + 7H_2O$.

Easily sol. in H_2O . (Debray, C. R. 66. 705.)

Sodium phosphomolybdate.

Sol. in H_2O and $HNO_3 + Aq$. (Sonnen-
 schein, A. 104. 45.)

Na_2O , $5H_2O$, P_2O_5 , $18MoO_3 + xH_2O$.

$2Na_2O$, $4H_2O$, P_2O_5 , $18MoO_3 + xH_2O$.

$3Na_2O$, P_2O_5 , $18MoO_3 + 26H_2O$. (Fried-
 heim.)

$3Na_2O$, P_2O_5 , $24MoO_3 + 42H_2O$. (Rosen-
 heim and Pinsker, Z. anorg. 1911, 70. 79.)

Sodium diphosphopentamolybdate, $3Na_2O$,
 P_2O_5 , $5MoO_3 + 14H_2O$.

Easily sol. in H_2O . (Debray.)

Sodium auramine phosphomolybdate, Na_2O ,
 $5Au_2O_3$, $2P_2O_5$, $11MoO_3$, $15NH_3$.

Sol. in hot H_2O . Very sol. in hot HCl .
 (Gibbs, Am. Ch. J. 1895, 17. 171.)

Metaphosphomolybdic acid.

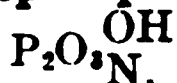
Ammonium monometaphosphomolybdate,
 $3(NH_4)_2O$, $4NH_4PO_3$, $10MoO_3 + 9H_2O$.

Very sol. in H_2O . (Gibbs, Am. Ch. J. 7.
 392.)

Barium hexametaphosphomolybdate, BaO ,
 $Ba_3(PO_3)_6$, $14MoO_3 + 55H_2O$.

Sol. in H_2O . (Gibbs.)

Pyrophosphonitrylic acid, $P_2HNO_4 =$



Not known in free state.

Ammonium pyrophosphonitrylate,



Insol. but gradually decomp. by H_2O .
 (Gladstone.)

Potassium —, KP_2NO_4 .

Insol. in H_2O . (Gladstone.)

Silver —, AgP_2NO_4 .

Ppt.

Triphosphonitrilic chloramide,

Sol. in H_2O without decomp.; sol. in ether, alcohol; sl. sol. in benzene. (Stokes, Am. Ch. J. 1895, 17. 287.)

Triphosphonitrilic tetrachlorhydrin,

Sol. in ether, alcohol, and H_2O ; insol. in benzene and CS_2 . (Stokes, Am. Ch. J. 1895, 17. 286.)

Triphosphonitrilic chloride, $P_3N_3Cl_4$.

Sol. in glacial acetic acid and H_2SO_4 . 100 pts. ether dissolve 46.5 pts. at 20° . (Liebig.)

Insol. in light petroleum; sol. in benzene. (Stokes, Am. Ch. J. 1897, 19. 783.)

Tetraphosphonitrilic chloride, $P_4N_4Cl_4$.

Sol. in alcohol, ether, benzene, H_2SO_4 . Sl. sol. in H_2O with decomp. 100 pts. ether dissolve 12.3 pts. at 20° . (Stokes, Am. Ch. J. 1895, 17. 281.)

Pentaphosphonitrilic chloride, $(PNCl_2)_5 = P_5N_5Cl_{10}$.

Sol. in benzene, light petroleum, acetic acid, ether, CS_2 ; insol. in H_2O . (Stokes, Am. Ch. J. 1897, 19. 790.)

Hexaphosphonitrilic chloride, $(PNCl_2)_6 = P_6N_6Cl_{12}$.

Sol. in benzene, light petroleum, ether, CS_2 ; insol. in H_2O . (Stokes.)

Heptaphosphonitrilic chloride, $(PNCl_2)_7 = P_7N_7Cl_{14}$.

Sol. in benzene, light petroleum, ether, CS_2 ; insol. in H_2O ; sol. in alcohol with decomp. (Stokes.)

Polyphosphonitrilic chloride, $(PNCl_2)_x$.

Depolymerizes on distillation; insol. in benzene and light petroleum and all neutral solvents; sol. in H_2O with decomp. (Stokes.)

Phosphonitrilochloramide, $P_2N_2Cl_2(NH_2)_2$.

Slowly sol. in H_2O with decomp. Insol. in ether and CS_2 . Sl. sol. in boiling CCl_4 . (Besson and Rosset, C. R. 1908, 146. 1149.)

Phosphonium bromide, PH_4Br .

Decomp. violently by H_2O .

Phosphonium chloride, PH_4Cl .

(Ogier, Bull. Soc. (2) 32. 483.)

Phosphonium titanium chloride, $2PH_4Cl, 3TiCl_4$.

Decomp. by H_2O , HCl , or alkalis + Aq. (Rose.)

Phosphonium iodide, PH_4I .

Decomp. by H_2O , alkalis, alcohol (Rose, Pogg. 46. 636.)

Decomp. by PCl_3 . (Wilde, B. 18. 21)

Phosphonium sulphate (?)

Deliquescent; very unstable. (C. R. 109. 644.)

Phosphoramidate, $P(NH_2)_3$.

Insol. in NH_4Br , $3NH_3$; sol. in NH_4I (Hugot, C. R. 1905, 141. 1235.)

Phosphortriamide, PON_3H_3 .

See Phosphoryl triamide.

Phosphoric acid, anhydrous, P_2O_5 .

See Phosphorus pentoxide.

Metaphosphoric acid, HPO_3 .

Sol. in H_2O . Not isolated. (Fl Pogg. 78. 362.)

Deliquescent. Sol. in H_2O , but solution decomp. into H_3PO_4 , slow cold, but more rapidly on heating solutions decomp. more rapidly than dil. (Giran, A. ch. 1903, (7) 30. 203)

Insol. in liquid CO_2 . (Büchner, Ch. 1906, 54. 674.)

Dimetaphosphoric acid, $H_2P_2O_5$.

Not isolated. (Fleitmann.)

Trimetaphosphoric acid, $H_3P_3O_7$.

Sol. in H_2O ; the solution is perm the cold, but on evaporation it is qu comp. to H_3PO_4 .

Tetrametaphosphoric acid, $H_4P_4O_{13}$.

Not isolated.

Hexametaphosphoric acid, $H_6P_6O_{21}$ (Glacial phosphoric acid.)

Deliquescent; easily sol. in H_2O with tion of heat and conversion into H_3PO_4 easily sol. in presence of slight impu

Insol. in liquid NH_3 . (Gore, Az 1898, 20. 828.)

Orthophosphoric acid, H_3PO_4 .

Very sol. in H_2O .

100 pts. of the solution contain at

26.23°	27.02°	29.42°	29.77°
95.9	95.98	96.15	96.11 pt

37.65°	39.35°	42.30° (mpt.
97.8	98.48	100 pts. H_2O

(Smith and Menzies, J. Am. Chem. 31. 1186.)

See also $10H_3PO_4 + H_2O$, and $2 H_2O$.

Sp. gr. of $H_3PO_4 + Aq$ containing

10	20	30	40	50
1.1	1.23	1.39	1.6	1.85

(Dalton.)

gr. of $H_2PO_4 + Aq.$

Sp. gr.	% P_2O_5	Sp. gr.	% P_2O_5
1.328	36.15	1.144	17.89
1.315	34.82	1.136	16.95
1.302	33.49	1.124	15.64
1.293	32.71	1.113	14.33
1.285	31.94	1.109	13.25
1.276	31.03	1.095	12.18
1.268	30.13	1.081	10.44
1.257	29.16	1.073	9.53
1.247	28.24	1.066	8.62
1.236	27.30	1.056	7.39
1.226	26.36	1.047	6.17
1.211	24.79	1.031	4.15
1.197	23.23	1.022	3.03
1.185	22.07	1.014	1.91
1.173	20.91	1.006	0.79
1.162	19.73
1.153	18.81

atta, C. N. 12. 160.)

city of $H_2PO_4 + Aq$ containing:12 18 % H_2PO_4 ,
0.0388 1.106536 54 % H_2PO_4 ,
0.2338 1.3840

chiff, A. 113. 183.)

 $d_4 + Aq$ at 15° . $a = \text{sp. gr. if } \%$
 $b = \text{sp. gr. if } \% \text{ is } H_2PO_4$.

b	%	a	b
1.0054	31	1.288	1.1962
1.0109	32	1.299	1.2036
1.0164	33	1.310	1.2111
1.0220	34	1.321	1.2186
1.0276	35	1.333	1.2262
1.0333	36	1.345	1.2338
1.0390	37	1.357	1.2415
1.0449	38	1.369	1.2493
1.0508	39	1.381	1.2572
1.0567	40	1.393	1.2651
1.0627	41	1.407	1.2731
1.0688	42	1.420	1.2812
1.0749	43	1.432	1.2894
1.0811	44	1.445	1.2976
1.0874	45	...	1.3059
1.0937	46	...	1.3143
1.1001	47	...	1.3227
1.1065	48	...	1.3313
1.1130	49	...	1.3399
1.1196	50	...	1.3486
1.1262	51	...	1.3573
1.1329	52	...	1.3661
1.1397	53	...	1.3750
1.1465	54	...	1.3840
1.1534	55	...	1.3931
1.1604	56	...	1.4022
1.1674	57	...	1.4114
1.1745	58	...	1.4207
1.1817	59	...	1.4301
1.1889	60	...	1.4395

ed by Gerlach, Z. anal. 8. 292.)

Sp. gr. of $H_2PO_4 + Aq$ at 17.5° .

% P_2O_5	Sp. gr.	% P_2O_5	Sp. gr.	% P_2O_5	Sp. gr.
1	1.007	24	1.208	47	1.476
2	1.014	25	1.219	48	1.491
3	1.021	26	1.229	49	1.506
4	1.028	27	1.240	50	1.521
5	1.036	28	1.250	51	1.536
6	1.044	29	1.261	52	1.551
7	1.053	30	1.272	53	1.566
8	1.061	31	1.282	54	1.581
9	1.070	32	1.293	55	1.597
10	1.078	33	1.304	56	1.613
11	1.086	34	1.315	57	1.629
12	1.095	35	1.326	58	1.645
13	1.103	36	1.338	59	1.661
14	1.112	37	1.350	60	1.677
15	1.120	38	1.362	61	1.693
16	1.129	39	1.374	62	1.709
17	1.139	40	1.386	63	1.725
18	1.148	41	1.398	64	1.741
19	1.158	42	1.410	65	1.758
20	1.168	43	1.423	66	1.775
21	1.178	44	1.436	67	1.792
22	1.188	45	1.448	68	1.809
23	1.198	46	1.460

(Hager, Adjumenta varia, Leipzig, 1876.)

Table for correction to be added or subtracted
for 1° change in temperature.

% P_2O_5	Corr.	% P_2O_5	Corr.
10-14	0.00035	36-45	0.00068
15-25	0.0004	46-55	0.00082
26-35	0.00052	56-68	0.001

(Hager.)

Sp. gr. of $H_2PO_4 + Aq.$

G. equivalents H_2PO_4 per litre	t°	Sp. gr. t°/t°
0.002572	17.714	1.001552
0.005142	17.706	1.0003051
0.01025	17.685	1.000595
0.02042	17.683	1.001158
0.03056	17.687	1.001708
0.04065	17.704	1.002252
0.0507	17.663	1.002790
0.10046	17.696	1.005412
0.19951	17.749	1.010560
0.29716	17.701	1.015584
0.49057	17.719	1.025469
0.5070	17.58	1.02627
5.0700	17.84	1.25162

(Kohlrausch, W. Ann. 1894, 53. 29.)

Miscible with conc. $HC_2H_3O_2 + Aq.$ Sol.
in 30 pts. warm creosote.1 cc. of a sat. solution of ether in H_2O dis-
solves 0.0886 grams H_2PO_4 .

1 cc. of a sat. solution of H_2O in ether dissolves 0.000033 grams H_3PO_4 .

Solutions of H_3PO_4 in H_2O containing less than 0.434 grams acid per 1 cc. lose an insignificant amount of acid to ether when agitated therewith. (Berthelot, C. R. 1896, 123. 345.)

The composition of the hydrates formed by H_3PO_4 at different dilutions is calculated from determinations of the lowering of the fr.-pt. produced by H_3PO_4 and of the conductivity and sp. gr. of $\text{H}_3\text{PO}_4 + \text{Aq.}$ (Jones, Am. Ch. J. 1905, 34. 331.)

$10\text{H}_3\text{PO}_4 + \text{H}_2\text{O}$. Solubility in H_2O .

100 pts. of the solution contain at:

24.11°	24.38°	24.40°
94.78	94.80	94.84 pts. H_3PO_4 ,

24.81°	25.41°	25.85°
94.95	95.26	95.54 pts. H_3PO_4 .

(Smith and Menzies, J. Am. Chem. Soc. 1909, 31. 1186.)

$2\text{H}_3\text{PO}_4 + \text{H}_2\text{O}$. Solubility in H_2O .

100 pts. of the solution contain at:

—16.3°	0.5°	14.95°	24.03°	27.0°
76.7	78.7	81.7	85.7	87.7 pts. H_3PO_4 ,

29.15°	29.35° (mpt.)
90.5	91.6 pts. H_3PO_4 ,

28.5°	27.0°	25.41°
92.5	93.4	94.1 pts. H_3PO_4 .

Retroflex part of curve.

(Smith and Menzies, J. Am. Chem. Soc. 1909, 31. 1186.)

Pyrophosphoric acid (Diphosphoric acid),
 $\text{H}_4\text{P}_2\text{O}_7$.

Very sol. in H_2O . The solution may be kept without change, but on heating it is converted into H_3PO_4 .

The acid in solution gradually changes to H_3PO_4 ; the conversion being more rapid with more concentrated solutions. (Montemartini and Egidi, Gazz. ch. it. 1902, 32. (1) 381.)

Phosphoric acid, $\text{H}_3\text{P}_2\text{O}_5$ (?)

Sol. in H_2O . (Joly, C. R. 100. 447.)

Phosphates.

The phosphates of NH_4 , K , Na , Li , Cs , and Rb are sol. in H_2O , with the exception of certain metaphosphates; the other phosphates excepting neutral Tl salts, are nearly insol. in H_2O , excepting when an excess of H_3PO_4 is present. The latter are all sol. in $\text{HNO}_3 + \text{Aq.}$

(a) **Metaphosphates.**

Monometaphosphates. Only alkali monometaphosphates are known, and they are all insol. in H_2O .

Dimetaphosphates. Alkali dimetaphosphates and some double salts containing alkali as one of the bases are sol. in H_2O , the rest are sl. sol. or insol. in H_2O .

Trimetaphosphates. All salts are insol. in H_2O .

Tetrametaphosphates. The alkali salts are sol. in H_2O , the others are insol.

Hexametaphosphates. The alkali salts are sol. in H_2O , the others insol., in H_2O , but sol. in $\text{Na hexametaphosphate} + \text{Aq.}$

(b) **Orthophosphates.** K , Na , Li , Rb orthophosphates are sol. in H_2O , the others are insol. in H_2O , but sol. in H_3PO_4 , and $\text{HNO}_3 + \text{Aq.}$; less easily in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ Pb , Al , and Fe orthophosphates are insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ Sl. sol. in H_2O , especially $\text{NH}_4\text{Cl} + \text{Aq.}$ solution they are pptd. by NH_3 . Orthophosphates insol. in H_2O are in an excess of alkali orthophosphates. All orthophosphates are insol. in alcohol.

(c) **Pyrophosphates.** Alkali pyrophosphates are sol. in H_2O ; the other pyrophosphates are insol. in H_2O , but are mostly sol. in an excess of $\text{pyrophosphate} + \text{Aq.}$

Aluminum metaphosphate, $\text{Al}_2(\text{PO}_3)_3$

Insol. in H_2O and conc. acids. (A. 61. 59.)

Aluminum orthophosphate, basic, $\text{P}_2\text{O}_5 + 18\text{H}_2\text{O}$.

Min. *Evansite*.

$4\text{Al}_2\text{O}_3, 3\text{P}_2\text{O}_5 + 18\text{H}_2\text{O}$. Ppt. in H_2O . (Rammelsberg.)

$2\text{Al}_2\text{O}_3, \text{P}_2\text{O}_5$.

+ $3\text{H}_2\text{O}$. Min. *Angelite*.

+ $5\text{H}_2\text{O}$. Min. *Kalaite* (Turquie) in $\text{HCl} + \text{Aq.}$

+ $6\text{H}_2\text{O}$. Decomp. by H_2O . (H. J. pr. (2) 37. 111.)

Min. *Peganite*. More or less sol. in $\text{HNO}_3 + \text{Aq.}$

+ $8\text{H}_2\text{O}$. Ppt. (Munroe, A. 11)

Min. *Fischerite*. Sl. attacked by $\text{HNO}_3 + \text{Aq.}$; sol. in $\text{H}_2\text{SO}_4 + \text{Aq.}$

$3\text{Al}_2\text{O}_3, 2\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}$, or $12\text{H}_2\text{O}$, acids, even after ignition. (Miller, 89.)

+ $10\text{H}_2\text{O}$. Min. *Caruleolactite* acids.

+ $12\text{H}_2\text{O}$. Min. *Wavellite*.

Aluminum orthophosphate, $\text{Al}_2(\text{PO}_4)_3$

Crystalline. Not attacked by $\text{HNO}_3 + \text{Aq.}$, difficultly by hot HCl (de Schulten, C. R. 98. 1583.)

Ignited $\text{Al}_2(\text{PO}_4)_3$ is sl. decomp. that solubility determinations are not reliable. For an extended discussion, see *Orthophosphates* (Cameron and Hurst, 1904, 22. 86)

Easily sol. in mineral acids, insol. in other organic acids. Easily sol. in H_2O , but is reprecipitated by $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in amount of alum + Aq (Rose), in acetate and other aluminum salts (Berzeliusscher, Z. anal. 6. 28). More sol. in phosphate in ammonium oxalate + Aq . (Milot.)

$\text{Citrate} + \text{Aq}$ dissolves 3% of the Al_2O_3 ; $\text{NH}_4\text{ citrate} + \text{Aq}$ dissolves 6.6% ; ammoniacal $\text{NH}_4\text{ citrate} + \text{Aq}$ completely in 25 min. (Erlenmeyer, Z. anal. 6. 28). $\text{LiOH} + \text{Aq}$, especially in presence of phosphates. (de-Koninck, Z. anal. 6. 28).

in presence of alkali tartrates or sugar, glycerine, etc.

ethyl acetate. (Naumann, B. 4. 1.)

scite. Very quickly sol. in warm Aq .

Min. *Zepharovitchite*.

Min. *Gibbsite*.

orthophosphate, acid, $2\text{Al}_2\text{O}_3, 16\text{H}_2\text{O}$.

insol. in acids after being ignited. (Milot, Z. anal. 22. 244.)

insol. in H_2O or alcohol. (Hautefeuille and Margottet, J. pr. (2) 16. 289.)

$\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}$. Insol. in acids after being ignited. (Milot.)

$\text{P}_2\text{O}_5 + 14\text{H}_2\text{O}$. Decomp. by cold

$\text{P}_2\text{O}_5 + 9\text{H}_2\text{O}$. Decomp. by hot water. (Meyer, A. 194. 200.)

$\text{P}_2\text{O}_5 + 3\text{H}_2\text{O} = \text{Al}_2(\text{H}_2\text{PO}_4)_3$. Deliquescent. Completely sol. in a little cold water. solution can be boiled without decomposition. (1 : 20) separates on boiling, which redissolves on more quickly the more dilute the solution. (Erlenmeyer, A. 194. 198.)

pyrophosphate, $\text{Al}_4(\text{P}_2\text{O}_7)_3 +$

2. Sol. in mineral acids, and insol. in acetic acid. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$, but when $\text{HCl} + \text{Aq}$ is reprecipitated by LiOH , and is not redissolved in an Aq . (Schwarzenberg, A. 65. 147.)

$\text{Al}_2\text{O}_3 + \text{Aq}$. (Rose, Pogg. 76. 19.)

metaphosphate, $\text{Al}_2\text{O}_3, 2\text{P}_2\text{O}_5$. (Hautefeuille and Margottet, C. R. 96. 1142.)

ammonium dihydrogen orthophosphate, $\text{H}_4\text{H}_2\text{PO}_4, \text{AlPO}_4$.

Decomp. by H_2O . Sol. in acids. Nearly insol. in 50% acetic acid. (M. Chem. Soc. 1907, 29. 720.)

Aluminum calcium phosphate, $\text{Al}_2\text{O}_3, 3\text{CaO}, \text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$.

Min. *Tavistockite*.

$2\text{Al}_2\text{O}_3, 6\text{CaO}, 3\text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$. Min. *Kirrolite*.

Aluminum calcium phosphate sulphate, $3\text{Al}_2\text{O}_3, \text{SO}_3, \text{Ca}_3(\text{PO}_4)_2 + 6\text{H}_2\text{O}$.

Min. *Svanbergite*. Scarcely attacked by $\text{HCl} + \text{Aq}$, and only sl. by $\text{H}_2\text{SO}_4 + \text{Aq}$.

Aluminum ferrous magnesium phosphate, $(\text{Mg}, \text{Fe})_2\text{Al}_2\text{P}_2\text{O}_7 + 4\text{H}_2\text{O}$.

Min. *Childrenite*. Slowly sol. in $\text{HCl} + \text{Aq}$.

Min. *Eosphorite*. Sol. in HNO_3 or $\text{HCl} + \text{Aq}$.

$(\text{Mg}, \text{Fe})\text{Al}_2\text{P}_2\text{O}_7 + \text{H}_2\text{O}$. Min. *Lazulite*. Only sl. attacked by acids, when not previously ignited.

Aluminum lithium phosphate, $\text{Al}_2(\text{PO}_4)_3, 4\text{Li}_2\text{PO}_4 + 30\text{H}_2\text{O}$.

Precipitate. (Berzelius.)

Insol. in H_2O ; easily sol. in acids.

Aluminum magnesium phosphate.

Min. *Lazulite*.

See Phosphate, aluminum ferrous magnesium.

Aluminum potassium phosphate, $\text{Al}_2\text{O}_3, \text{K}_2\text{O}, 2\text{P}_2\text{O}_5$.

Insol. in acids. (Ouvrard, A. ch. (6) 16. 289.)

$2\text{Al}_2\text{O}_3, 2\text{K}_2\text{O}, 3\text{P}_2\text{O}_5$. (Ouvrard.)

Aluminum silver metaphosphate, $2\text{Al}_2\text{O}_3, \text{Ag}_2\text{O}, 4\text{P}_2\text{O}_5$.

(Hautefeuille and Margottet, C. R. 96. 849, 1142.)

Aluminum sodium pyrophosphate,

$\text{Al}_2\text{Na}_2(\text{P}_2\text{O}_7)_2$.

Insol. in H_2O and acids. (Wallroth.)

Nearly insol. in acids. (Ouvrard, A. ch. (6) 16. 338.)

$2\text{Al}_2\text{O}_3, 3\text{Na}_2\text{O}, 3\text{P}_2\text{O}_5$. Sol. in $\text{HNO}_3 + \text{Aq}$. (Ouvrard.)

$\text{Al}_4(\text{P}_2\text{O}_7)_3, 2\text{Na}_4\text{P}_2\text{O}_7 + 30\text{H}_2\text{O}$.

Very difficultly sol. in H_2O . (Pahl, Bull. Soc. (2) 22. 122.)

Aluminum phosphate lithium fluoride,

$2\text{Al}_2(\text{PO}_4)_3, 3\text{LiF}$.

Min. *Amblygonite*. Sl. attacked by $\text{HCl} + \text{Aq}$, more easily by $\text{H}_2\text{SO}_4 + \text{Aq}$.

Ammonium metaphosphate, NH_4PO_3 .

Insol. in H_2O . (Fleitmann, Pogg. 78. 345.)

Ammonium dimetaphosphate, $(\text{NH}_4)_2(\text{PO}_3)_2$.

Sol. in 1.15 pts. cold or hot H_2O . (Fleitmann, Pogg. 78. 245.) More sol. in dil. alcohol than Na or K salt.

Sol. in 0.9 pts. H₂O; easily sol. in acids especially by boiling with conc. H₂SO₄. (Glatzel, Dissert. 1880.)

Ammonium trimetaphosphate, (NH₄)₃P₃O₉.
Very sol. in H₂O. (Lindbom, Acta Lund. 1873. 15.)

Ammonium tetrametaphosphate, (NH₄)₄P₄O₁₂.
Sol. in H₂O. (Warschauer, Z. anorg. 1903, 36. 177.)
+4H₂O. Much more sol. in H₂O than the K or Na salt. 1 pt. is sol. in 8 pts. H₂O. (Glatzel, Dissert. 1880.)

Ammonium pentametaphosphate, (NH₄)₅P₅O₁₅.
Sol. in H₂O. (Tammann, J. pr. 1892, (2) 45. 455.)

Ammonium dekametaphosphate, (NH₄)₁₀P₁₀O₃₀.
Very sl. sol. in H₂O; 100 g. H₂O dissolved 1.20-1.54 g. in 2 months. Easily sol. in hot H₂O with decomp. (Tammann, J. pr. 1892, (2) 45. 448.)
+12H₂O. (Tammann, J. pr. 1892, (2) 45. 465.)

Ammonium orthophosphate, (NH₄)₃PO₄+3H₂O.
Difficultly sol. in H₂O.
Less sol. in H₂O than (NH₄)₂HPO₄. (Berzelius.)
Insol. in alkalies+Aq. (Berzelius.)
Sl. sol. in H₂O. Decomp. in the air. (Schottländer, Z. anorg. 1894, 7. 344.)

Solubility in H₃PO₄+Aq at 25°.

In 1000 g. of the solution, mols.	
NH ₄	PO ₄
7.42	0.084
5.02	0.20
2.95	0.46
3.04	1.02
3.32	1.32
4.78	2.32

(D'Ans and Schreiner, Z. phys. Ch. 1910, 75. 105.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)
+5H₂O. (Sestini, Gazz. ch. it. 9. 298.)

Ammonium hydrogen orthophosphate, (NH₄)₂HPO₄.
Easily sol. in H₂O. Effloresces to form NH₄H₂PO₄. (Schiff, A. 112. 88.)
Sol. in 4 pts. cold, and less hot H₂O. Solution loses NH₃ by boiling. Insol. in alcohol.

100 g. H₂O dissolve 131 g. at 15°; sp. of sat. solution = 1.343. (Greenish Smith, Pharm. J. 1901, 66. 774.)

Solubility in H₃PO₄+Aq at 25°.

In 1000 g. of the solution, mols.	
NH ₄	PO ₄
6.42	3.23
6.46	3.74
6.56	4.01
6.78	4.34
7.26	4.83
7.16	4.82

(D'Ans and Schreiner, Z. phys. Ch. 19105.)

Insol. in acetone. (Eidmann, C. C. II, 1014; Naumann, B. 1904, 37. 4329.)

Ammonium dihydrogen orthophosphate, NH₄H₂PO₄.
Does not effloresce.
Less easily sol. in H₂O than (NH₄)₂HPO₄. (Mitscherlich, A. ch. 19. 385.)
Sol. in 5 pts. cold, and less hot H₂O.

Solubility in H₃PO₄+Aq at 25°.

In 1000 g. of the solution, mols.	
NH ₄	PO ₄
6.72	4.54
5.62	3.88
4.62	3.36
2.72	2.59
2.50	2.54
2.58	4.29
2.76	6.21
3.06	7.70
3.10	7.86

(D'Ans and Schreiner, Z. phys. Ch. 19106.)

Insol. in acetone. (Eidmann, C. C. II. 1014; Naumann, B. 1904, 37. 4329.)

Ammonium orthophosphate, acid.
Decomp. into NH₄H₂PO₄ at 7 (Parravano and Mieli, Gazz. ch. it. 190II. 536.)

Ammonium pyrophosphate, (NH₄)₂P₂O₇
Easily sol. in H₂O. Alcohol precipit from the aqueous solution. (Schwarzenberg, A. 65. 141.)
Insol. in acetone. (Eidmann, C. C. II. 1014; Naumann, B. 1904, 37. 4329.)

Ammonium hydrogen pyrophosphate (NH₄)₂H₂P₂O₇.
Very sol. in H₂O. Insol. in al (Schwarzenberg, A. 65. 141.)

m barium trimetaphosphate,
 $(\text{BaP}_3\text{O}_9)_2 + \text{H}_2\text{O}.$

sol. in H_2O . (Lindbom.)

m cadmium dimetaphosphate,
 $(\text{CdP}_2\text{O}_7)_2, \text{CdO}, 2\text{P}_2\text{O}_5 + 3\text{H}_2\text{O} =$
 $(\text{Cd}(\text{P}_2\text{O}_6)_2)_2.$

cent. (Fleitmann, Pogg. 78. 347.)

m cadmium orthophosphate,
 $\text{CdPO}_4 + 1\frac{1}{4}\text{H}_2\text{O}.$

sol. in $\text{NH}_4\text{OH} + \text{Aq}$ and acids.
 Gm. K. Handb. 6^{te} Aufl. III. 74.)

m calcium dimetaphosphate,
 $(\text{CaP}_2\text{O}_6)_2 + 2\text{H}_2\text{O}.$

sol. in H_2O . Not decomp. by
 leitmann, Pogg. 78. 344.)

m calcium phosphate, $\text{NH}_4\text{CaPO}_4 +$

Herzfeld and Feuerlein, Z. anal. 20.

Not completely decomp. by cold
 hour; rapidly by hot H_2O . (Lasne,
 1902, (3) 27. 131.)

m chromium orthophosphate, basic,
 $(\text{H}_2\text{PO}_4)_2, 2\text{CrPO}_4, 4\text{Cr}(\text{OH})_3.$

J. Am. Chem. Soc. 1907, 29. 1196.)
 $\text{HPO}_4, 2\text{CrPO}_4 + 3\text{H}_2\text{O}.$ Ppt.

m chromic pyrophosphate,
 $\text{CrP}_2\text{O}_7 + 6\text{H}_2\text{O}.$

in cold H_2O . Decomp. by boiling
 senheim, B. 1915, 48. 586.)

m cobaltous metaphosphate.

ely sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq}$.
 pr. 3. 215.)

m cobaltous orthophosphate,
 $\text{CoPO}_4 + \text{H}_2\text{O}.$

comp. by boiling H_2O . (Debray, J.
) 46. 121.)

Ppt. (Chancel, 1862.)
 $(\text{H}_2\text{PO}_4)_2 + 4\text{H}_2\text{O}.$ Insol. in H_2O .

m copper dimetaphosphate,
 $(\text{P}_2\text{O}_6)_2, \text{CuP}_2\text{O}_6 + 2\text{H}_2\text{O}.$

sol. in H_2O ; insol. in alcohol.
 n, Pogg. 78. 345.)

Efflorescent. Very sl. sol. in
 l. in alcohol. (F.)

50 pts. H_2O . Slowly attacked by
 nly boiling H_2SO_4 attacks easily.
 Dissert. 1880.)

n glucinum orthophosphate,
 $\text{HfPO}_4.$

cold, sl. sol. in hot H_2O . (Rössler,
 148.)

Ammonium glucinum sodium orthophosphate,
 $(\text{NH}_4)_2\text{GlNa}_2(\text{PO}_4)_2 + 7\text{H}_2\text{O}.$

(Scheffer, A. 109. 146.)

Ammonium iron (ferrous) orthophosphate,
 $\text{NH}_4\text{FePO}_4 + \text{H}_2\text{O}.$

Insol. even in boiling H_2O . When still
 moist, easily sol. in dil. acids, but sparingly
 and slowly sol. after drying, even in conc.
 acids. Decomp. by NH_4OH , KOH , and
 $\text{NaOH} + \text{Aq}$. Insol. in alcohol. (Otto, J. pr.
 2. 409.)

$(\text{NH}_4)_2\text{FeH}_2(\text{PO}_4)_2 + 4\text{H}_2\text{O}.$ (Debray.)

Ammonium iron (ferric) hydrogen orthophos-
phate, basic, $2(\text{NH}_4)_2\text{HPO}_4, 3\text{FePO}_4,$
 $3\text{Fe}(\text{OH})_3.$

Ppt. Insol. in 95% alcohol. (Cohen, J.
 Am. Chem. Soc. 1907, 29. 719.)

Ammonium iron (ferric) hydrogen orthophos-
phate, $\text{NH}_4\text{H}_2\text{Fe}(\text{PO}_4)_2.$

Ppt. Same properties as Na salt. (Wein-
 land, Z. anorg. 1913, 84. 356.)

Partially hydrolyzed by H_2O . Readily sol.
 in HCl , HNO_3 , H_2SO_4 and H_3PO_4 . Partially
 hydrolyzed by cold $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in
 excess of hot $\text{NH}_4\text{OH} + \text{Aq}$. Completely
 hydrolyzed by caustic alkalis. Practically
 insol. in 50% acetic acid. (Cohen, J. Am.
 Chem. Soc. 1907, 29. 718.)

Ammonium lead dimetaphosphate,
 $(\text{NH}_4)_2\text{Pb}(\text{P}_2\text{O}_6)_2.$

Very difficultly sol. in H_2O and acids.
 (Fleitmann, Pogg. 78. 343.)

Ammonium lithium metaphosphate, $\text{Li}_2\text{O},$
 $2(\text{NH}_4)_2\text{O}, 3\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}.$

Not appreciably sol. in cold H_2O but
 rapidly and abundantly sol. in H_2O at 70° .
 (Taminann, J. pr. 1892, (2) 45. 442.)

Ammonium lithium phosphate, $(\text{NH}_4)_2\text{LiPO}_4.$

Sl. sol. in H_2O . (Berzelius.)

Ammonium magnesium metaphosphate,
 $(\text{NH}_4)_2\text{O}, 2\text{MgO}, 2\text{P}_2\text{O}_5 + 9\text{H}_2\text{O} (?)$.

Sol. with difficulty in H_2O or acids when
 heated. Easily sol. in H_2O before heating.
 (Wach, Schw. J. 59. 29.)

Precipitated from aqueous solution by
 alcohol.

Ammonium magnesium dimetaphosphate,
 $(\text{NH}_4)_2\text{Mg}(\text{P}_2\text{O}_6)_2 + 6\text{H}_2\text{O}.$

Efflorescent. (Fleitmann, Pogg. 78. 346.)

Ammonium magnesium phosphate,
 NH_4MgPO_4 , and $+6\text{H}_2\text{O}.$

1 l. H_2O dissolves 66 mg. anhydrous
 NH_4MgPO_4 at 15° . (Fresenius, A. 55. 109.)

1 l. H_2O dissolves 74.1 mg. anhydrous
 NH_4MgPO_4 at $20.5-22.5^\circ$. (Ebermayer.)

1 l. H_2O dissolves 106 mg. anhydrous NH_4MgPO_4 . (Liebig.)

Insol. in H_2O , but when boiled with H_2O it loses NH_3 and H_2O . (Struve, Z. anal. 1898, 37. 485.)

Solubility of $NH_4MgPO_4 \cdot 6H_2O$ in H_2O at t° .

t°	G. salt in 100 g. H_2O
0	0.0231
20	0.0516
40	0.0359
50	0.0303
60	0.0401
70	0.0163
80	0.0195

(Wenger, Dissert. Geneva, 1911.)

Aqueous solution is precipitated by NH_4OH , but not by $Na_2HPO_4 + Aq$. (Fresenius.)

Sol. in 44,600 pts. H_2O containing ammonia. More sol. in H_2O containing NH_4Cl , and is sol. in 7548 pts. of a solution containing 1 pt. NH_4Cl to 5 pts. H_2O and ammonia, and in 15,627 pts. of a solution containing 1 pt. of NH_4Cl to 7 pts. H_2O and ammonia. (Fresenius.)

According to Kremers (J. pr. 55. 190), a solution of 3 pts. H_2O to 1 pt. $NH_4OH + Aq$ of 0.96 sp. gr. is best suited for washing the precipitated NH_4MgPO_4 .

According to Ebermayer (J. pr. 60. 41), 1 pt. anhydrous salt is sol. in 13,497 pts. H_2O at 23° ; in 31,038 pts. $NH_4OH + Aq$ (4 pts. H_2O : 1 pt. $NH_4OH + Aq$ of 0.961 sp. gr.) at 21.25° ; in 36,764 pts. $NH_4OH + Aq$ (3 pts. H_2O : 1 pt. $NH_4OH + Aq$) at 20.6° ; in 43,089 pts. $NH_4OH + Aq$ (1 pt. H_2O : 1 pt. $NH_4OH + Aq$) at 22.5° ; in 45,206 pts. $NH_4OH + Aq$ (1 pt. H_2O : 2 pts. $NH_4OH + Aq$) at 22.5° ; in 52,412 pts. $NH_4OH + Aq$ (1 pt. H_2O : 3 pts. $NH_4OH + Aq$) at 22.5° ; in 60,883 pts. pure $NH_4OH + Aq$ (sp. gr. 0.961) at 22.5° .

Almost absolutely insol. in H_2O containing $\frac{1}{2}$ vol. $NH_4OH + Aq$ (sp. gr. 0.96) and NH_4Cl , i. e., much more insol. than given by Fresenius. (Kubel, Z. anal. 8. 125.)

According to Kissel (Z. anal. 8. 173), 1 l. $NH_4OH + Aq$ (3 pts. H_2O : 1 pt. $NH_4OH + Aq$ of 0.96 sp. gr.) dissolves 4.98 mg. in 24 hours, while 13.9 mg. are dissolved if 18 g. NH_4Cl to a litre of H_2O are also present.

$(NH_4)_2SO_4 + Aq$ containing 2.2 g. per litre dissolves 71.7 mg.; 3.0 g., 113 mg.; 10 g., 147 mg.; $NaCl + Aq$ containing 2 g. $NaCl$ per l. dissolves 123.4 mg.; $NaNO_3 + Aq$ containing 3 g. $NaNO_3$ per l. dissolves 93.1 mg. (Liebig, A. 106. 196.)

Completely insol. in water containing ammonium phosphate or ammonium sodium phosphate. (Berzelius.)

800 ccm. H_2O , sat. with CO_2 , dissolve 1.425 g. (Liebig.)

Easily sol. in $H_2SO_4 + Aq$, acetic and other

acids, also in boiling solution of citrate. (Millot, Bull. Soc. (2) 18. 20)

When in presence of Fe or Al salt to a considerable extent in $H_2C_2H_3O_2$ 6 g. NH_4Cl in 100 ccm. H_2O cost ccm. 6.34% $NH_4OH + Aq$ dissolve = 0.0029 g. $Mg_3P_2O_7$; 1 g. (NH_4 100 ccm. H_2O , and $NH_4OH + Aq$ 0.0061 g. $Mg_3P_2O_7$; 2 g. citric acid of $NH_4OH + Aq$ dissolve = 0.0147 g. Solubility prevented by excess of mixture. (Lindo, C. N. 48. 217.)

Solubility of $NH_4MgPO_4 \cdot 6H_2O$ in at t° .

(G. salt dissolved in 100 g. sol

t°	5% $NH_4NO_3 + Aq$	5% $NH_4Cl + Aq$
0	0.1100	0.0597
20	0.0463	0.1000
30	0.0546	0.1133
40	0.0645	0.0713
50	0.0723	0.0031
60	0.0846	0.1728
70	0.0834	0.1239
80	0.1000	0.1913

t°	4% $NH_4OH + Aq$ and 5% $NH_4Cl + Aq$	4% NH_4OH 10% NH_4Cl
20	0.0165	0
60	0.0274	0

(Wenger, Dissert. Geneva, 1

About 3 times as sol. in $Ca(C_2H_3O_2)_2$ as in $NaC_2H_3O_2 + Aq$, but solubility prevented by excess of $MgCl_2$. (Ville (2) 18. 316.)

Sl. sol. in ammonium citrate + 1 mg. 400 g. ammonium citrate Solubility = 0.457% at ord. temp. at 50° . (Bolus, Ch. Z. 1903, 27. 11 Min. Struete.

+ H_2O . Insol. in H_2O or citric (Millot and Maquenne, Bull. Soc. (

Ammonium magnesium hydrogen phosphate, $(NH_4)_2MgH_2(PO_4)_2 \cdot 3H_2O$ (Graham.)

Ammonium magnesium phosphate $5MgO, (NH_4)_2O, 2P_2O_5 + 24H_2O$ (Gawalovsky, C. C. 1888. 721.)

Ammonium magnesium sodium phosphate, $(NH_4)_2Mg_2Na(P_2O_7)_4$

Insol. in H_2O and not decomposed by acids. (Berthelot and André, A. ch. 18 1885.)

Manganous dimetaphosphate,
 $\text{Mn}(\text{PO}_3)_2 + 4\text{H}_2\text{O}$.

Easily attacked by acids. (Glatzel, 1880.)

(Fleitmann, Pogg. 78. 346.)

Manganous orthophosphate,
 $\text{Mn}_3(\text{PO}_4)_2 + \text{H}_2\text{O}$.

92 pts. cold, and 20,122 pts. and in 17,755 pts. $\text{NH}_4\text{Cl} + \text{Aq}$. (Fresenius.)

Insol. in H_2O below 70° ; at 70° dissolve 0.0052 g. salt; at 80° , larger, Dissert. Geneva, 1911.)

Insol. in dil. acids. Decomp. by heat but not by $\text{NH}_4\text{OH} + \text{Aq}$ or HCl . Insol. in NH_4OH or NH_4 salts (3.)

Solubility in salts + Aq at t° .
 (1 g. in 100 g. solvent.)

5% H_2NO_2 + Aq.	5% $\text{NH}_4\text{Cl} + \text{Aq}$.	1 pt. NH_4OH D = 0.96 + 4 pts. H_2O .
0.0206	0.0020	0.0116
0.0200	0.0255	0.0122
0.0226	0.0345	
0.0209	0.0386	0.0118
0.0226	0.0355	0.0132
0.0270	0.0384	0.0193
0.0281	0.0414	0.0191
0.0326	0.0451	0.0197

(Wenger, l. c.)

Insol. in alcohol.
 Insol. in acetone. (Naumann, B. 1904, 37.)

Manganic pyrophosphate,
 $\text{Mn}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Very cold H_2O with separation of MnO_2 . (Enheim, B. 1915, 48. 584.)

Manganous sodium pyrophosphate,
 $\text{Na}_4\text{Mn}_2(\text{P}_2\text{O}_7)_2 + 3\text{H}_2\text{O}$.

Insol. in water or alcohol. Easily sol. in very hot, J. pr. 2. 418.)

$\text{Na}_4(\text{NH}_4)_4\text{Mn}_2(\text{P}_2\text{O}_7)_2 + 3\text{H}_2\text{O}$ according to Berzelius.

Mercuric metaphosphate.

Insol. in water, or at least in $\text{NH}_4\text{OH} + \text{Aq}$. (3. 216.)

Nickel metaphosphate.

Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$, from which it is obtained on evaporation of the NH_3 . (3. 215.)

Nickel dimetaphosphate,
 $\text{Ni}_2(\text{PO}_3)_2 + 4\text{H}_2\text{O}$.

Insol. in H_2O . (Glatzel, Dissert.

Ammonium nickel orthophosphate,
 $\text{NH}_4\text{NiPO}_4 + 2\text{H}_2\text{O}$.

Ppt. (Debray, C. R. 59. 40.)
 $+ 6\text{H}_2\text{O}$. Decomp. by boiling H_2O . (Debray.)

Ammonium potassium dimetaphosphate,
 $(\text{NH}_4)_3\text{K}_3(\text{P}_2\text{O}_6)_2$.

More sol. in H_2O than following salt. (Fleitmann, Pogg. 78. 341.)

$\text{NH}_4\text{K}_3\text{P}_4\text{O}_{12} + 2\text{H}_2\text{O}$. Difficultly sol. in H_2O . (Fleitmann.)

Ammonium potassium pyrophosphate,
 $\text{NH}_4\text{K}_2\text{HP}_2\text{O}_7 + \frac{1}{2}\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O . Decomp. on boiling. (Schwarzenberg.)

Ammonium sodium dimetaphosphate,
 $\text{NH}_4\text{NaP}_2\text{O}_6 + \text{H}_2\text{O}$.

More sol. in H_2O than $\text{Na}_2\text{P}_2\text{O}_6$, but less than $(\text{NH}_4)_2\text{P}_2\text{O}_6$. Less sol. in alcohol than in H_2O . (Fleitmann, Pogg. 78. 340.)

Ammonium sodium orthophosphate,
 $(\text{NH}_4)_2\text{NaPO}_4 + 4\text{H}_2\text{O}$.

Decomp. by H_2O . Cryst. from $\text{NH}_4\text{OH} + \text{Aq}$ of 0.96 sp. gr. From H_2O solution, $\text{NaNH}_4\text{HPO}_4 + 4\text{H}_2\text{O}$ separates out. (Uelsmann, Arch. Pharm. (2) 99. 138.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)
 $+ 5\text{H}_2\text{O}$.

$\text{NH}_4\text{Na}_2\text{PO}_4 + 12\text{H}_2\text{O}$. (Herzfeld, Z. anal. 20. 191.)

$(\text{NH}_4)_3\text{Na}(\text{PO}_4)_2 + 6\text{H}_2\text{O}$. Sol. in H_2O with decomp. Cryst. from hot conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Uelsmann, Arch. Pharm. (2) 99. 138.)

Ammonium sodium hydrogen phosphate (Microcosmic salt), $\text{NH}_4\text{NaHPO}_4 + 4\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O . Sol. in 6 pts. cold, and 1 pt. boiling H_2O . Insol. in alcohol.

Aqueous solution gives off NH_3 , especially if hot.

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

Min. *Stercorite*.

$+ 5\text{H}_2\text{O}$. (Uelsmann.)

The composition of the hydrates formed by this salt at different dilutions is calculated from determinations of the lowering of the fr.-pt. produced by the salt and of the conductivity and sp. gr. of its aqueous solutions. (Jones, Am. Ch. J. 1905, 34. 319.)

$(\text{NH}_4)_3\text{Na}_3\text{H}_3(\text{PO}_4)_4 + 3\text{H}_2\text{O}$. Decomp. by H_2O . (Filhol and Senderens, C. R. 93. 388.)

Ammonium sodium pyrophosphate,
 $(\text{NH}_4)_2\text{Na}_2\text{P}_2\text{O}_7 + 5\text{H}_2\text{O}$.

Easily sol. in H_2O . Aqueous solution decomp. by boiling. (Schwarzenberg, A. 65. 142.)

$+ 6\text{H}_2\text{O}$. (Rammelsberg.)

Ammonium sodium glucinum orthophosphate,
 $(\text{NH}_4)_2\text{Na}_2\text{Gl}(\text{PO}_4)_2 + 7\text{H}_2\text{O}$.

Precipitate. (Scheffer.)

Ammonium thallos orthophosphate,
 $(\text{NH}_4)_2\text{PO}_4$, $(\text{NH}_4)_2\text{TiPO}_4$, or $\text{H}_2\text{NH}_4\text{PO}_4$,
 HTl_2PO_4 .

Sol. in H_2O . (Lamy; Rammelsberg.)

Ammonium uranyl phosphate,
 $\text{NH}_4(\text{UO}_2)\text{PO}_4 + x\text{H}_2\text{O}$.

Insol. in H_2O and $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in mineral acids, from which it is precipitated by $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$, in which it is insol. (Knop.)

+ $3\text{H}_2\text{O}$. Insol. in H_2O and acetic acid. Sol. in all mineral acids, oxalic acid and $\text{M}_2\text{CO}_3 + \text{Aq}$. (Lienau, Dissert. 1898.)

Ammonium vanadium phosphate.

See Phosphovanadate, ammonium.

Ammonium zinc dimetaphosphate,
 $(\text{NH}_4)_2\text{Zn}(\text{P}_2\text{O}_5)_2 + 6\text{H}_2\text{O}$.

Efflorescent. (Fleitmann, Pogg. 78. 347.)

+ $4\text{H}_2\text{O}$. Sol. in 70 pts. H_2O . Decomp. by H_2SO_4 . (Glatzel, Dissert. 1880.)

Ammonium zinc orthophosphate, basic,
 3NH_3 , 2ZnO , $\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}$.

(Rother, A. 1867, 143. 356.)

$4(\text{NH}_4)_2\text{O}$, 6ZnO , $3\text{P}_2\text{O}_5 + 4\text{H}_2\text{O}$. (Schweikert, A. 1868, 145. 57.)

Ammonium zinc orthophosphate, NH_4ZnPO_4 ,
 $+ \text{H}_2\text{O}$.

Insol. in H_2O . Sol. in acids, and caustic alkalies. (Bette, A. 15. 129.)

Ammonium zinc hydrogen phosphate,
 $\text{NH}_4\text{H}_2\text{PO}_4$, $\text{ZnHPO}_4 + \text{H}_2\text{O}$.

Insol. in H_2O . (Debray.)

$4(\text{NH}_4)_2\text{O}$, 6ZnO , $3\text{P}_2\text{O}_5$. (Schweikert, A. 145. 57.)

$3(\text{NH}_4)_2\text{O}$, 4ZnO , $2\text{P}_2\text{O}_5 + 13\text{H}_2\text{O}$. (Rother, A. 143. 356.)

Ammonium phosphate selenate.

See Selenophosphate, ammonium.

Barium triphosphate, 5BaO , $3\text{P}_2\text{O}_5$.

Insol. in H_2O ; insol. in acids after heating to a high temp. (Schwarz, Z. anorg. 1895, 9. 264.)

Barium metaphosphate, $\text{Ba}(\text{PO}_3)_2$.

Insol. in H_2O or dil. acids. (Maddrell, A. 61. 61.)

Not decomp. by boiling with acids or alkali carbonates + Aq . (Fleitmann, Pogg. 78. 352.)

Barium dimetaphosphate, $\text{BaP}_2\text{O}_6 + 2\text{H}_2\text{O}$.

More difficultly sol. in H_2O than $\text{Ba}_3(\text{P}_3\text{O}_{10})_2$. Slightly attacked by boiling conc. $\text{HCl} + \text{Aq}$ or $\text{HNO}_3 + \text{Aq}$. Easily decomp. by H_2SO_4 . (Fleitmann, Pogg. 78. 254.)

Barium trimetaphosphate, $\text{Ba}_3(\text{P}_3\text{O}_{10})_2 + 2\text{H}_2\text{O}$.

Somewhat sol. in H_2O . (Fleitmann, 313.)

+ $6\text{H}_2\text{O}$. Easily sol. in $\text{HCl} + \text{Aq}$. (Fleitmann, 313.)

1 l. H_2O dissolves 2.589 g. at ord. (Wiesler, Z. anorg. 1901, 28. 198.)

Barium hexametaphosphate, $\text{Ba}_3\text{P}_6\text{O}_{36}$.

Sol. in H_2O only after boiling several

Nearly insol. in H_2O . (Lüder, Z. 5. 15.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wackenrodt)

Sol. in $\text{Na}_4\text{P}_6\text{O}_{36} + \text{Aq}$. Sol. in HNO_3

After ignition it is nearly insol. in HNO_3

Barium orthophosphate, $\text{Ba}_3(\text{PO}_4)_2$.

Precipitate. Very sl. sol. or insol. in H_2O . (Graham, Pogg. 32. 49.)

Sol. in $\text{HCl} + \text{Aq}$. Decomp. by SCl_2

Insol. in methyl acetate. (Naum, 1909, 42. 3790.)

Barium hydrogen phosphate, BaHPO_4 .

Sol. in 10,000 pts. H_2O . (Malaguti, 3) 51. 346.)

Sol. in 20,570 pts. H_2O at 20° . (Fleitmann, 1833.)

Not completely soluble in water with CO_2 , but BaCl_2 causes no ppt. in Na_2HPO_4 Aq containing 7.16 g. or less Na_2HPO_4 per litre after it has been saturated with CO_2 . (Setschenow, C. C. 1875. 97.)

Easily sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$, and dil. Aq . $\text{HNO}_3 + \text{Aq}$ of 1.275 sp. gr. if not has scarcely any solvent action, but it dissolves on dilution until a maximum is reached when 10 vols. of H_2O have been added. (Bischof, Schw. J. 67. 39.)

Sol. in 367–403 pts. acetic acid (1.1 gr.) at 22.5° . (Bischof, l. c.)

Easily sol. in H_2O containing NH_4NO_3 , or NH_4 succinate, from which it is completely pptd. by NH_4OH . (Rose.)

Insol. in Na_2HPO_4 or $\text{BaCl}_2 + \text{Aq}$. (Pogg. 76. 23.)

More sol. in BaCl_2 or $\text{NaCl} + \text{Aq}$ than in H_2O , 1 pt. BaHPO_4 being sol. in 43 H_2O containing 1.2% NaCl and 0.8% BaCl_2 . (Ludwig, Arch. Pharm. (2) 56. 265.)

Sol. in Na citrate + Aq . (Spiller.)

Barium tetrahydrogen phosphate,
 $\text{BaH}_4(\text{PO}_4)_2$.

Sol. in H_2O . (Mitscherlich, 1831.)

Decomp. by much H_2O into BaHPO_4 . Sol. in phosphoric, and certain other acids. (Berzelius, A. ch. 2. 153.)

Barium pyrophosphate, $\text{Ba}_2\text{P}_2\text{O}_7 + x\text{H}_2\text{O}$.

Somewhat sol. in H_2O , in much H_2O , also in $\text{HCl} + \text{Aq}$ or $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ or $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Fleitmann, 1833.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wackenrodt)

hydrogen pyrophosphate, $\text{BaH}_2\text{P}_2\text{O}_7$, $\text{P}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Knorre and Oppelt, B. 21. 773.)

traphosphate, $\text{Ba}_3\text{P}_4\text{O}_{13}$.

in H_2O or acids when strongly heated. in and Henneberg, A. 65. 331.)

indianic pyrophosphate,

InP_2O_7 , $\text{P}_2\text{O}_7 + 5\text{H}_2\text{O}$.

insol. in H_2O . (Rosenheim, B. 585.)

potassium trimetaphosphate,

$\text{P}_3\text{O}_9 + \text{H}_2\text{O}$.

less sol. in H_2O than $\text{NH}_4\text{BaP}_3\text{O}_9$ or P_3O_9 . (Lindbom.)

$\text{HCl} + \text{Aq}$ after ignition.

potassium orthophosphate, BaKPO_4 .

in H_2O . (Ouvrard, A. ch. (6) 16.

P_2O_5 . (de Schulten, C. R. 96. 706.)

sodium dimetaphosphate,

$\text{Na}_2(\text{P}_2\text{O}_6)_2 + 4\text{H}_2\text{O}$.

Al, Dissert. 1880.)

sodium trimetaphosphate, BaNaP_3O_9 , P_3O_9 .

easily sol. in H_2O than $\text{Ba}_3(\text{P}_3\text{O}_9)_2$. acids, unless ignited. (Fleitmann and Arg, A. 65. 314.)

insol. in $\text{HCl} + \text{Aq}$ after ignition by long boiling. When fused it is sol. in $\text{HCl} + \text{Aq}$. (Lindbom, Acta 73. 21.)

sodium orthophosphate, $\text{BaNaPO}_4 + \text{P}_2\text{O}_5$.

de Schulten, C. R. 96. 706.)

attacked by cold, but decomp. by hot HCl . (Williers, C. R. 104. 1103.)

insol. in H_2O . (Quartaroli, C. A. 1911.)

sodium pyrophosphate,

$\text{Na}_4(\text{P}_2\text{O}_7)_2$.

insol. in hot HCl and HNO_3 . (Tammann, Z. 2, (2) 45. 469.)

P_2O_7 , $\text{Na}_4\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$. Completely sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, but not insol. in H_2O or $\text{H} + \text{Aq}$. Easily sol. in HNO_3 or HCl . Insol. in alcohol. (Baer, Pogg. 75.

uranous metaphosphate, UO_2 , BaO ,

in H_2O . (Meyer, A. ch. 1907, (8) 12. 142.)

Barium uranyl orthophosphate,

$\text{Ba}(\text{UO}_2)_2(\text{PO}_4)_2 + 8\text{H}_2\text{O}$.

Min. *Uranocircite*.

Barium phosphate chloride, $3\text{Ba}_3(\text{PO}_4)_2$,

BaCl_2 .

Min. *Barytapatite*. (Deville and Caron, A. ch. (3) 67. 451.)

$4\text{BaH}_4(\text{PO}_4)_2$, BaCl_2 . (Erlenmeyer, J. B. 1857. 145.)

15BaO , $6\text{P}_2\text{O}_5$, $\text{BaCl}_2 + 6\text{H}_2\text{O}$ (?). Sol. in 18,000 pts. cold H_2O . Much more sol. in H_2O containing BaCl_2 , NH_4Cl , and NH_4OH . (Ludwig, Arch. Pharm. (2) 56. 271.)

Bismuth orthophosphate, basic, 2BiPO_4 ,

$3\text{Bi}_2\text{O}_3$.

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. (Cavazzi, Gazz. ch. it. 14. 289.)

Bismuth orthophosphate, BiPO_4 .

Insol. in H_2O or $\text{HNO}_3 + \text{Aq}$. Sl. sol. in NH_4 salts + Aq . (Chancel, C. R. 50. 416.)

Not decomp. by H_2O . Other phosphates of Bi are decomp. by H_2O . (Montmartini, C. C. 1900, II. 1256.)

Not hydrolyzed by hot H_2O ; sl. sol. in $\text{BiCl}_3 + \text{Aq}$; decomp. by boiling alkali. (Caven, J. Soc. Chem. Ind. 1897, 16. 30.)

More sol. in $\text{HCl} + \text{Aq}$ than in $\text{HNO}_3 + \text{Aq}$. (Rose.)

Sol. in $\text{UO}_2(\text{NO}_3)_2 + \text{Aq}$. (M'Curdy, Am. J. Sci. (2) 31. 282.)

Insol. in $\text{MNO}_3 + \text{Aq}$.

Insol. in Bi salts + Aq . (Rose, Pogg. 76. 26.)

Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$, but insol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett, 1837.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. (Kühn.)

+ $3\text{H}_2\text{O}$. Ppt. Decomp. by H_2S or $\text{KOH} + \text{Aq}$. (Vanino, J. pr. 1906, (2) 74. 151.)

Bismuth pyrophosphate, basic, $2\text{Bi}_2\text{O}_3$, P_2O_5 .

Insol. in H_2O and $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$; sol. in hot HCl and $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, and NH_4 citrate + Aq . (Passerini, Cim. 9. 84.)

Bismuth pyrophosphate, $\text{Bi}_4(\text{P}_2\text{O}_7)_3$.

Insol. in H_2O or $\text{HNO}_3 + \text{Aq}$. (Chancel, C. R. 50. 416.)

Decomp. by H_2O . (Wallroth, Bull. Soc. (2) 39. 316.)

Sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Stromeyer.)

Bismuth sodium pyrophosphate, $\text{NaBiP}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Insol. in H_2O . (Rosenheim, B. 1915, 48. 588.)

Boron phosphate, BPO_4 .

Insol. in H_2O . Not attacked by boiling alkalies. (Meyer, B. 22. 2919.)

Bromomolybdenum phosphate.

See under Bromomolybdenum comps.

Cadmium triphosphate, $\text{Cd}_3(\text{P}_3\text{O}_{10})_2$.

Insol. in H_2O and acids. (Glühmann, Dissert. 1899.)

Cadmium tetraphosphate, $6\text{CdO}, 4\text{P}_2\text{O}_5 + 18\text{H}_2\text{O}$.

Insol. in acids. (Glühmann.)

Cadmium metaphosphate.

Very sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Persoz, A. ch. 56. 334.)

Cadmium dimetaphosphate, $\text{Cd}(\text{PO}_3)_2 + 2\text{H}_2\text{O}$.

Sol. in 32 pts. H_2O . Scarcely attacked by acids, especially conc. H_2SO_4 . (Glatzel, Dissert. 1880.)

Cadmium tetrametaphosphate.

Insol. in H_2O . Easily decomp. by $\text{Na}_2\text{S} + \text{Aq.}$ (Fleitmann, Pogg. 78. 358.)

$\text{Cd}_2(\text{PO}_3)_4 + 10\text{H}_2\text{O}$. Not so very difficultly attacked by acids but insol. after ignition. (Glatzel, Dissert. 1880.)

Cadmium orthophosphate, $\text{Cd}_3(\text{PO}_4)_2$.

Ppt. Insol. in H_2O . Sol. in Cd salts + Aq. (Stromeyer.)

Easily sol. in NH_4 sulphate, chloride, nitrate, or succinate + Aq. (Wittstein, Repert. 57. 32.)

$\text{H}_2\text{Cd}_3(\text{PO}_4)_4 + 4\text{H}_2\text{O}$. Sol. in dil. $\text{H}_3\text{PO}_4 + \text{Aq.}$ (de Schulten, Bull. Soc. (3) 1. 473.)

Cadmium tetrahydrogen phosphate, $\text{CdH}_4(\text{PO}_4)_2 + 2\text{H}_2\text{O}$.

Decomp. by great excess of H_2O . (de Schulten.)

Cadmium pyrophosphate, $\text{Cd}_2\text{P}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in NH_4OH , $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq.}$ or acids. Insol. in $\text{KOH} + \text{Aq.}$ Sol. in $\text{SO}_2 + \text{Aq.}$ (Schwarzenberg, A. 65. 183.)

Cadmium hydrogen orthophosphate hydrazine, $\text{CdHPO}_4, 2\text{N}_2\text{H}_4$.

Decomp. by light. (Franzen, Z. anorg. 1908, 60. 283.)

Cadmium potassium tetrametaphosphate, $\text{CdK}_3(\text{PO}_3)_4 + 3\text{H}_2\text{O}$.

Sol. in 135 pts. H_2O . Difficultly decomp. by acids. (Glatzel, Dissert. 1880.)

Cadmium potassium orthophosphate, CdKPO_4 .

Insol. in H_2O ; sol. in dil. $\text{HCl} + \text{Aq.}$ (Ouvrard, A. ch. (6) 16. 321.)

Cadmium potassium pyrophosphate, $\text{CdK}_2\text{P}_2\text{O}_7$.

Insol. in H_2O ; sol. in dil. $\text{HCl} + \text{Ac. rard.}$)

$5\text{Cd}_2\text{P}_2\text{O}_7, 4\text{K}_4\text{P}_2\text{O}_7 + 30\text{H}_2\text{O}$. More easily sol. in H_2O than the CdNa salt. Sv. V. A. F. 30, 7. 39.)

Cadmium sodium triphosphate, $\text{Na}_3\text{Cd}(\text{PO}_3)_3 + 12\text{H}_2\text{O}$.

Sol. in acids even after ignition. (Glühmann, Dissert. 1899.)

Cadmium sodium trimetaphosphate, $\text{Na}_4\text{Cd}(\text{PO}_3)_4 + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O . Insol. in alcohol. Z. anorg. 1901, 28. 204.)

Cadmium sodium tetrametaphosphate, $\text{Na}_2\text{Cd}(\text{PO}_3)_4 + 3\text{H}_2\text{O}$.

Completely insol. in H_2O . (Glatzel, Dissert. 1880.)

Cadmium sodium orthophosphate, $\text{CdNa}_4(\text{PO}_4)_2$.

Insol. in H_2O ; very sol. in dil. ac. CdNaPO_4 . As above. (Ouvrard, A. ch. (6) 16. 321.)

Cadmium sodium pyrophosphate, $\text{CdNa}_2\text{P}_2\text{O}_7$.

Sol. in dil. acids, even acetic acid. (Ouvrard, A. ch. (6) 16. 321.)

$+ 4\text{H}_2\text{O}$. Insol. in H_2O . (Pahl, F. 30, 7. 39.)

Cadmium phosphate bromide, 3CdBr_2 .

Sol. in cold very dil. $\text{HNO}_3 + \text{Aq.}$ (de Schulten, Bull. Soc. (3) 1. 472.)

Cadmium phosphate chloride, 3CdCl_2 .

Sol. in dil. $\text{HNO}_3 + \text{Aq.}$ (de Schulten, Bull. Soc. (3) 1. 472.)

Cæsium metaphosphate, CsPO_3 .

Sol. in H_2O . (von Berg, B. 1901, 30. 183.)

Cæsium orthophosphate, $\text{Cs}_3\text{PO}_4 + 12\text{H}_2\text{O}$.

Deliquescent; very sol. in H_2O . (von Berg, B. 1901, 30. 183.)

Cæsium hydrogen orthophosphate, $\text{CsH}_2\text{PO}_4 + \text{H}_2\text{O}$.

Very sol. in H_2O . (von Berg, B. 1901, 30. 183.)

Cæsium dihydrogen orthophosphate, CsH_2PO_4 .

Sol. in H_2O ; insol. in alcohol. (von Berg, B. 1901, 30. 183.)

Cæsium pyrophosphate, $\text{Cs}_2\text{P}_2\text{O}_7$.

Very sol. in H_2O ; very hygroscopic. (von Berg, B. 1901, 30. 183.)

triphosphate, 5CaO, 3P₂O₅.
in H₂O. (Schwarz, Z. anorg. 1895,

monometaphosphate, Ca(PO₃)₂.
in H₂O and dil. acids. (Maddrell, A.
ecomp. by digestion with alkali car-
+Aq. (Fleitmann.)

dimetaphosphate, Ca₂(P₂O₆)₂+
O.
in H₂O. Decomp. by warm H₂SO₄,
appreciably by conc. HCl or HNO₃+
eitmann, Pogg. 78. 255.)

hexametaphosphate (?).
in H₂O. Sol. in Na₃P₆O₁₃+Aq and
-Aq. (Rose, Pogg. 76. 3.)
O₁₈. Nearly insol. in H₂O; sol. in dil.
Lüder, Z. anorg. 5. 15.)

orthophosphate, basic, 3Ca₃(PO₄)₂+
H₂.
agton, J. B. 1873. 253.)
P₂O₅. (Hilgenstock.)

orthophosphates,
brium in system CaO+P₂O₅+H₂O.

solubility of CaO in P₂O₅+Aq at 25°.

g. P ₂ O ₅ per l. of solution	Solid phase
4.69	CaHPO ₄ , 2H ₂ O
22.39	
23.37	
36.14	
41.24	
59.35	
63.03	
75.95	
79.10	
109.8	
129.8	
139.6	
142.7	
154.6	
191.0	
216.5	
234.6	CaH ₄ (PO ₄) ₂ , H ₂ O
279.7	
351.9	
361.1	
380.3	
395.1	
419.7	
424.6	
428.0	
451.7	
475.3	
505.8	
528.9	
538.3	

n and Seidell, J. Am. Chem. Soc.
1905, 27. 1508.)

Solubility of CaO in P₂O₅+Aq at 25°.

g. CaO per l. of solution	g. P ₂ O ₅ per l. of solution	Solid phase
7.61	19.96	CaHPO ₄
6.51	16.52	
5.01	12.82	
3.42	8.16	
2.42	5.75	
1.58	3.66	
0.544	1.516	Solid phases are evidently solid solutions
0.400	1.108	
0.291	0.773	
0.232	0.662	
0.145	0.381	
0.062	0.109	
0.049	0.088	Ca ₃ (PO ₄) ₂ Solid phase is prob- ably a solid solution
0.034	0.015	
0.587	0.013	
0.789	0.012	

(Cameron and Seidell, J. Am. Chem. Soc.
1905, 27. 1513.)

Solubility of CaO in P₂O₅+Aq at 50.7°.

100 g. of the solution contain		Solid phase
g. P ₂ O ₅ .	g. CaO	
62.01	0.336	CaH ₄ P ₂ O ₆ +CaH ₄ P ₂ O ₆ , H ₂ O
58.08	0.635	CaH ₄ P ₂ O ₆ , H ₂ O
54.67	0.939	"
50.25	1.428	"
46.15	2.100	"
41.92	2.974	"
37.33	3.898	"
33.18	4.880	"
29.61	5.725	CaH ₄ P ₂ O ₆ , H ₂ O+CaHPO ₄
15.48	3.507	CaHPO ₄
9.465	2.328	"
6.157	1.563	"
2.946	0.852	"
2.281	0.692	"
0.1521	0.0588	"
0.1527	0.0596	CaHPO ₄ , 2H ₂ O
0.1331	0.0514	Ca ₃ P ₂ O ₈ , H ₂ O
0.0942	0.0351	"
0.0309	0.0106	"
0.00068	0.00071	"

(Bassett, Z. anorg. 1908, 59. 15.)

Solubility of CaO in P ₂ O ₅ + Aq at 40°.		
100 g. of the solution contain		Solid phase
g. P ₂ O ₅	g. CaO	
45.42	1.768	CaH ₄ P ₂ O ₇ , H ₂ O
41.33	2.588	
36.79	3.584	
32.46	4.505	
28.27	5.501	
21.67	4.813	CaHPO ₄
17.78	4.100	
16.35	3.810	
9.905	2.536	
6.979	1.847	
4.397	1.267	Ca ₃ P ₂ O ₈ , H ₂ O
1.819	0.576	
0.423	0.156	
0.294	0.110	
0.158	0.0592	
0.146	0.0519	Ca ₄ P ₂ O ₉ , 4H ₂ O
0.128	0.0508	
0.0262	0.0098	
trace	0.0709	
"	0.0814	
"	0.0829	"
"	0.0840	

(Bassett, Z. anorg. 1908, 59. 18.)

Solubility of CaO in P ₂ O ₅ + Aq at 25°.		
100 g. the solution contain		Solid phase
g. P ₂ O ₅	g. CaO	
36.11	3.088	CaH ₄ P ₂ O ₇ , H ₂ O
31.97	4.128	
28.34	4.908	
27.99	4.930	
25.45	5.489	
22.90	5.523	CaHPO ₄
17.55	4.499	
15.34	4.027	
9.10	2.638	
6.049	1.878	
3.613	1.181	CaHPO ₄ + CaHPO ₄ , 2H ₂ O
2.387	0.826	
0.417	0.165	
0.178	0.0696	
0.0332	0.0126	
0.0948	0.0352	Probably Ca ₃ P ₂ O ₈ , H ₂ O
0.0571	0.0211	
0.0525	0.0175	
0.0468	0.0186	
trace	0.1131	
	0.118	Ca ₄ P ₂ O ₉ , 4H ₂ O + Ca(OH) ₂ Ca(OH) ₂

(Bassett, Z. anorg. 1908, 59. 20.)

Calcium orthophosphate, Ca₃(PO₄)₂.
Decomp. by long boiling with H₂O into basic salt, 3Ca₃(PO₄)₂, CaO, H₂. This decomp. begins with cold H₂O, so that the solubility at 6–8° varies from 9.9 to 28.6 mg. in a litre. (Warrington, Chem. Soc. (2) 11. 983.)

1 l. cold H₂O dissolves in 7 day ignited, and 79 mg. freshly pr Ca₃(PO₄)₂. (Völcker, J. B. 1882. 1.
100,000 pts. H₂O dissolve 2.36 pt
ous Ca phosphate; 2.56 pts. igniter
phate; 3.00 pts. Ca phosphate from
(Maly and Donath, J. pr. (2) 7. 41
Solubility of bones in various
given by Maly and Donath, l. c.
0.009 g. Ca₃(PO₄)₂ is sol. in
0.153 " " " " "
sat. with CO₂. (Joffre, Bull. Soc
19. 372.)
Determinations of solubility
stated in the literature vary becaus
is apparently a solid solution of C
CaO. When placed in contact
more PO₄ ions dissolve than C
resulting solution is acid and
richer in Ca than before additi
For material of the approximate c
Ca₃(PO₄)₂, the amt. dissolved by C
at ord. temp. is 0.01–0.10 g. per l
on conditions of experiment. H
CO₂ dissolves 0.15–0.30 g. per l.
and Hurst, J. Am. Chem. Soc
903.)
The decomposition of Ca₃(PO₄)₂
increased by presence of CaSO₄; d
presence of CaCO₃ or of CaSO₄ and
increases the amount of PO₄ diss
solution of water alone and the
solutions, but has no other effect
crease the amount of Ca in the
contact with CaCO₃. (Cameron
J. Am. Chem. Soc. 1904, 26. 1458
Sol. in CO₂ + Aq.
1 l. H₂O containing 1 vol. CO₂
12 hours at 10°, 0.75 g. precipitated
0.166 g. Ca₃(PO₄)₂ from bone as
Ca₃(PO₄)₂ from bones which had
20 years. (Lassaigne, J. ch. méd.
1 l. H₂O containing 0.8 vol. C
0.61 g. Ca₃(PO₄)₂. (Liebig, A. 10
H₂O sat. with CO₂ at 5–10° at
pressure dissolves 0.527–0.60 g. Ca
if containing 1% NH₄Cl, 0.739 g.
(Warrington, Chem. Soc. (2) 9. 80
Solubility varies according t
Ca₃(PO₄)₂.
In apatite, 1 pt. Ca₃(PO₄)₂ i
222,222 pts. H₂O sat. with CO₂; in
in 5698 pts.; in bone ash, in 81
So. Carolina phosphate, in 6983 p
phatic guano from Orchilla Id., i
(Williams, C. N. 24. 306.)
Al₂O₃, H₂ and Fe₂O₃, H₂ prevent t
of Ca₃(PO₄)₂ in H₂O containing C
ington, l. c.)
1 l. H₂O dissolves 0.22848 g.
under a CO₂-pressure of 2 atm
(Ehlert, Z. Elektrochem. 1912, 14
Sol. in SO₂ + Aq. forming a liqu
gr. at 9° from freshly precipitated
and of 1.188 sp. gr. from bone ash
Sol. in H₂S + Aq. 1 l. H₂O an

es 190–240 mg. $\text{Ca}_3(\text{PO}_4)_2$. (Béchamp, (4) 16. 241.)
ly sol. in HNO_3 or $\text{HCl} + \text{Aq}$.
pts. very dil. $\text{HCl} + \text{Aq}$ dissolve 198–225
 $\text{Ca}_3(\text{PO}_4)_2$. (Crum, A. 63. 294.)
pts. HCl of 1.153 sp. gr. (containing
 Cl) dissolve at 17° when diluted with:
1 4 7 pts. H_2O ,
45.0 62.3 64.7 pts. $\text{Ca}_3(\text{PO}_4)_2$,
13 16 19 pts. H_2O .
71.9 69.5 69.7 pts. $\text{Ca}_3(\text{PO}_4)_2$.
(Bischof, Schw. J. 67. 39.)

mp. by H_2SO_4 .
pletely decomp. to CaSO_4 and H_3PO_4 .
xture of H_2SO_4 and alcohol.

Solubility in $\text{HNO}_3 + \text{Aq}$.
 $\text{Ca}_3(\text{PO}_4)_2$ dissolves at $16.25\text{--}17.5^\circ$ in
. $\text{HNO}_3 + \text{Aq}$ which contain pts. H_2O
1 pt. HNO_3 (sp. gr. = 1.23).

Aq	Pts. H_2O	Pts. $\text{HNO}_3 + \text{Aq}$	Pts. H_2O
	0	30.64	10.754
	0.827	26.48	13
	3.309	32.14	13.236
	5.791	36.06	15.718
	8.273	127.81	40
	10

(Bischof, 1833.)

sol. in acetic, lactic, malic, and tar-
cids than in HCl or $\text{HNO}_3 + \text{Aq}$.

H_2PO_4 in 100 cc. of $\text{H}_2\text{PO}_4 + \text{Aq}$	G. $\text{Ca}_3(\text{PO}_4)_2$ dissolved by 100 cc. of solvent
5	3.85
10	7.28
15	9.45
20	12.50
25	13.79
30	15.10

(Causse, C. R. 1892, 114. 414.)

small quantities of the salts of the
metals increase the solubility in H_2O .
gne, J. chim. méd. (3) 3. 11.)
e cold H_2O with 2 g. NaCl dissolves
g. $\text{Ca}_3(\text{PO}_4)_2$; with 3 g. NaNO_3 , 33 mg.
 $\text{Ca}_3(\text{PO}_4)_2$. (Liebig.)
e H_2O containing 8.75% NaCl dissolves
mg. $\text{Ca}_3(\text{PO}_4)_2$. (Lassaigne.)
salts have even more effect, especially
 $+ \text{Aq}$, which dissolves $\text{Ca}_3(\text{PO}_4)_2$ in
d; also ammonium nitrate and suc-
(Wittstein.)
 $\text{Ca}_3(\text{PO}_4)_2 + \text{Aq}$ dissolves $\text{Ca}_3(\text{PO}_4)_2$ as
as CaSO_4 . (Liebig, A. 61. 128.)
e H_2O containing 2 g. NaCl dissolves
 1.3° 45.7 mg. $\text{Ca}_3(\text{PO}_4)_2$; 3 g. NaNO_3 ,
33 mg. $\text{Ca}_3(\text{PO}_4)_2$; 2.2 g. $(\text{NH}_4)_2\text{SO}_4$,
g. $\text{Ca}_3(\text{PO}_4)_2$. (Liebig, A. 106. 185.)

Dry $\text{Ca}_3(\text{PO}_4)_2$ also dissolves by long boiling
with solutions of ammonium chloride, nitrate,
succinate (Wittstein), or sulphate (Delkes-
kamp).

Sol. in 89,448 pts. H_2O (boiled) at 7° ;
19,628 pts. H_2O (boiled) containing 1%
 NH_4Cl at 10° ; 4324 pts. H_2O (boiled) con-
taining 10% NH_4Cl at 17° ; 1788 pts. H_2O sat.
with CO_2 and containing 10% NH_4Cl at 10°
and 751 mm. pressure; 1351 pts. H_2O sat.
with CO_2 and containing 1% NH_4Cl at 12°
and 745 mm. pressure; 42,313 pts. H_2O sat.
with CO_2 and containing CaCO_3 at 21° and
756.3 mm. pressure; 18,551 pts. H_2O sat. with
 CO_2 and containing CaCO_3 and 1% NH_4Cl at
 16° and 746.1 mm. pressure. (Warington,
Chem. Soc. (2) 4. 296.)

Aqueous solutions of the following NH_4
salts dissolve the given amts. of $\text{Ca}_3(\text{PO}_4)_2$,
calculated for 100 pts. of the corresponding
acid: NH_4Cl , 0.655 pt.; NH_4NO_3 , 0.306 pt.;
 $(\text{NH}_4)_2\text{SO}_4$, 1.050 pts.; $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$, 0.255 pt.;
 NH_4 tartrate, 4.56 pts.; NH_4 citrate, 7.015
pts.; NH_4 malate, 1.125 pts. $\text{Ca}_3(\text{PO}_4)_2$. (Ter-
reil, Bull. Soc. (2) 35. 578.)

Solubility in various salts + Aq under a CO_2
pressure of 2 atmospheres, at 14° .

Salt	G. salt per 100 g. H_2O	G. $\text{Ca}_3(\text{PO}_4)_2$ sol. in 1 l. of the solvent.
H_2O		0.22848
NaCl	50 conc.	1.3208 0.64089
$\text{MgCl}_2 + 6\text{H}_2\text{O}$	86.9 conc.	1.2873 2.8923
$\text{KMgCl}_2 + 6\text{H}_2\text{O}$	79.2 conc.	1.5771 1.1536
K_2SO_4 , MgSO_4 , $\text{MgCl}_2 + 6\text{H}_2\text{O}$	70.95 conc.	1.7777 2.4911
NaNO_3	72.7 conc.	1.5827 0.8638
K_2SO_4	74.5 conc.	4.9041 4.7649
$(\text{NH}_4)_2\text{SO}_4$	56.5 conc.	2.4131 5.8849
$\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$	137.7 conc.	2.4911 3.2267
$\text{MgSO}_4 + 7\text{H}_2\text{O}$	105.3 conc.	1.9728 3.6001
NH_4Cl	45.74 conc.	1.3710 1.2929

(Ehlert and Hempel, Z. Elektrochem. 1912,
18. 728.)

$\text{Ca}_3(\text{PO}_4)_2$ is sol. in $\text{K}_2\text{C}_2\text{O}_4 + \text{Aq.}$ 100 ccm. $\text{K}_2\text{C}_2\text{O}_4 + \text{Aq}$ (1½% $\text{K}_2\text{C}_2\text{O}_4$) dissolves 57.1% of the P_2O_5 from phosphorite, 71% from guano by boiling 25 min. At ord. temp. bone meal gives up 50–80% of its P_2O_5 to $\text{K}_2\text{C}_2\text{O}_4 + \text{Aq}$ in 36 hours. (Liebig, Landw. J. B. 1881. 603.)
Sol. in Ca sucrate + Aq. (Bobierre, C. R. 32. 859.)
More sol. in H_2O containing starch, glue, or other animal substances than in pure H_2O . (Vauquelin, Pogg. 85. 126.)
Sol. in H_2O containing organic matter, therefore when bones decay under H_2O , $\text{Ca}_3(\text{PO}_4)_2$ is dissolved in considerable quantity. (Hayes, Edin. Phil. J. 5. 378.)
Sol. in sodium citrate + Aq. (Spiller.)
Solubility in NH_4 citrates + Aq.
Ammonium citrate solution of 1.09 sp. gr. at 30–35° dissolves precipitated $\text{Ca}_3(\text{PO}_4)_2$ completely, but not phosphorite. (Frese-
nius.)
Dried on the air, with 2½ H_2O . Sol. in 40 min. in diammonium citrate + Aq (sp. gr. = 1.09); triammonium citrate + Aq (sp. gr. =

1.09) dissolves 55.3% of the P_2O_5 ; citric acid + Aq (¼%) dissolves 83.8% of the P_2O_5 (Erlenmeyer, B. 14. 1253.)
Dried at 50°, with 1½ H_2O . Sol. in 45 min. in diammonium citrate + Aq (sp. gr. = 1.09); triammonium citrate + Aq dissolves 52.3% of the P_2O_5 . (Erlenmeyer.)
Ignited. Diammonium citrate + Aq (sp. gr. 1.09) dissolves 93% of the P_2O_5 ; triammonium citrate + Aq (sp. gr. 1.09) dissolves 32% of the P_2O_5 ; citric acid (¼%) dissolves 53.4% of the P_2O_5 . (Erlenmeyer.)
Insol. in liquid NH_3 . (Franklin, Am. J. 1898, 20. 827.)
Insol. in alcohol and ether.
Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)
Insol. in acetone. (Eidmann, C. C. II. 1014; Naumann, B. 1904, 37. 4329)
Min. Apatite.
0.002 g. is sol. in 1 l. H_2O .
0.014 “ “ “ “ 1 l. H_2O sat. with CO_2 . (Joffre, Bull. Soc. 1898, (3) 19. 374.)

+ H_2O . Solubility in H_2O , in H_2O sat. with CO_2 , and in H_2O containing $\text{CO}_2 + \text{CaH}_2$. Temp. 16°–20°.

Solvent			In 1 l. of the solvent	
			P_2O_5 mg.	C
(1) Boiled distilled H_2O .			0.74	
(2) 1200 cc. distilled $\text{H}_2\text{O} + 50$ cc. H_2O sat. with CO_2 .			6.9	
(3) 1000 cc. “ “ + 250 cc. “ “ “ “			48.5	
(4) 1250 cc. H_2O sat. with CO_2 .			91.9	
Solutions of $\text{CO}_2 + \text{CaH}_2(\text{CO}_3)_2$. 1 l. (filtered) contains:				
(5) { Calcium carbonate	{ Carbonate	13 mg.	0.38	1
{ Bicarbonate	{ Carbonic acid	166 “		
{ Free carbonic acid		73 “		
		9 “		
(6) { Calcium carbonate	{ Carbonate	13 “	1.1	1
{ Bicarbonate	{ Carbonic acid	277 “		
{ Free carbonic acid		122 “		
		49 “		
(7) { Calcium carbonate	{ Carbonate	13 “	0.80	1
{ Bicarbonate	{ Carbonic acid	376 “		
{ Free carbonic acid		165 “		
		105 “		
(8) { Calcium carbonate	{ Carbonate	13 “	1.77	1
{ Bicarbonate	{ Carbonic acid	475 “		
{ Free carbonic acid		209 “		
		206 “		
(9) { Calcium carbonate	{ Carbonate	13 “	1.30	1
{ Bicarbonate	{ Carbonic acid	545 “		
{ Free carbonic acid		240 “		
		301 “		

(Schloesing, C. R. 1900, 131. 151.)

m hydrogen phosphate, CaHPO_4 , and $2\text{H}_2\text{O}$.
l. or nearly so in H_2O . Gradually p. by cold, more quickly by hot H_2O .
pts. H_2O dissolve 0.135–0.152 pt. $\text{O}_4+2\text{H}_2\text{O}$. Solution clouds up on boil-
Birnbaum.)
pts. H_2O dissolve 0.28 pt., and if sat. O_2 , 0.66 pt. $\text{CaHPO}_4+2\text{H}_2\text{O}$. (Dusart
louze.)
n this salt dissolves in H_2O , decomp. place and a very considerable time is
ry to establish equilibrium. (Rindell, 1902, 134. 112.)
h less decomp. by H_2O than $\text{Ca}_3(\text{PO}_4)_2$, $\text{H}_4(\text{PO}_4)_2$, and the decomposition of
lt in water depends only slightly upon
lative amounts of solid and solvent
are present. The decomposition is in-
by the addition of CO_2 . The presence
 O_4 or of CaCO_3 decreased the amount
osphoric acid which dissolved. See
l paper. (Cameron and Seidell, J. Am. Soc. 1904, 26. 1460.)
n the ratio of $\text{P}_2\text{O}_5 : \text{CaO}$ is above 1.0
w 1.27, H_2O dissolves 0.40–0.54 g. CaO
11–1.52 g. P_2O_5 (see original paper).
on and Bell, J. Am. Chem. Soc. 1905, 2.)

Solubility in $\text{H}_3\text{PO}_4+\text{Aq}$.

H_3PO_4 in 100 cc. $\text{H}_3\text{PO}_4+\text{Aq}$	G. CaHPO_4 dissolved by 100 cc. of solvent
5	4.30
10	7.15
15	9.30
20	11.86
25	13.40
30	15.10

(Causse, C. R. 1892, 114. 415.)

H_2O containing 2.2 g. $(\text{NH}_4)_2\text{SO}_4$, 2 g. or 3 g. NaNO_3 dissolves 79.2, 66.3, or g. CaP_2O_7 , which is present in form of O_4 . (Liebig, A. 106. 185.) Slowly but
tely sol. in boiling $\text{NH}_4\text{Cl}+\text{Aq}$.
, Arch. Pharm. (2) 111. 102.) Easily
 $\text{H}_2\text{SO}_4+\text{Aq}$. (Gerland, J. pr. (2) 4.
Very sol. in HCl or HNO_3+Aq . Less
 $\text{HC}_2\text{H}_3\text{O}_2$. (Berzelius.) More sol. in
in conc. $\text{HC}_2\text{H}_3\text{O}_2+\text{Aq}$, but 60 pts.
 O_2 (1 mol.) dissolve at most 23.1 pts.
mol.=142 pts.) from this compound.
is solution of sodium acetate dissolves
asily than H_2O , and becomes turbid
ing. (Birnbaum.)
pletely sol. in $\text{K}_2\text{C}_2\text{O}_4+\text{Aq}$. (Liebig,
J. B. 1881. 603.)
f sat. solution in N/200 acid K tartrate
t 25° contains 0.235 g. CaHPO_4 .
. in alcohol. Sol. in many organic
ces, as starch or gelatine+ Aq .
. in acetone. (Eidmann, C. C. 1899,
t.)

$+1/3\text{H}_2\text{O}$. (Vorbringer, Z. anal. 9. 457.)
 $+\text{H}_2\text{O}$. (Gerlach, J. pr. (2) 4. 104.)
 $+2\text{H}_2\text{O}$. Min. Brushite.
 $+3\text{H}_2\text{O}$. Min. Metabrushite.
 $+5\text{H}_2\text{O}$. (Dusart, C. R. 66. 327.)

Calcium tetrahydrogen orthophosphate,
 $\text{CaH}_4(\text{PO}_4)_2+\text{H}_2\text{O}$.

Very deliquescent. Crystals take up 97.7
pts. H_2O in 16 days, and 226 pts. H_2O in 28
days from air saturated with moisture.
(Birnbaum, Zeit. Ch. (2) 7. 131.)

Not hygroscopic when pure. (Stocklasa,
B. 23. 626 R.)

Completely sol. in 100 pts. H_2O , but de-
comp. by 10–40 pts. H_2O with separation of
 CaHPO_4 , which slowly dissolves. (Erlen-
meyer, J. B. 1873. 254.)

Later (B. 9. 1839) Erlenmeyer says
 $\text{CaH}_4(\text{PO}_4)_2+\text{H}_2\text{O}$ is sol. in 700 pts. H_2O and
decomp. into CaHPO_4 by a less amount of
 H_2O . Wattenberg (Z. anal. 19. 243) says that
the decomposition by small amts. of H_2O
down to 144 pts. H_2O to 1 pt. salt is inappre-
ciable.

Completely sol. in 200 pts. H_2O if pure, and
in less H_2O in presence of H_3PO_4 . (Stocklasa.)

Sol. in 25 pts. H_2O at 15° . Solution begins
to decompose when warmed to 50° . (Otto,
C. C. 1887. 1563.)

Greatly decomp. by H_2O and the resulting
solution is to be regarded as a solution of the
decomposition products rather than of the
substance itself. The presence of an excess
of CaSO_4 does not materially affect the
amount of phosphoric acid entering the solu-
tion. (Cameron, J. Am. Chem. Soc. 1904, 26.
1462.)

Violently decomp. by H_2O in conc. solu-
tion; only sl. decomp. when dissolved in 200
pts. H_2O . (Stocklasa, Z. anorg. 1892, 1. 310.)

Solubility of $\text{CaH}_4\text{P}_2\text{O}_8$ in $\text{H}_3\text{PO}_4+\text{Aq}$ at
pressure of 745 mm. at high temp.

m. pt.	100 g. of the solution contain		Solid phase
	G. P_2O_5	G. CaO	
115°	43.60	5.623	$\text{CaH}_4\text{P}_2\text{O}_8, \text{H}_2\text{O}+\text{CaHPO}_4$
132°	53.43	4.327	$\text{CaH}_4\text{P}_2\text{O}_8+\text{CaH}_4\text{P}_2\text{O}_8, \text{H}_2\text{O}$
169°	63.95	4.489	$\text{CaH}_4\text{P}_2\text{O}_8$

(Bassett, Z. anorg. 1908, 59. 26.)

Glacial $\text{HC}_2\text{H}_3\text{O}_2$ ppt. it completely from
aqueous solution even in presence of HNO_3 .
(Persoz.)

Decomp. by 50 pts. absolute alcohol at b.-
pt. in 1 hour; by 30 pts. in 2 hours. Sol. in
absolute ether. (Erlenmeyer, l. c.)

Calcium pyrophosphate, $\text{Ca}_2\text{P}_2\text{O}_7 + 4\text{H}_2\text{O}$.

Somewhat sol. in H_2O ; completely sol. in mineral acids; less sol. in acetic acid, and insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Schwarzenberg, A. 65. 145.) Less sol. in warm than in cold acetic acid. (Baer, Pogg. 75. 155.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wackenroder, A. 41. 316.)

Insol. in $\text{CaCl}_2 + \text{Aq}$.

Min. *Pyrophosphorite*.

Calcium hydrogen pyrophosphate, $\text{CaH}_2\text{P}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Pahl, B. 7. 478.)

$2\text{CaH}_2\text{P}_2\text{O}_7, \text{Ca}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$. Decomp. by boiling with H_2O into—

$\text{CaH}_2\text{P}_2\text{O}_7, \text{Ca}_2\text{P}_2\text{O}_7 + 3\text{H}_2\text{O}$. Insol. in hot H_2O . (Knorre and Oppelt, B. 21. 771.)

Tetracalcium hydrogen phosphate, $\text{Ca}_4\text{H}(\text{PO}_4)_3 + \text{H}_2\text{O}$.

Ppt. Insol. in H_2O , but decomp. by boiling therewith. Sol. in acids. (Warington, Chem. Soc. (2) 4. 296.)

$+ 2\text{H}_2\text{O}$.

Calcium tetraphosphate, $\text{Ca}_3\text{P}_4\text{O}_{18}$.

Insol. in acids when ignited. (Fleitmann and Henneberg, A. 65. 331.)

Calcium lithium phosphate, CaLiPO_4 .

Insol. in H_2O . (Rose, Pogg. 77. 298.)

Calcium potassium dimetaphosphate, $\text{CaK}_2(\text{P}_2\text{O}_5)_2 + 4\text{H}_2\text{O}$.

As BaK comp. (Glatzel, Dissert. 1880.)

Calcium potassium orthophosphate, CaKPO_4 .

Insol. in H_2O . (Rose, Pogg. 77. 291.)

Easily sol. in acids. (Ouvrard, A. ch. (6) 16. 308.)

Calcium potassium pyrophosphate, $\text{CaK}_2\text{P}_2\text{O}_7$.

Insol. in H_2O ; easily sol. in dil. acids. (Ouvrard, C. R. 106. 1599.)

Calcium sodium dimetaphosphate, $\text{CaNa}_2(\text{P}_2\text{O}_5)_2 + 4\text{H}_2\text{O}$.

As BaNa comp. (Glatzel.)

Calcium sodium trimetaphosphate, $\text{CaNaP}_3\text{O}_9 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Fleitmann, A. 65. 315.)

Easily sol. in H_2O . Difficultly sol. in $\text{HCl} + \text{Aq}$ when heated to redness. Easily sol. in boiling $\text{HCl} + \text{Aq}$ after being fused. (Lindbom.)

Calcium sodium orthophosphate, CaNaPO_4 .

Insol. in H_2O . (Rose, Pogg. 77. 292.)

Easily sol. in dil. acids. (Ouvrard, A. ch. (6) 16. 308.)

$3\text{CaO}, 3\text{Na}_2\text{O}, 2\text{P}_2\text{O}_5$. Sol. in dil. acids. (Ouvrard, C. R. 1888, 106. 1599.)

Calcium sodium pyrophosphate, $\text{CaNa}_2\text{P}_2\text{O}_7 + 4\text{H}_2\text{O}$.

Insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. Easily sol. $\text{HCl} + \text{Aq}$, $\text{HNO}_3 + \text{Aq}$, and also in $\text{HCl} + \text{Aq}$. (Baer, Pogg. 75. 159.)

$\text{Ca}_{10}\text{Na}_{16}(\text{P}_2\text{O}_7)_9$. Sol. in acids. (Wall Bull. Soc. (2) 39. 316.)

$3\text{CaO}, 3\text{Na}_2\text{O}, 2\text{P}_2\text{O}_5$. Easily sol. in (Ouvrard, A. ch. (6) 16. 307.)

Calcium thorium metaphosphate, $\text{ThO}_2\text{P}_2\text{O}_5$.

(Colani, C. R. 1909, 149. 209.)

Calcium uranous metaphosphate, $\text{UO}_2\text{P}_2\text{O}_5$.

Insol. in acids. (Colani, A. ch. 19 12. 140.)

Calcium uranyl phosphate, $\text{Ca}(\text{UO}_2)_2\text{H} + 2, 3, \text{ or } 4\text{H}_2\text{O}$.

Sol. in $\text{HNO}_3 + \text{Aq}$. (Debray.)

$\text{Ca}(\text{UO}_2)_2(\text{PO}_4)_2 + 8\text{H}_2\text{O}$. Min. Sol. in $\text{HNO}_3 + \text{Aq}$.

$3\text{CaO}, 5\text{UO}_3, 2\text{P}_2\text{O}_5 + 16\text{H}_2\text{O}$. (B Dissert. 1900.)

Calcium phosphate chloride, $\text{Ca}_3(\text{PO}_4)_2$.

(Deville and Caron, A. ch. (3) 67. 4

$3\text{Ca}_3(\text{PO}_4)_2, \text{CaCl}_2$. *Chlorapatite*. L H_2O . (Daubrée, Ann. Min. (4) 19. 68

$7\text{CaH}_4(\text{PO}_4)_2, \text{CaCl}_2 + 14\text{H}_2\text{O}$. 8 $\text{HCl} + \text{Aq}$.

$4\text{CaH}_4(\text{PO}_4)_2, \text{CaCl}_2 + 8\text{H}_2\text{O}$.

$\text{CaH}_4(\text{PO}_4)_2, \text{CaCl}_2 + 2\text{H}_2\text{O}$. Partly H_2O with decomp. Also with $8\text{H}_2\text{O}$. meyer, J. B. 1657. 145.)

Calcium phosphate chloride fluoride, $3\text{Ca}_3(\text{PO}_4)_2, \text{CaClF}$.

Min. *Apatite*. Boiling H_2O dissolves CaCl_2 ; dil. mineral acids dissolve easily acid with more difficulty. Easily sol. molten NaCl , crystallizing on cooling. (hammer.)

Calcium phosphate silicate, $\text{Ca}_3(\text{PCa}_2\text{SiO}_4)$.

Insol. in H_2O ; decomp. by HC (Carnot and Richard, C. R. 97. 316.)

$4\text{Ca}_3(\text{PO}_4)_2, \text{Ca}_2\text{SiO}_3$. (Bücking and C. C. 1887. 562.)

$4\text{Ca}_3(\text{PO}_4)_2, 3\text{Ca}_2\text{SiO}_3$. (B. and L.) $\text{Ca}(\text{PO}_3)_2, \text{CaSiO}_3$. (Stead and R. Chem. Soc. 51. 601.)

Calcium dihydrogen phosphate, $\text{CaH}_2(\text{PO}_4)_2, \text{CaSO}_3 + \text{H}_2\text{O}$.

Not decomp. by cold, slowly by H_2O . Slightly sol. in $\text{NH}_4\text{OH} + \text{Aq}$. mineral acids. Insol. in cold, slowly boiling acetic acid. More sol. in a sol. oxalic acid. (Gerland, C. N. 20. 263.)

Metaphosphate, $\text{Ce}(\text{PO}_3)_2$.

(Ramsdellberg.)

$5\text{P}_2\text{O}_5$. Insol. in H_2O or acids. (B. 22. 976.)

Orthophosphate, CePO_4 .

in H_2O . Easily sol. in acids. (A. ch. (6) 8. 193.)

in acids. (Hartley, Proc. Roy. Soc.

Insol. in H_2O . Sol. in acids.

in $\text{H}_3\text{PO}_4 + \text{Aq}$; sl. sol. in HCl or $\text{HNO}_3 + \text{Aq}$. (Hisinger.)

Pyrotholite. Completely decomp. by heat finely powdered. Insol. in dil. $\text{HNO}_3 + \text{Aq}$. (Boussingault, A. ch. (6) 8. 193.)

Pyrotholite. Completely decomp. by heat finely powdered. Insol. in dil. $\text{HNO}_3 + \text{Aq}$. (Boussingault, A. ch. (6) 8. 193.)

Pyrophosphate, $4\text{CeO}_2, 6\text{P}_2\text{O}_5 + 26\text{H}_2\text{O}$. (Hartley, Proc. Roy. Soc. 41. 202.)

Pyrophosphate, $\text{Ce}_2\text{H}_2(\text{P}_2\text{O}_7)_2 + 6\text{H}_2\text{O}$. (Hartley, Proc. Roy. Soc. 41. 202.)

Pyrophosphate, $\text{Ce}_2\text{H}_2(\text{P}_2\text{O}_7)_2 + 6\text{H}_2\text{O}$. (Hartley, Proc. Roy. Soc. 41. 202.)

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Pyrophosphate, $\text{Ce}_2\text{H}_2(\text{P}_2\text{O}_7)_2 + 6\text{H}_2\text{O}$. (Hartley, Proc. Roy. Soc. 41. 202.)

Pyrophosphate, $\text{Cr}_2(\text{PO}_4)_2$.

in H_2O . Easily sol. in citric, tartaric acids. Sl. sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$. (A. ch. 1882, (5) 25. 415.)

Precipitate. Easily sol. in acids. (Moissan, A. ch. (5) 21. 199.)

Pyrophosphate, $\text{Cr}_2(\text{PO}_4)_2$.

in H_2O or conc. acids. (Maddrell, A.

Pyrophosphate, CrPO_4 .

decomp. by hot H_2O . Somewhat sol. in $\text{HCl} + \text{Aq}$ and in $\text{Cr}_2(\text{SO}_4)_3 + \text{Aq}$. (Soc. Chem. Ind. 1897, 16. 29.)

in methyl acetate. (Naumann, B.

1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Chromic phosphate, $\text{Cr}_2(\text{PO}_4)_2 + 12\text{H}_2\text{O}$.

Violet modification. Precipitate. (Ramsdellberg, Pogg. 68. 383.)

+6H₂O. Green modification. Very sl. sol. in H_2O and still less in NH_4NO_3 or $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Carnot, C. R. 94. 1313.)

Insol. in acetic, but easily sol. in mineral acids. Easily sol. in cold KOH or $\text{NaOH} + \text{Aq}$, from which it is separated on boiling. (Dowling and Plunkett, Chem. Gaz. 1858. 220.)

Chromic hydrogen phosphate, $\text{Cr}_2\text{H}_2(\text{PO}_4)_2 + 16\text{H}_2\text{O}$.

Sol. in H_2O . (Haushofer.)

Chromic pyrophosphate, $\text{Cr}_4(\text{P}_2\text{O}_7)_2$.

Anhydrous. Insol. in H_2O or acids. (Ouvrard, A. ch. (6) 16. 344.)

+7H₂O. Precipitate. Sol. in strong mineral acids, $\text{SO}_2 + \text{Aq}$, $\text{KOH} + \text{Aq}$, and $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Schwarzenberg, A. 66. 149.)

Insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Stromeyer.)

Chromic potassium phosphate, $\text{Cr}_2\text{O}_3, \text{K}_2\text{O}, 2\text{P}_2\text{O}_5$.

Insol. in H_2O and in acids. (Ouvrard, A. ch. (6) 16. 289.)

Chromic potassium pyrophosphate, $\text{K}(\text{CrP}_2\text{O}_7) + 5\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . Decomp. by boiling H_2O . (Rosenheim, B. 1915, 48. 586.)

$\text{Cr}_2\text{K}_2\text{H}_4(\text{P}_2\text{O}_7)_2$. Insol. in H_2O , acids, or alkalis. Sl. decomp. by boiling conc. H_2SO_4 . (Schjerning, J. pr. (2) 45. 515.)

Chromic silver phosphate, $2\text{Cr}_2\text{O}_3, 2\text{Ag}_2\text{O}, 5\text{P}_2\text{O}_5$.

(Hautefeuille and Margottet, C. R. 96. 1142.)

Chromic sodium orthophosphate, $\text{Na}_2\text{HPO}_4, 2\text{CrPO}_4 + 5\text{H}_2\text{O}$.

Decomp. by H_2O . (Cohen, J. Am. Chem. Soc. 1907, 29. 1197.)

Chromic sodium pyrophosphate, $\text{Cr}_2\text{Na}_2(\text{P}_2\text{O}_7)_2$.

Insol. in acids. (Wallroth, Bull. Soc. (2) 39. 316.)

+10H₂O, and 16H₂O. Sl. sol. in cold H_2O . Decomp. by boiling H_2O . (Rosenheim, B. 1915, 48. 586.)

Cobaltous monometaphosphate, $\text{Co}(\text{PO}_3)_2(?)$.

Insol. in H_2O and dil. acids. Sol. in conc. $\text{HCl} + \text{Aq}$. (Maddrell, A. 58. 61.)

Cobaltous dimetaphosphate, $\text{Co}_2(\text{P}_2\text{O}_6)_2$.

Insol. in cold conc. H_2SO_4 ; sl. sol. on warming, but sol. in H_2O after treating with H_2SO_4 . Sol. in conc. $\text{NH}_4\text{OH} + \text{Aq.}$ Scarcely attacked by boiling $\text{Na}_2\text{S} + \text{Aq.}$ (Fleitmann.)

Cobaltous hexametaphosphate (?).

Ppt. Sol. in sodium hexametaphosphate + Aq. (Rose, Pogg. 76. 4.)

Cobaltous orthophosphate, $\text{Co}_3(\text{PO}_4)_2 + x\text{H}_2\text{O}$.

Sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$ or $\text{NH}_4\text{OH} + \text{Aq}$; sl. sol. in NH_4Cl or $\text{NH}_4\text{NO}_3 + \text{Aq.}$ (Salvetat, C. R. 48. 295.) Sol. in Co salts + Aq. + $2\text{H}_2\text{O}$. (Debray, A. ch. (3) 61. 438.) + $8\text{H}_2\text{O}$. (Reynoso, C. R. 34. 795.)

Cobaltous hydrogen orthophosphate, $\text{CoHPO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$.

Ppt. (Debray.) + $2\frac{1}{2}\text{H}_2\text{O}$. Ppt. Insol. in H_2O . Sol. in $\text{H}_3\text{PO}_4 + \text{Aq.}$ (Bödeker, A. 94. 357.)

Cobaltous tetrahydrogen orthophosphate, $\text{CoH}_4(\text{PO}_4)_2$.

Sol. in H_2O . (Reynoso.)

Cobaltous pyrophosphate.

Ppt. Sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq.}$ (Stromeyer.) Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Schwarzenberg.)

Cobaltous pyrometaphosphate, $3\text{CoO}, 2\text{P}_2\text{O}_5$.

(Braun.) $6\text{CoO}, 5\text{P}_2\text{O}_5$. (Braun.)

Cobaltous potassium phosphate, CoKPO_4 .

Insol. in H_2O ; easily sol. in dil. acids. (Ouvrard, C. R. 106. 1729.) $3\text{CoO}, 3\text{K}_2\text{O}, 2\text{P}_2\text{O}_5$. As above.

Cobaltous sodium triphosphate, $\text{NaCo}_2\text{P}_3\text{O}_{10}$.

(Schwarz, Z. anorg. 1895, 9. 260.) $\text{Na}_2\text{CoP}_3\text{O}_{10} + 12\text{H}_2\text{O}$. Very sol. in H_2O ; decomp. in aq. solution. Sol. in acids. (Schwarz, Z. anorg. 1895, 9. 258.)

Cobaltous sodium metaphosphate, $\text{Co}_2\text{Na}_2(\text{PO}_3)_4$.

Insol. in H_2O or acids, even conc. H_2SO_4 . (Watts' Dict.)

Cobaltous sodium monometaphosphate, $6\text{Co}(\text{PO}_3)_2, 2\text{NaPO}_3$.

Insol. in H_2O and dil. acids. Sol. in conc. H_2SO_4 . (Maddrell, A. 61. 57.)

Cobaltous sodium trimetaphosphate, $\text{CoNa}_4(\text{PO}_3)_3 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Fleitmann and Henneberg, A. 65. 315.)

Cobaltous sodium orthophosphate, $\text{Co}_3(\text{PO}_4)_2$.

Insol. in H_2O . (Ouvrard, C. R. 106. 1729.) $\text{Co}_3(\text{PO}_4)_2, 2\text{Na}_2\text{HPO}_4 + 8\text{H}_2\text{O}$. (J. Pharm. (3) 46. 119.)

Cobaltous sodium pyrophosphate, $\text{Co}_{10}\text{Na}_{16}(\text{P}_2\text{O}_7)_6$.

Insol. in H_2O . Sol. in acids. (V + $x\text{H}_2\text{O}$. Sol. in H_2O . (Stromeyer.)

Cobaltous zinc phosphate, $\text{Co}_2(\text{PO}_4)_2 + 3\text{Zn}_3(\text{PO}_4)_2 + 12\text{H}_2\text{O}$.

Ppt. Sol. in acids. (Gentile.) $\text{CoZn}_2(\text{PO}_4)_2 + 6\text{H}_2\text{O}$. Insol. in H_2O .

Columbium phosphate (?).

Insol. in H_2O . (Blomstrand.)

Cupric dimetaphosphate, $\text{Cu}_2(\text{P}_2\text{O}_6)_2$.

Insol. in H_2O . Sol. in conc. H_2SO_4 . (drell, A. 61. 62.) Insol. in most acids and in alkalis, except hot NH_4OH . Sol. in conc. H_2SO_4 , in which it is moderately sol. Not decomp. by H_2S , but by $(\text{NH}_4)_2\text{S}$, less easily by Na_2S , and $\text{K}_2\text{S} + \text{Aq.}$ (mann, Pogg. 78. 242.) + $8\text{H}_2\text{O}$. Completely insol. in H_2O . (mann.)

Cupric hexametaphosphate (?).

Sol. in $\text{Na}_6\text{P}_6\text{O}_{18} + \text{Aq}$ or C (Rose, Pogg. 76. 5.) $\text{Cu}_2\text{P}_6\text{O}_{18}$. Easily sol. in H_2O especially when freshly pptd. (J. anorg. 5. 15.)

Cupric orthophosphate, basic, $6\text{CuO}, 3\text{H}_2\text{O}$.

Min. *Phosphocalcite*. $5\text{CuO}, \text{P}_2\text{O}_5 + 2\text{H}_2\text{O}$. Min. *Dibasic*. + $3\text{H}_2\text{O}$. Min. *Eskite*. Easily sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ and $\text{HNO}_3 + \text{Aq.}$ $4\text{CuO}, \text{P}_2\text{O}_5 + \text{H}_2\text{O}$. Slowly sol. or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$; insol. in cold Aq. (Steinschneider, C. C. 1891, 1. 10.) Sl. sol. in $\text{CuCl}_2 + \text{Aq}$ and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ Decomp. by boiling H_2O and potash. (Caven, J. Soc. Chem. 16. 29.)

Min. *Libethenite*. Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ + $2\text{H}_2\text{O}$. Min. *Pseudolibethenite*. Sol. in acids and $\text{NH}_4\text{OH} + \text{Aq.}$ + $3\text{H}_2\text{O}$. Min. *Tagilite*. Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$

Cupric triphosphate, $5\text{CuO}, 3\text{P}_2\text{O}_5$.

Sol. in H_2O . Sol. in HNO_3 . (J. anorg. 1895, 9. 262.)

Cupric dimetaphosphate, CuP_2O_6 .

Sol. in 78 pts. H_2O . Easily sol. in hot conc. H_2SO_4 . (Glatzel, D. 1895, 1. 10.)

phosphate, $\text{Cu}_3(\text{P}_2\text{O}_7)_2 + 9\text{H}_2\text{O}$.
n H_2O (0.04 g. in 1 l. at 20°).
pr. 1892, (2) 45. 425.)

phosphate, $\text{Cu}_2\text{P}_4\text{O}_{12}$.
and in HCl . Sl. sol. in boiling
sol. in boiling conc. H_2SO_4 .

early insol. in H_2O . Slowly
acids except conc. H_2SO_4 .

osphate, $\text{Cu}_3(\text{PO}_4)_2 + 3\text{H}_2\text{O}$.
); easily sol. in acids, even
 O_2 , or $\text{H}_2\text{SO}_3 + \text{Aq}$. Sol. in
Sl. sol. in NH_4 salts + Aq .
salts + Aq . (Rose, Pogg. 76.

$\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Steinschnei-
II, 51.)

id NH_3 . (Gore, Am. Ch. J.

hyl acetate. (Naumann, B.

n phosphate, $\text{CuHPO}_4 +$

); sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$, and
Insol. in NH_4Cl , and
(Brett, Phil. Mag. (3) 10.

osphate, basic, $\text{Cu}_2\text{P}_2\text{O}_7$,
+ $3\text{H}_2\text{O}$.

(Pahl, J. B. 1873. 229.)

phate, $\text{Cu}_2\text{P}_2\text{O}_7$.

Insol. in H_2O , and very sl.
ids. (Fleitmann, Pogg. 78.

Cu metaphosphate, but de-
(Rose, Pogg. 76. 14.)

in mineral acids, and NH_4OH
 $\text{Na}_4\text{F}_2\text{O}_7 + \text{Aq}$. (Schwarzen-

.)
 $\text{I}_2\text{SO}_3 + \text{Aq}$ without decomp.,
on boiling.

oiling $\text{KOH} + \text{Aq}$.

ccess of $\text{CuSO}_4 + \text{Aq}$.

Pahl, Sv. V. A. F. 30, 7. 40.)

y sl. sol. in H_2O .

ds. (Wiesler, Z. anorg. 1901,

rric) pyrophosphate,
 $7)_2 + 12\text{H}_2\text{O}$.

C. R. 1908, 146. 233.)

m phosphate, $4\text{CuO}, \text{K}_2\text{O}$,

(Ouvrard, C. R. 111. 177.)

above.

Cupric potassium tetrametaphosphate,
 $\text{K}_2\text{CuP}_4\text{O}_{12} + 4\text{H}_2\text{O}$.

Sol. in 58 pts. H_2O . Easily attacked by
acids. (Glatzel, Dissert. 1880.)

Cupric potassium pyrophosphate, $\text{CuK}_2\text{P}_2\text{O}_7$.

Extremely easily sol. in H_2O . (Persoz, A.
ch. (3) 20. 315.)

$\text{Cu}_2\text{P}_2\text{O}_7, 3\text{K}_4\text{P}_2\text{O}_7 + 4\text{H}_2\text{O}$. Insol. in H_2O .
(Pahl, Sv. V. A. F. 30, 7. 44.)

Cupric sodium phosphate, $\text{Cu}_2\text{Na}_6(\text{PO}_4)_4$.

Insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in conc.
acids. (Wallroth, Bull. Soc. (2) 39. 316.)

Cupric sodium triphosphate,

$\text{CuNa}_3\text{P}_3\text{O}_{10} + 12\text{H}_2\text{O}$.

Sl. sol. in H_2O ; very unstable.

Easily sol. in acids. (Stange, Z. anorg.
1896, 12. 458.)

Cupric sodium tetrametaphosphate,

$\text{CuNa}_2\text{P}_4\text{O}_{12}$.

As insol. in H_2O as Cu dimetaphosphate.
Difficultly decomp. by digestion with $\text{Na}_2\text{S} +$
 Aq . (Fleitmann, Pogg. 78. 355.)

+ $4\text{H}_2\text{O}$. Sol. in 45 pts. H_2O . (Glatzel,
Dissert. 1880.)

Cupric sodium orthophosphate, $3\text{Cu}_2(\text{PO}_4)_3$,
 NaH_2PO_4 .

Decomp. by H_2O to 4CuO , P_2O_5 . (Stein-
schneider, C. C. 1891, II. 52.)

$2\text{Cu}_2(\text{PO}_4)_3, \text{Na}_2\text{HPO}_4$. Decomp. by H_2O
into—

$3\text{Cu}_2(\text{PO}_4)_3, \text{Na}_2\text{HPO}_4$. Decomp. by H_2O .
(S.)

$\text{Cu}_2(\text{PO}_4)_3, \text{NaH}_2\text{PO}_4$. Decomp. by H_2O .
(S.)

$6\text{Cu}_2(\text{PO}_4)_3, 2\text{Na}_2\text{PO}_4$. Decomp. by H_2O .
(S.)

Cupric sodium pyrophosphate, $\text{CuNa}_2\text{P}_2\text{O}_7$.

Insol. in H_2O . (Fleitmann and Henne-
berg, A. 65. 387.)

+ $\frac{2}{3}\text{H}_2\text{O}$. (F. and H.) Much more sol.
than the next salt. (Pahl.)

+ $6\text{H}_2\text{O}$. (Persoz, A. ch. (3) 20. 315.)

$\text{Cu}_2\text{P}_2\text{O}_7, \text{CuNa}_2\text{P}_2\text{O}_7 + 3\frac{1}{2}\text{H}_2\text{O}$. Very ef-
florescent; insol. in H_2O . (F. and H.)

+ $10\frac{1}{2}\text{H}_2\text{O}$. (Pahl, Sv. V. A. F. 30, 7. 42.
 $\text{CuNa}_2\text{P}_2\text{O}_7, \text{Na}_4\text{P}_2\text{O}_7$. Sol. in H_2O . (F.

and H.)

+ $2\text{H}_2\text{O}$. (F. and H.)

+ 12, and 16 H_2O . Very efflorescent, and
sol. in H_2O . (Pahl.)

$\text{Cu}_2\text{Na}_2\text{P}_4\text{O}_{14} + 10\text{H}_2\text{O}$. Insol. in H_2O ;
sol. in HCl and HNO_3 even after heating.

(Stange, Z. anorg. 1896, 12. 456.)

Cupric uranyl phosphate, $(\text{UO}_2)_2\text{Cu}(\text{PO}_4)_2 +$
 $8\text{H}_2\text{O}$.

Insol. in H_2O ; easily sol. in acids. (De-
bray.)

Min. *Chalcolite*. Sol. in $\text{HNO}_3 + \text{Aq}$.

Cupric orthophosphate ammonia, $\text{Cu}_2(\text{PO}_4)_2 \cdot 4\text{NH}_3$.

Sl. sol. in H_2O . Easily sol. in H_2O containing NH_4OH . (Schiff, A. 123. 41.)

2CuO , $3\text{P}_2\text{O}_5$, $20\text{NH}_3 + 21\text{H}_2\text{O}$. Easily sol. in cold H_2O , with subsequent decomp. (Metzner, A. 149. 66.)

2CuO , P_2O_5 , 6NH_3 . (Maumené.)

Cupric pyrophosphate ammonia, 8CuO , $3\text{P}_2\text{O}_5$, $4\text{NH}_3 + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Schwarzenberg, A. 65. 133.)

$\text{Cu}_2\text{P}_2\text{O}_7$, $4\text{NH}_3 + \text{H}_2\text{O}$. Sl. sol. in H_2O . (Schiff, A. 123. 1.)

Didymium metaphosphate, $\text{Di}(\text{PO}_3)_2$.

Precipitate. (Smith.)

Di_2O_3 , $5\text{P}_2\text{O}_5$. Insol. in H_2O . (Cleve.)

Didymium phosphate, $2\text{Di}_2\text{O}_3$, $3\text{P}_2\text{O}_5$.

Insol. in H_2O . (Ouvrard, C. R. 107. 37.)

Didymium orthophosphate, DiPO_4 .

Insol. in H_2O . Very sl. sol. in dil., easily sol. in conc. acids. (Marignac.) Insol. in H_2O . (Wallroth, Bull. Soc. (2) 39. 316.)

$+ \text{H}_2\text{O}$. (Frerichs and Smith, A. 191. 355.)

Didymium trihydrogen phosphate,

$\text{Di}_2\text{H}_3(\text{PO}_4)_3$.

Precipitate. (Frerichs and Smith.)

Existence is doubtful. (Cleve, B. 12. 910.)

Didymium hexahydrogen phosphate,

$\text{DiH}_6(\text{PO}_4)_3 + \text{H}_2\text{O}$.

Precipitate. (Hermann.)

Didymium pyrophosphate, $\text{Di}_4(\text{P}_2\text{O}_7)_3 + 6\text{H}_2\text{O}$.

Precipitate. (Cleve.)

Didymium hydrogen pyrophosphate,

$\text{Di}_2\text{H}_4(\text{P}_2\text{O}_7)_3$.

Precipitate. Sol. in disodium pyrophosphate + Aq. (Frerichs and Smith, A. 191. 355.)

Does not exist. (Cleve.)

Didymium potassium phosphate, $2\text{Di}_2\text{O}_3$, $3\text{K}_2\text{O}$, $3\text{P}_2\text{O}_5 = 2\text{DiPO}_4$, K_3PO_4 .

Insol. in H_2O . (Ouvrard, C. R. 107. 37.)

Didymium sodium orthophosphate, Di_2O_3 , $3\text{Na}_2\text{O}$, $2\text{P}_2\text{O}_5 = \text{DiPO}_4$, Na_3PO_4 .

Insol. in H_2O . (Ouvrard.)

Didymium sodium pyrophosphate, Di_2O_3 , Na_2O , $2\text{P}_2\text{O}_5 = \text{DiNaP}_2\text{O}_7$.

Insol. in H_2O . (Ouvrard, C. R. 107. 37.)

Dysprosium orthophosphate, DyPO_4 .

Nearly insol. in H_2O .

Easily sol. in dil. acids or ac. (Jantsch, B. 1911, 44. 1276.)

Erbium phosphate, $\text{ErPO}_4 + \text{H}_2\text{O}$.

Precipitate.

Erbium pyrophosphate, ErHP_2O_7 .

Scarcely sol. in boiling H_2O . : in acids.

Erbium sodium pyrophosphate, E.

Precipitate. (Wallroth.)

Glucinum metaphosphate, $\text{Gl}(\text{PO}_3)_2$.

Insol. in H_2O and acids. (Bleyer 1912, 79. 274.)

Glucinum orthophosphate, basic.

$2\text{Gl}_2\text{P}_2\text{O}_7$, $\text{GlO} + 13\text{H}_2\text{O}$.

Ppt. (Bleyer, Z. anorg. 1912, 1)

Glucinum orthophosphate, $\text{Gl}_2(\text{PO}_4)_3$.

Precipitate. Insol. in H_2O . So (Atterberg, Sv. V. A. Handl. 12, 1)

1 l. 2% $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ dissolves the anhydrous salt; 1 l. 10% $\text{HC}_2\text{H}_3\text{O}_2$ dissolves 1.725 g. (Sestini, Gazz. 313.)

$+ 7\text{H}_2\text{O}$. (Atterberg.)

Glucinum hydrogen orthophosphate $+ 3\text{H}_2\text{O}$.

$\text{GlH}_4(\text{PO}_4)_2$ hygroscopic. (Bl anorg. 1912, 79. 266.)

Precipitated by alcohol. (Atte

Glucinum phosphate, 5GlO , $2\text{P}_2\text{O}_5$.

Ppt. Sol. in H_2O with decomp.

3GlO , P_2O_5 , $3\text{H}_2\text{O} + \text{H}_2\text{O}$. (Sch. ch. it. 20. 313.)

Glucinum pyrophosphate, $\text{Gl}_2\text{P}_2\text{O}_7$.

Precipitate. (Scheffer.)

Sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Strom

Glucinum potassium phosphate,

Insol. in H_2O . (Ouvrard, C. R.

Glucinum sodium phosphate, C

Sl. sol. in cold, easily sol. in hot (Wallroth.) Insol. in acetic acid.

Min. *Beryllonite*.

GlO , $2\text{Na}_2\text{O}$, P_2O_5 . Insol. in H rard, C. R. 110. 1333.)

Gold (Auric) sodium pyrophosph

$\text{Au}_4(\text{P}_2\text{O}_7)_3$, $2\text{Na}_4\text{P}_2\text{O}_7 + \text{H}_2\text{O}$.

Sol. in H_2O . (Persoz.)

Iron pyrophosphate, ammonia, $\text{Fe}_2\text{P}_2\text{O}_7, 6\text{P}_2\text{O}_5, 3\text{Na}_2\text{O}, 14\text{NH}_3 + 24\text{H}_2\text{O}$.
 1 H_2O . (Gibbs, Am. Ch. J. 1895,

Iron (ferric) trimetaphosphate, $\text{Fe}(\text{P}_3\text{O}_9)_3 +$
 3 H_2O .

sl. sol. in cold, more easily in hot
 water ignition sol. in $\text{HCl} + \text{Aq}$ only
 boiling. (Lindbom, Acta Lund.

Iron hexametaphosphate, $\text{Fe}_3\text{P}_6\text{O}_{18}$.

Freshly pptd. is sol. in H_2O , and very
 little traces of acids, or $\text{Na}_4\text{P}_6\text{O}_{18} + \text{Aq}$.
 (L. anorg. 5. 15.)

Iron phosphate, basic, $7\text{FeO}, 2\text{P}_2\text{O}_5 +$
 3 H_2O .

Iron pyrochroite. Sol. in dil. H_2SO_4 or
 decomposed by boiling KOH or
 Aq .

Iron orthophosphate, $\text{Fe}_3(\text{PO}_4)_2$.

1 H_2O ; sol. in acids.
 1000 pts. H_2O containing more than
 1 pt. (Pierre.)

in excess of ferrous salts + Aq .

560 pts. H_2O containing $\frac{1}{100}$ pt.

Sol. in 1666 pts. H_2O containing
 1 $\text{H}_4\text{C}_2\text{H}_3\text{O}_7$. (Pierre, A. ch. (3) 36.

NH_4 salts + Aq .

$\text{NH}_4\text{OH} + \text{Aq}$. Not pptd. in pres-
 ence of citrate.

in acetone. (Naumann, B. 1904, 37.

(Debray, A. ch. (3) 61. 437.)

Min. *Vivianite*. Easily sol. in
 $\text{NO}_3 + \text{Aq}$. Boiling $\text{KOH} + \text{Aq}$ dis-
 solves phosphoric acid. Sol. in cold citric
 acid. (Bolton, C. N. 37. 14.)

1 H_2O . Sol. in acids. (Evans, C. C.
 30.)

Iron hydrogen orthophosphate, $\text{FeHPO}_4 +$
 2 H_2O .

Debray, A. ch. (3) 61. 437.)

pure $\text{Fe}_3(\text{PO}_4)_2$. (Erlenmeyer and
 A. 194. 176.)

Iron tetrahydrogen orthophosphate,
 $(\text{PO}_4)_3 + \text{H}_2\text{O}$.

sol. in H_2O . Not changed by al-
 teration. (Erlenmeyer and Heinrichs, A. 194.

Iron pyrophosphate.

sol. in an excess of $\text{Na}_4\text{P}_2\text{O}_7$ or FeSO_4 .
 (Schwarzenberg, A. 65. 153.)

Iron orthophosphate, $\text{Fe}_3(\text{PO}_4)_2$ or $\text{Fe}(\text{PO}_4)_2$.

in H_2O or dil. acids. Sol. in conc.
 acids. (Maddrell, Phil. Mag. (3) 30. 322.)

Iron (ferric) orthophosphate, basic, $2\text{Fe}_2\text{O}_3,$
 $\text{P}_2\text{O}_5 + x\text{H}_2\text{O}$.

Insol. in NH_4 citrate, sol. in NH_4 tartrate
 + Aq . (Wittstein.)

+ 3 H_2O . Min. *Krawurite*. Easily sol. in
 $\text{HCl} + \text{Aq}$.

+ 4 H_2O . Ppt. (Milot, C. R. 82. 89.)

+ 5 H_2O . Min. *Dufrenite*.

+ 12 H_2O . Min. *Cacoxene*. Sol. in $\text{HCl} +$
 Aq .

+ 18, or 24 H_2O . Min. *Delvauxite*.

5 $\text{Fe}_2\text{O}_3, 3\text{P}_2\text{O}_5 + 14\text{H}_2\text{O}$. Min. *Beraunite*.
 Sol. in $\text{HCl} + \text{Aq}$.

3 $\text{Fe}_2\text{O}_3, 2\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}$. Min. *Eleonorite*.
 Sol. in $\text{HCl} + \text{Aq}$.

Ferric orthophosphate, $\text{Fe}_2(\text{PO}_4)_2 + x\text{H}_2\text{O}$, or
 $2\text{Fe}_2\text{O}_3, 3\text{P}_2\text{O}_5 + x\text{H}_2\text{O}$.

+ 4, or 8 H_2O . (Pptd. ferric phosphate.)

Insol. in H_2O . Sol. in 1500 pts. boiling H_2O .
 (Bergmann, 1815.) Sol. in pure H_2O when

all traces of soluble salts are absent. (Frese-
 nius.) Very sl. sol. in, but decomp. by H_2O .

(Lachowicz, W. A. B. 101, 2b. 374.) For an
 extended discussion of solubility in and de-

composition by H_2O and effect of salts see
 Cameron and Hurst, (J. Am. Chem. Soc. 1904,
 26. 888.)

Easily sol. in dil. mineral acids, excepting
 $\text{H}_3\text{PO}_4 + \text{Aq}$. Insol. in cold $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

(Wittstein.) 100 ccm. cold H_2O containing
 10% $\text{HC}_2\text{H}_3\text{O}_2$ dissolve 0.007 g. salt. (See-

tini, Gazz. ch. it. 5. 252.) When freshly pptd.
 easily sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$, or $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$.

(Berthier.) Easily sol. in tartaric or citric
 acid + Aq , also in NH_4 salts of those acids,

and Na citrate + Aq . (Heydenreich, C. N. 4.
 158.) See below.

Sol. in 12,500 pts. H_2O sat. with CO_2 .
 (Pierre, A. ch. (3) 36. 78.)

Insol. in NH_4 salts + Aq . (Wittstein.)
 Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ in presence of Na_2HPO_4 ;

insol. in hot $\text{Na}_2\text{HPO}_4 + \text{Aq}$; sol. in $(\text{NH}_4)_2\text{CO}_3$
 + Aq (Berzelius). NH_4OH , KOH , or

$\text{NaOH} + \text{Aq}$ dissolve out H_3PO_4 .
 Sol. in ferric salts + Aq , even ferric acetate,

but insol. in ferrous acetate + Aq .
 Partially sol. in large amt. of $\text{Na}_2\text{CO}_3 + \text{Aq}$.

Not pptd. in presence of Na citrate. (Spiller.)
 Arth (Bull. Soc. (3) 2. 324) obtained a

modification of $\text{Fe}_2(\text{PO}_4)_2$, insol. in $\text{HNO}_3 +$
 Aq , but sol. in hot conc. $\text{HCl} + \text{Aq}$.

+ 4 H_2O . Min. *Strengite*. Easily sol. in
 $\text{HCl} + \text{Aq}$; insol. in $\text{HNO}_3 + \text{Aq}$.

+ 5 H_2O . Only sl. sol. in H_2O . Slowly sol.
 in HNO_3 , easily sol. in HCl . (Weinland, Z.

anorg. 1913, 84. 361.)

Diammonium citrate + Aq dissolves 4.8%
 of the P_2O_5 ; triammonium citrate, 5.8% P_2O_5 ;

and with an excess of NH_4OH , 21.2% P_2O_5
 is dissolved. (Erlenmeyer, B. 14. 1253.)

+ 9 H_2O . Dissolves in 35 min. in diam-
 monium citrate + Aq (sp. gr. 1.09); in 55

min. in triammonium citrate + Aq (sp. gr.
 1.09); citric acid + Aq ($\frac{1}{4}$ % citric acid) dis-

solves 17.5% of the P_2O_5 . (Erlenmeyer, l. c.)

Iron (ferric) phosphate, acid, $8\text{Fe}_2\text{O}_3, 9\text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$.

Insol. in H_2O . (Rümpfer, Z. anal. 12. 151.)
 $6\text{Fe}_2\text{O}_3, 7\text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$.

$4\text{Fe}_2\text{O}_3, 5\text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$.

$2\text{Fe}_2\text{O}_3, 3\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}$. Ppt. Decomp. by H_2O finally into $\text{Fe}_2(\text{PO}_4)_2$. (Erlenmeyer and Heinrich, A. 194. 176.)

$8\text{Fe}_2\text{O}_3, 11\text{P}_2\text{O}_5 + 9\text{H}_2\text{O}$. As above. (E. and H.)

$4\text{Fe}_2\text{O}_3, 7\text{P}_2\text{O}_5 + 9\text{H}_2\text{O}$. As above. (E. and H.)

$\text{Fe}_2\text{O}_3, 2\text{P}_2\text{O}_5 + 8\text{H}_2\text{O}$. Insol. in H_2O or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$; sol. in NH_4 citrate, alkali hydrates, or carbonates + Aq . (Winkler.) Slowly decomp. by H_2O . (E. and H.)

+ $10\text{H}_2\text{O}$. (Waine, C. N. 36. 132.)

$2\text{Fe}_2\text{O}_3, 5\text{P}_2\text{O}_5 + 17\text{H}_2\text{O}$.

$\text{Fe}_2\text{O}_3, 3\text{P}_2\text{O}_5 + 6\text{H}_2\text{O} = \text{FeH}_2(\text{PO}_4)_2$. Deliquescent. Insol. in H_2O , but decomp. into $\text{Fe}_2(\text{PO}_4)_2$. (E. and H.)

+ $4\text{H}_2\text{O}$. (Hautefeuille and Margottet, C. R. 106. 135.)

Ferric pyrophosphate, $\text{Fe}_4(\text{P}_2\text{O}_7)_3$.

Two modifications.—(a) Sol. in acids, $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, $\text{FeCl}_3 + \text{Aq}$, $\text{NH}_4\text{OH} + \text{Aq}$, and in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.

Insol. in acetic, sulphurous acid, or $\text{NH}_4\text{Cl} + \text{Aq}$. Sol. in NH_4 citrate + Aq . (Schwarzenberg, A. 65. 153.)

(b) Insol. in dil. acids, $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, $\text{FeCl}_3 + \text{Aq}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Gladstone, Chem. Soc. (2) 5. 435.)

Solubility of $\text{Fe}_4(\text{P}_2\text{O}_7)_3$ in $\text{NH}_4\text{OH} + \text{Aq}$ at 0° .

100 g. sat. solution contain		100 g. sat. solution contain	
G. NH_3	G. $\text{Fe}_4(\text{P}_2\text{O}_7)_3$	G. NH_3	G. $\text{Fe}_4(\text{P}_2\text{O}_7)_3$
0.884	5.606	5.92	14.71
1.59	9.75	8.26	13.89
3.71	14.85	10.55	7.40
4.72	15.94	15.96	2.52
5.93	13.92	18.83	0.445
7.91	14.61		

(Pascal, A. ch. 1909, (8) 16. 374.)

Insol. in acetone. (Krug and M'Elroy, J. Anal. Appl. Ch. 6. 184.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Ferroferric orthophosphate, $2\text{Fe}_3(\text{PO}_4)_2, 3(\text{Fe}_2\text{O}_3, 2\text{P}_2\text{O}_5) + 16\text{H}_2\text{O}$.

Ppt. Sol. in $\text{HCl} + \text{Aq}$. (Rammelsberg.)

$4\text{Fe}_2\text{O}_3, 6\text{FeO}, 5\text{P}_2\text{O}_5 + 40\text{H}_2\text{O}$. Sol. in 40 min. in diammonium citrate + Aq (sp. gr. = 1.09); triammonium citrate + Aq (sp. gr. = 1.09) dissolves 55.7% of the P_2O_5 . (Erlenmeyer, B. 14. 1253.)

Ferrous lithium phosphate, $\text{Li}_3\text{PO}_4, \text{Fe}_3(\text{PO}_4)_2$.

Min. *Triphylline*. Easily sol. in acids; not wholly decomp. by $\text{KOH} + \text{Aq}$.

Iron (ferrous) manganous phosphate, $\text{Fe}_3(\text{PO}_4)_2, \text{Mn}_2(\text{PO}_4)_2$.

Min. *Triplite*. Easily sol. in HCl 5(Mn, Fe) O , $2\text{P}_2\text{O}_5 + 5\text{H}_2\text{O}$. *Maulite*. Sol. in acids.

Ferric manganous sodium phosphate, $\text{FePO}_4, (\text{Na}_2, \text{Mn})_2\text{PO}_4 + \frac{1}{2}\text{H}_2\text{O}$. Min.—(?).

Ferrous manganous phosphate, $3(\text{Mn}, \text{Fe})_2(\text{PO}_4)_2, \text{MnCl}_2$. (Deville and Caron.)

Ferrous manganous phosphate f, $(\text{Mn}, \text{Fe})_2(\text{PO}_4)_2, (\text{Mn}, \text{Fe})\text{F}_2$.

Min. *Triplite*, *Zurriesite*. Sol. in 3(Mn, Fe) $_2(\text{PO}_4)_2, \text{MnF}_2$. (D Caron, C. R. 47. 985.)

Ferric potassium phosphate, $2\text{Fe}_2, 3\text{P}_2\text{O}_5$.

Not attacked by boiling H_2O . A. ch. (6) 16. 289.)

$\text{Fe}_2\text{O}_3, \text{K}_2\text{O}, 2\text{P}_2\text{O}_5$. Insol. in H attacked by acids. (Ouvrard.)

Ferric silver metaphosphate, $2\text{Fe}_2, 5\text{P}_2\text{O}_5$.

(Hautefeuille and Margottet, 1142.)

Ferric silver pyrophosphate, $\text{Fe}_2, \text{A} 4\text{H}_2\text{O}$.

Ppt. (Pascal, C. R. 1908, 146.)

Ferric sodium phosphate, $2\text{Fe}_2, 3\text{P}_2\text{O}_5$.

Decomp. by H_2O . (Ouvrard.)

Ferrous sodium triphosphate, $\text{Fe}_2, 11\frac{1}{2}\text{H}_2\text{O}$.

Stable dry; sol. in HNO_3 ; decomp. by H_2O . (Stange, Z. anorg. 451.)

Ferric sodium hydrogen orthophosphate, $\text{Fe}(\text{PO}_4)_2, \text{H}_2\text{Na} + \text{H}_2\text{O}$.

Difficultly sol. in H_2O . Slowly boiling with H_2O . Sol. in dil. HNO_3 . Decomp. by alkalis and carbonates. (Weinland, Z. anorg. 1913, 84. 358.)

$\text{Fe}(\text{PO}_4)_2, \text{H}_2\text{Na} + \text{H}_2\text{O}$. Difficultly sol. in H_2O . Decomp. by boiling with dil. HCl and in dil. HNO_3 . Decomp. by alkalis and alkali carbonates. (V anorg. 1913, 84. 358.)

Ferric sodium pyrophosphate, $2\text{Na}_4\text{P}_2\text{O}_7 + 7\text{H}_2\text{O}$.

Slowly but completely sol. in H_2O by alcohol. (Milck, J. B. 1885, 2.)

Very sol. in H_2O . (Fleitmann berg.)

H_2O . Easily sol. in H_2O , espe-
(Pahl, J. B. 1873. 229.)

Insol. in H_2O , dil. HCl , or
sl. sol. in conc. $\text{HCl} + \text{Aq}$; de-
c. hot H_2SO_4 without solution.
pr. (2) 16. 342.)

etone. (Naumann, B. 1904, 37.

$\text{P}_2\text{O}_5 + 9\text{H}_2\text{O}$. Decomp. by H_2O .
3. 1915, 48. 586.)

$5\text{Na}_4\text{P}_2\text{O}_7 + 7\text{H}_2\text{O}$. (Pahl, J. B.

phosphate sulphate, $3\text{Fe}_2(\text{PO}_4)_3$,
 P_2O_5 , $2\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$.

ite.

etaphosphate, $\text{La}_2(\text{PO}_3)_6$.

(Frerichs and Smith.)

P_2O_5 . Insol. in H_2O , dil., or conc.
son, B. 22. 976.)

thophosphate, LaPO_4 .

(Hermann.)

O and acids. (Ouvrard, C. R.

hydrogen phosphate,

P_2O_5 .

(Frerichs, B. 7. 799.)

doubtful. (Cleve, B. 11. 910.)

osphate, acid, La_2O_3 , $2\text{P}_2\text{O}_5$.

(Hermann.)

rophosphate, $\text{LaHP}_2\text{O}_7 + 3\text{H}_2\text{O}$.

P_2O_5 . Precipitate. (Frerichs and

ist. (Cleve.)

otassium orthophosphate,

K_2O , $3\text{P}_2\text{O}_5 = 2\text{LaPO}_4$, K_3PO_4 .

O . (Ouvrard, C. R. 107. 37.)

odium orthophosphate, La_2O_3 ,

P_2O_5 .

O . (Ouvrard.)

dium pyrophosphate,

7.

tic, and dil. cold mineral acids.
lil. acids. (Wallroth.)

osphate, PbP_2O_6 .

st insol. in H_2O . Sol. in HNO_3 ,
nann, Pogg. 78. 253.)

osphate, $\text{Pb}_3(\text{P}_3\text{O}_{10})_2 + 3\text{H}_2\text{O}$.

l. in H_2O . Less sol. in H_2O
sponding Ag salt. (Fleitmann
g, A. 65. 304.)

. of the trimetaphosphates.
a Lund. 1873. 12.)

Anhydrous salt is insol. in H_2O ; easily sol.
in $\text{HNO}_3 + \text{Aq}$. (Lindbom.)

Lead tetrametaphosphate, $\text{Pb}_4\text{P}_4\text{O}_{12}$.

Insol. in H_2O .

More easily decomp. by acids than the other
insol. metaphosphates. Easily decomp. by
alkali hydrosulphides + Aq in the cold.
(Fleitmann, Pogg. 78. 353.)

Lead hexametaphosphate, $\text{Pb}_6\text{P}_6\text{O}_{18}$.

Nearly insol. in H_2O ; sol. in acids. (Lü-
dert, Z. anorg. 5. 15.)

Lead orthophosphate, basic, 4PbO , P_2O_5 .

(Gerhardt, A. 72. 85.)

Lead orthophosphate, $\text{Pb}_3(\text{PO}_4)_2$.

Insol. in H_2O ; sol. in $\text{HNO}_3 + \text{Aq}$. Insol.
in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

Sl. sol. in H_2O . 1.35×10^{-4} g. is contained
in 1 litre of sat. solution at 20° . (Böttger, Z.
phys. Ch. 1903, 46. 604.)

Not hydrolyzed by boiling H_2O . Sol. in
boiling $\text{KOH} + \text{Aq}$; insol. in $\text{NH}_4\text{OH} + \text{Aq}$.
Insol. in $\text{Pb}(\text{NO}_3)_2 + \text{Aq}$. (Caven, J. Soc.
Chem. Ind. 1897, 16. 30.)

Sol. in 782.9 pts. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ containing
38.94 pts. pure $\text{HC}_2\text{H}_3\text{O}_2$. (Bertrand, Monit.
Scient. (3) 10. 477.)

Insol. in methyl acetate. (Naumann, B.
1909, 42. 3790); ethyl acetate. (Naumann,
B. 1910, 43. 314.)

Lead hydrogen phosphate, PbHPO_4 .

Insol. in H_2O . Decomp. by H_2SO_4 , or HCl
+ Aq . Sol. in HNO_3 , or in KOH or $\text{NaOH} +$
 Aq . Insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in cold
 $\text{NH}_4\text{Cl} + \text{Aq}$ (Brett), from which it can be
completely precipitated by a great excess of
 $\text{NH}_4\text{OH} + \text{Aq}$.

More sol. in $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 + \text{Aq}$ at $18.8-25^\circ$
than in pure H_2O . (Wappen.)

Sol. in sat. $\text{NaCl} + \text{Aq}$, but less than PbSO_4 .
(Becquerel, C. R. 20. 1524.)

Insol. in Pb salts + Aq .

Not pptd. in presence of Na citrate
(Spiller.)

Lead pyrophosphate, $\text{Pb}_2\text{P}_2\text{O}_7 + \text{H}_2\text{O}$.

Insol. in H_2O . Sol. in HNO_3 , or $\text{KOH} + \text{Aq}$.
Insol. in $\text{NH}_4\text{OH} + \text{Aq}$, $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{SO}_2 +$
 Aq . (Schwarzenberg, A. 65. 133.) Sol. in
 $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Stromeyer.)

Insol. in acetone. (Naumann, B. 1904, 37.
4329.)

Lead potassium phosphate, PbKPO_4 .

Decomp. by hot H_2O . (Ouvrard, C. R. 110.
1333.)

Lead sodium phosphate, PbNaPO_4 .

Very sol. in dil. acids. (Ouvrard, C. R.
110. 1333.)

10PbO , $8\text{Na}_2\text{O}$, $9\text{P}_2\text{O}_5$. (Ouvrard.)

Lead sodium pyrophosphate, $\text{PbNa}_2\text{P}_2\text{O}_7$.

Insol. in hot H_2O . (Gerhardt, A. ch. (3) 22. 506.)

Lead triphosphate sodium pyrophosphate, $\text{Pb}_3\text{Na}_4\text{P}_3\text{O}_{17} + 10\text{H}_2\text{O}$.

Sol. in HNO_3 after melting. (Stange, Z. anorg. 1896, 12. 459.)

Lead phosphate chloride, $2\text{PbHPO}_4, \text{PbCl}_2$.

Insol. in boiling H_2O ; sol. in dil. HNO_3 + Aq. (Gerhardt, A. ch. (3) 22. 505.)

$2\text{Pb}_3(\text{PO}_4)_2, \text{PbCl}_2$. Ppt. (Heintz, Pogg. 73. 119.)

$3\text{Pb}_3(\text{PO}_4)_2, \text{PbCl}_2$. Min. *Pyromorphite*. Sol. in HNO_3 , and KOH + Aq.

Sl. sol. in cold citric acid + Aq. (Bolton, C. N. 37. 14.)

+ H_2O . Insol. in H_2O . Sol. in dil. HNO_3 + Aq. (Heintz.)

Lithium metaphosphate, LiPO_3 .

Insol. in boiling H_2O . Scarcely sol. in acetic acid. Easily sol. in mineral acids. (Merling, Z. anal. 1879, 18. 565.)

Lithium tetrametaphosphate, $\text{Li}_4\text{P}_4\text{O}_{12} + 4\text{H}_2\text{O}$.

Very sol. in H_2O . (Warschauer, Z. anorg. 1903, 36. 180.)

Lithium orthophosphate, Li_3PO_4 .

Very slightly sol. in H_2O .

Sol. in 2539 pts. pure H_2O and 3920 pts. ammoniacal H_2O ; much more readily in H_2O containing NH_4 salts. Easily sol. in HCl + Aq or HNO_3 + Aq. (Mayer, A. 98. 193.)

Easily sol. in carbonic acid water. (Troost.) Sol. in dil. acids or acetic acid. (de Schulten, Bull. Soc. (3) 1. 479.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

+ $\frac{1}{2}\text{H}_2\text{O}$, or H_2O .

Lithium hydrogen phosphate, Li_2HPO_4 .

Nearly insol. in H_2O . (Gmelin.) Sol. in 833 pts. H_2O at 12° . (Rammelsberg.)

$\text{Li}_2\text{H}(\text{PO}_4)_2 + \text{H}_2\text{O}$. Sol. in 200 pts. H_2O . (Rammelsberg.)

Lithium dihydrogen phosphate, LiH_2PO_4 .

Deliquescent, and very sol. in H_2O . (Rammelsberg.)

Heptalithium dihydrogen phosphate,

$\text{Li}_7\text{H}_2(\text{PO}_4)_3$.

+ H_2O , or $2\text{H}_2\text{O}$. Sol. in H_2O . (Rammelsberg.)

Lithium pentahydrogen phosphate,

$\text{LiH}_5(\text{PO}_4)_2 + \text{H}_2\text{O}$.

Deliquescent, and sol. in H_2O .

Lithium pyrophosphate, $\text{Li}_2\text{P}_2\text{O}_7 + 2\text{H}_2\text{O}$

(Rammelsberg, B. A. B. 1883. 21.)

Lithium manganous phosphate, $\text{Mn}_2(\text{PO}_4)_2$.

Min. *Lithiophilite*.

Lithium potassium metaphosphate, $2\text{K}_2\text{O}, 3\text{P}_2\text{O}_5 + 4\text{H}_2\text{O}$.

As NH_4 comp. (Tammann, J. (2) 45. 443.)

Lithium potassium pyrophosphate, (Kraut, A. 1876, 182. 170.)

Lithium sodium phosphate, $3\text{Li}_2\text{P}_2\text{O}_5$.

Insol. in H_2O . Sol. in dil. acids. C. R. 110. 1333.)

$2\text{Li}_2\text{O}, \text{Na}_2\text{O}, 2\text{P}_2\text{O}_5$. As above

Lithium sodium pyrophosphate, $\text{Li}_2\text{P}_2\text{O}_5$.

$5\text{Li}_2\text{O}, \text{Na}_2\text{O}, 3\text{P}_2\text{O}_5$.

$4\text{Li}_2\text{O}, 6\text{Na}_2\text{O}, 5\text{P}_2\text{O}_5$. (Kraut, 182. 168.)

Magnesium metaphosphate, $\text{Mg}(\text{PO}_3)_2$.

Insol. in H_2O or dil. acids, but sol + Aq. (Maddrell, A. 61. 62.)

Not decomp. by very long digest alkali carbonates, or orthophosphates (Fleitmann.)

Magnesium dimetaphosphate, $\text{Mg}_2\text{P}_2\text{O}_7$.

Insol. in H_2O ; decomp. by acid (Mann, Pogg. 78. 259.)

Magnesium trimetaphosphate, $\text{Mg}_3\text{P}_3\text{O}_{10}$.

Sl. sol. in cold H_2O , more easily in boiling. When ignited, insol. in boiling (Lindbom.)

Cryst. with 12, or $15\text{H}_2\text{O}$.

Magnesium tetrametaphosphate, $\text{Mg}_4\text{P}_4\text{O}_{13}$.

Insol. in H_2O , somewhat sol. in More easily sol. in HNO_3 + Aq, easily sol. in conc. H_2SO_4 . (Glatz 1880.)

+ $10\text{H}_2\text{O}$. Sol. in 70 pts. H_2O .

Magnesium orthophosphate, $\text{Mg}_3(\text{PO}_4)_2$.

+ 5, or $7\text{H}_2\text{O}$.

1 litre H_2O dissolves 0.1 $\text{Mg}_3(\text{PO}_4)_2$ in 7 days, but 0.205 g precipitated. (Völcker, J. B. 1

1 l. H_2O with 2 g. NaCl dissolve 1 l. H_2O with 3 g. NaNO_3 dissolve

$\text{Mg}_3(\text{PO}_4)_2$. (Liebig, A. 106. 185.)

Easily sol. in acids, except in H_2SO_4 (Schaffner, A. 50. 145.)

Easily sol. in H_2O in presence of

H_2O . Sol. in 30 min. in diammonium Aq (sp. gr. = 1.09); triammonium Aq (sp. gr. = 1.09) dissolves 37.5% O_2 . (Erlenmeyer, B. 14. 1253.)
 O . Sol. in 10 min. in diammonium Aq (sp. gr. = 1.09); triammonium Aq (sp. gr. = 1.09) dissolves 23.2% O_2 ; sol. in 15 min. in $\frac{1}{4}\%$ citric acid Erlenmeyer, l. c.)
 in liquid NH_3 . (Franklin, Am. Ch. 10. 828.)

magnesium hydrogen phosphate, $\text{MgHPO}_4 + \text{H}_2\text{O}$.

322 pts. cold H_2O in several days. to 40° becomes milky, and separates out at 100° of same salt, so that at 100° contains only 1 pt. salt in 498

Much more sol. in H_2O containing acids, even dil. oxalic or acetic acids. (Phil. Mag. Ann. 2. 20.) Easily sol. + Aq. (Gerland, J. pr. (2) 4. 127.) aqueous solution of Mg salts, but $\text{Na}_2\text{HPO}_4 + \text{Aq}$. (Rose.) Sol. in water + Aq. (Spiller.) When freshly precipitated it is sol. in hot $\text{NH}_4\text{Cl} + \text{Aq}$, and + Aq does not completely reprecipitate. Sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett, J. (3) 10. 96.) Insol. in alcohol. (S.)

solubility in H_3PO_4 , see under MgO .

O . (Debray.)

Easily sol. in dil. acids. (de C. R. 100. 263.)

O . Sl. sol. in H_2O , easily in acids. (Z. anorg. 3. 67.)

H_2O . (Bergmann.)

O . (Debray.)

magnesium tetrahydrogen phosphate, $\text{Mg}_3(\text{PO}_4)_2$.

hygroscopic. Sol. in 5 pts. H_2O with- out. (Stoklasa, Z. anorg. 3. 67.)

O . Not hygroscopic. Sol. in H_2O decomp. (Stoklasa, Z. anorg. 1. 307.) precip. by alcohol into $\text{MgHPO}_4 + 3\text{H}_2\text{O}$.

magnesium pyrophosphate, $\text{Mg}_2\text{P}_2\text{O}_7$.

insol. in H_2O ; readily sol. in $\text{HCl} + \text{Aq}$. (Fresenius.)

O . Sl. sol. in H_2O , easily in HCl or Aq ; sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$, and $\text{Na}_4\text{P}_2\text{O}_7$ (Schwarzenberg.)

$\text{MgSO}_4 + \text{Aq}$, and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.

magnesium tetraphosphate, $\text{Mg}_3\text{P}_4\text{O}_{13}$.

in H_2O . (Fleitmann and Henneberg, l.)

magnesium potassium dimetaphosphate, $\text{Mg}_2(\text{P}_2\text{O}_6)_2$.

sol. in dil. acids. (Ouvrard, C. R. 1. 1729.)

O . Sol. in 10.2 pts. H_2O . (Glatzel.)

Magnesium potassium orthophosphate, MgKPO_4 .

Sl. sol. in H_2O . Decomp. by H_2O . Easily sol. in acids.

+ $6\text{H}_2\text{O}$.

2MgO , K_2O , $3\text{P}_2\text{O}_5$. Insol. in H_2O ; sol. in dil. $\text{HCl} + \text{Aq}$. (Ouvrard, C. R. 106. 1729.)

$\text{Mg}_2\text{HK}(\text{PO}_4)_2 + 15\text{H}_2\text{O}$. (Haushofer.)

Magnesium rubidium orthophosphate, $\text{RbMgPO}_4 + 6\text{H}_2\text{O}$.

Easily sol. in warm dil. $\text{HCl} + \text{Aq}$.

Not decomp. by boiling H_2O . (Erdmann, A. 1897, 294. 73.)

Magnesium sodium triphosphate, $\text{MgNa}_3\text{P}_3\text{O}_{10} + 13\text{H}_2\text{O}$.

Decomp. in the air. (Stange, Z. anorg. 1896, 12. 454.)

Magnesium sodium metaphosphate, 3MgO , Na_2O , $4\text{P}_2\text{O}_5$.

Insol. in H_2O or $\text{H}_3\text{PO}_4 + \text{Aq}$. Scarcely sol. in $\text{HCl} + \text{Aq}$, or aqua regia. Not decomp. by $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Sol. in conc. H_2SO_4 . (Maddrell, A. 61. 53.)

Magnesium sodium dimetaphosphate, $\text{MgNa}_2(\text{P}_2\text{O}_6)_2 + 4\text{H}_2\text{O}$.

Sol. in 25 pts. H_2O . (Glatzel, Dissert. 1880.)

Magnesium sodium trimetaphosphate, $\text{MgNa}_4(\text{P}_3\text{O}_9)_2 + 5\text{H}_2\text{O}$.

Sl. sol. in H_2O . After ignition is insol. in H_2O . (Lindbom.)

Magnesium sodium phosphate, 10MgO , $8\text{Na}_2\text{O}$, $9\text{P}_2\text{O}_5$.

Insol. in H_2O ; easily sol. in dil. acids. (Ouvrard, C. R. 106. 1729.)

Magnesium sodium orthophosphate, MgNaPO_4 .

Insol. in H_2O . (Rose.)

+ $9\text{H}_2\text{O}$. (Schoecker and Violet, A. 140. 232.)

MgO , $2\text{Na}_2\text{O}$, P_2O_5 . Insol. in H_2O . (Ouvrard.)

3MgO , $3\text{Na}_2\text{O}$, $2\text{P}_2\text{O}_5$. Insol. in H_2O . (Ouvrard.)

Magnesium sodium pyrophosphate, basic (?).

Precipitate; sl. sol. in H_2O . Easily in $\text{HCl} + \text{Aq}$, $\text{HNO}_3 + \text{Aq}$, and $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Baer, Pogg. 75. 168.)

Sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$, and in $\text{MgSO}_4 + \text{Aq}$.

Insol. in alcohol.

Magnesium phosphate chloride, $\text{Mg}_3(\text{PO}_4)_2$, MgCl_2 .

(Deville and Caron, A. ch. (3) 67. 455.)

Magnesium pyrophosphate nitrogen dioxide,
 $\text{Mg}_2\text{P}_2\text{O}_7, \text{H}_2\text{O}, \text{NO}_2.$

Scarcely sol. in water. (Luck, Z. anal. 13. 255.)

**Magnesium phosphate fluoride, $\text{Mg}_3(\text{PO}_4)_2,$
 $\text{MgF}_2.$**

Min. *Wagnerite*. Slowly sol. in hot HNO_3 , and H_2SO_4 .

Magnesium phosphate calcium fluoride,
 $2\text{Mg}_3(\text{PO}_4)_2, \text{CaF}_2.$

Min. *Kjerulfite*.

Manganous dimetaphosphate, $\text{Mn}_2(\text{P}_2\text{O}_6)_2.$

Anhydrous. Insol. in H_2O and dil. acids. (Fleitmann.) Sol. in conc. H_2SO_4 . (Madrell.) Scarcely attacked by warm $\text{Na}_2\text{S} + \text{Aq}$, and not much more by $(\text{NH}_4)_2\text{S} + \text{Aq}$. Decomp. by $\text{Na}_2\text{CO}_3 + \text{Aq}$.

+ $8\text{H}_2\text{O}$. Insol. in H_2O and dil. acids. (Fleitmann, Pogg. 78. 257.)

Manganous trimetaphosphate, $\text{Mn}_3(\text{P}_3\text{O}_9)_2 + 11\text{H}_2\text{O}.$

Difficultly sol. in cold or warm H_2O . More easily sol. in cold, very easily in warm $\text{HCl} + \text{Aq}$. When ignited, is insol. in acids, even aqua regia. (Lindholm.)

Manganous hexametaphosphate.

Sol. in sodium hexametaphosphate + Aq . (Rose, Pogg. 76. 4.)

$\text{Mn}_3\text{P}_6\text{O}_{18}$. Nearly insol. in H_2O ; easily sol. in acids. (Lüdert, Z. anorg. 5. 15.)

Manganic metaphosphate, $\text{Mn}(\text{PO}_3)_2.$

Insol. in H_2O or acids; decomp. by alkalis. (Schjerning, J. pr. (2) 45. 515.)

Insol. in H_2O ; sol. in HCl ; decomp. by alkalis + Aq . (Barbier, C. R. 1902, 135. 1055.)

+ H_2O . Insol. in H_2O or acids, except $\text{HCl} + \text{Aq}$. Sl. decomp. by boiling with H_2SO_4 . (Hermann, Pogg. 74. 303.)

Manganous tetrametaphosphate, $\text{Mn}_2(\text{PO}_3)_4.$

Not attacked by acids. (Glatzel, Dissert. 1880.)

+ $10\text{H}_2\text{O}$. Sol. only in boiling conc. H_2SO_4 . (Glatzel.)

Manganous dekametaphosphate,

$5\text{MnO}, 5\text{P}_2\text{O}_5 + 12\text{H}_2\text{O}.$

Ppt. (Tammann, J. pr. 1892, (2) 45. 450.)

Manganous orthophosphate, $\text{Mn}_2(\text{PO}_4)_2.$

+ H_2O . (Debray.)

+ $3\text{H}_2\text{O}$. Sol. in 20 min. in diammonium citrate + Aq (sp. gr. = 1.09); triammonium citrate + Aq (sp. gr. = 1.09) dissolves 30.2% of the P_2O_5 . (Erlenmeyer, B. 14. 1253.)

+ $4\frac{1}{2}$ – $5\frac{1}{2}\text{H}_2\text{O}$. Efflorescent. (Meyer and Heinrich, A. 190. 208.)

+ $7\text{H}_2\text{O}$. Very sl. sol. in H_2O . (B.) Easily sol. in mineral acids; sol. in + Aq .

Easily sol. in $\text{SO}_2 + \text{Aq}$. (Gerh. (2) 4. 97.)

Somewhat sol. in boiling $(\text{NH}_4)_2\text{CO}_3$, but deposited on cooling. (Berzel.)

Partly sol. in cold NH_4Cl , or NH_4OH . (Brett.)

Sol. in cold or hot solutions of sulphate or succinate. (Wittstein.)

Sl. sol. in Mn salts + Aq . (Rose 25.)

Insol. in alcohol.

Sol. in 10 min. in diammonium citrate (sp. gr. = 1.09); triammonium citrate (sp. gr. = 1.09) dissolves 53% of (Erlenmeyer, B. 14. 1253.)

Manganous dihydrogen orthophosphate, $\text{MnHPO}_4 + 3\text{H}_2\text{O}.$

Sl. sol. in H_2O . Solution decomp. (Debray.) Slowly decomp. by cold $\text{Mn}_2(\text{PO}_4)_2$. (Erlenmeyer and Heinrich, A. 190. 208.)

Easily sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Gerh.) Sl. sol. in $\text{HC}_2\text{H}_3\text{O}_2$, easily in cold acids. (Heintz.) Sol. in $(\text{NH}_4)_2\text{CO}_3$ from which it is reprecipitated on boiling by boiling $\text{KOH} + \text{Aq}$.

Insol. in alcohol.

$\text{Mn}_2(\text{PO}_4)_2, 2\text{MnHPO}_4 + 4\text{H}_2\text{O}$. (Ten, C. C. 1905, I. 188.)

Manganous tetrahydrogen phosphate, $\text{MnH}_4(\text{PO}_4)_2 + 2\text{H}_2\text{O}.$

Deliquescent. Easily sol. in water. Decomp. to MnHPO_4 . (Erlenmeyer and Heinrich, A. 190. 208.)

Not decomp. by H_2O . (Otto, C. 1563.)

H_2O decomp. it into MnHPO_4 , containing some dissolved salt. H_2O used, the more MnHPO_4 separates. Acid filtrate separates MnHPO_4 . At 0° the decomp. increases in proportion to the amt. of salt, but 1 g. of the salt changed in 100 g. H_2O . With less of salt to 100 g. H_2O the decomp. is to that of $\text{CaH}_4(\text{PO}_4)_2$, but with less of salt it is the opposite, becoming increasing amts. of the salt. (Vissler, 1899, 129. 412.)

Alcohol dissolves out H_2PO_4 . (L.)

Pentamanganous dihydrogen phosphate, $\text{Mn}_5\text{H}_2(\text{PO}_4)_4 + 4\text{H}_2\text{O}.$

Not decomp. by boiling H_2O . (Meyer and Heinrich, A. 190. 208.)

Manganic orthophosphate, basic, $\text{Mn}_2\text{H}_2(\text{PO}_4)_3 + \text{H}_2\text{O}.$

Sl. sol. in H_2O .

orthophosphate, $\text{MnPO}_4 + \text{H}_2\text{O}$.
acids. (Christensen, J. pr. (2)

pyrophosphate, $\text{Mn}_2\text{P}_2\text{O}_7$.
ous. (Lewis, Sill. Am. J. (3) 14.

Insol. in H_2O . Insol. in $\text{MnSO}_4 + \text{Aq}$.
sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Rose.)
ltly sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, but easily
sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Pahl.) Decomp. by
aq. Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Schwar-
n acetone. (Naumann, B. 1904, 37.

hydrogen pyrophosphate,
 $\text{H}_2\text{P}_2\text{O}_7 + 4\text{H}_2\text{O}$.
 H_2O . (Pahl.)

pyrophosphate,
 $\text{P}_6\text{O}_{21} + 14\text{H}_2\text{O}$.
 H_2SO_4 , and H_3PO_4 . (Auger, C. R.
b. 95.)
 P_2O_7 . Insol. in H_2O ; very sl. at-
y dil. $\text{HCl} + \text{Aq}$, easily by conc. Sol.
 H_2SO_4 . (Schjerning, J. pr. (2) 45.

potassium dimetaphosphate,
 $\text{K}_2(\text{PO}_3)_4 + 6\text{H}_2\text{O}$.
95 pts. H_2O . When ignited is not
by acids. (Glatzel, Dissert. 1880.)

potassium orthophosphate,
 KPO_4 .
in H_2O ; easily sol. in dil. acids.
l.)

potassium pyrophosphate,
 KP_2O_7 .
O. Sl. sol. in cold H_2O .
O. Sl. sol. in cold H_2O . (Rosen-
1915, 48. 584.)

potassium phosphate, $\text{MnK}_2\text{P}_2\text{O}_7$.
in H_2O ; sol. in dil. acids. (Ouvrard,
36. 1729.)
O. Sl. sol. in H_2O . (Pahl.)
 P_2O_7 , $2\text{K}_4\text{P}_2\text{O}_7 + 10\text{H}_2\text{O}$. Difficultly
sol. (Pahl.)

potassium pyrophosphate,
 KP_2O_7 .
in H_2O . Decomp. by acids and
(Schjerning.)

silver pyrophosphate,
 $\text{MnP}_2\text{O}_7 + 3\text{H}_2\text{O}$.
st insol. in H_2O . (Rosenheim, B.
. 585.)

Manganous sodium triphosphate,
 $\text{MnNa}_3\text{P}_3\text{O}_{10} + 12\text{H}_2\text{O}$.
Sl. sol. in H_2O ; the melt obtained by heating
the salt is readily sol. in H_2SO_4 . (Stange, Z.
anorg. 1896, 12. 455.)

Manganous sodium dimetaphosphate,
 $\text{MnNa}_2(\text{PO}_3)_4 + 6\text{H}_2\text{O}$.
Easily sol. in boiling H_2SO_4 , but not at-
tacked by acids after boiling. (Glatzel,
Dissert. 1880.)

Manganous sodium trimetaphosphate,
Sol. in H_2O . (Fleitmann and Henneberg.)
 $\text{MnNa}(\text{PO}_3)_3$. Insol. in H_2O , dil. acids, or
alkalies. (Schjerning, J. pr. (2) 45. 515.)

Manganous sodium octometaphosphate,
 $\text{Mn}_3\text{Na}_2(\text{PO}_3)_8$.
Insol. in acids except conc. H_2SO_4 . (Tam-
mann, J. pr. 1892, (2) 45. 469.)
 $+ 5\text{H}_2\text{O}$. Almost insol. in cold H_2O .
Decomp. by boiling H_2O with separation
of Mn_2O_3 . (Rosenheim, B. 1915, 48. 584.)

Manganous sodium orthophosphate,
 MnNaPO_4 .
Insol. in H_2O . (Ouvrard, C. R. 106. 1729.)
 MnO , $2\text{Na}_2\text{O}$, P_2O_5 . As above.

Manganous sodium pyrophosphate,
 $\text{MnNa}_2\text{P}_2\text{O}_7$.
Insol. in H_2O ; easily sol. in dil. acids.
(Wallroth.)
 $+ 4\frac{1}{2}\text{H}_2\text{O}$. Very sl. sol. in H_2O . (Pahl.)
 $3\text{Mn}_2\text{P}_2\text{O}_7$, $2\text{Na}_4\text{P}_2\text{O}_7 + 24\text{H}_2\text{O}$. Very sl.
sol. in H_2O . (Pahl.)

Manganic sodium pyrophosphate, MnNaP_2O_7
 $+ \text{H}_2\text{O}$.
(Christensen, J. pr. (2) 28. 1.)

Manganic dipyrophosphate ammonia,
 $\text{Mn}_2\text{P}_4\text{O}_{14}$, 2NH_3 .
Insol. in H_2O .
Decomp. by HCl and by alkalies. (Bar-
bier, C. R. 1902, 135. 1109.)

Manganous phosphate chloride, $\text{Mn}_3(\text{PO}_4)_2$,
 MnCl_2 .
Insol. in H_2O . (Deville and Caron, A. ch.
(3) 67. 459.)
 $3\text{Mn}_3(\text{PO}_4)_2$, MnCl_2 . Insol. in H_2O . (De-
ville and Caron.)

Mercurous hexametaphosphate (?).
Ppt. Sol. in sodium hexametaphosphate
 $+ \text{Aq}$. (Rose.)
 $\text{Hg}_6\text{P}_6\text{O}_{18}$. Insol. in H_2O ; very sl. sol. in
acids. (Lüdert, Z. anorg. 5. 15.)
Moderately sol. in H_2O when freshly pptd.
More sol. in acids than the mercurous salt.
(Lüdert.)

Mercurous orthophosphate, $(\text{Hg}_2)_2(\text{PO}_4)_2$.

Ppt. Decomp. by boiling with H_2O . (Gerhardt.)

Sol. in $\text{HNO}_3 + \text{Aq}$. Sol. in $\text{Hg}_2(\text{NO}_3)_2 + \text{Aq}$. Insol. in $\text{H}_3\text{PO}_4 + \text{Aq}$.

Mercuric orthophosphate, $\text{Hg}_2(\text{PO}_4)_2$.

Insol. in H_2O . Sl. sol. in hot H_2O , crystallizing out on cooling. (Haack, A. 262. 185. Slowly sol. in cold dil., quickly in hot dil. or cold conc. $\text{HCl} + \text{Aq}$. Less easily sol. in $\text{HNO}_3 + \text{Aq}$. Sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$. (Berzelius.) Insol. in $\text{H}_3\text{PO}_4 + \text{Aq}$. (Haack.) Decomp. by $\text{NaCl} + \text{Aq}$ into insol. HgCl_2 , 3HgO , but sol. in $\text{NaCl} + \text{Aq}$, containing HNO_3 . (Haack.)

Sol. in 6 pts. NH_4Cl in aqueous solution by heating. (Trommsdorff.)

Sol. in $(\text{NH}_4)_2\text{CO}_3$, $(\text{NH}_4)_2\text{SO}_4$, or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Wittstein.)

Insol. in alcohol.

Mercuromercuric orthophosphate, $7\text{Hg}_2\text{O}$, 14HgO , $2\text{P}_2\text{O}_5 + 20\text{H}_2\text{O}$.

(Brooks, Pogg. 66. 63.)

Mercurous pyrophosphate, $\text{Hg}_4\text{P}_2\text{O}_7 + \text{H}_2\text{O}$.

Sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, when recently pptd. Insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, when heated to 100° . Sol. in $\text{HNO}_3 + \text{Aq}$. Decomp. by $\text{HCl} + \text{Aq}$. (Schwarzenberg, A. 65. 133.)

Mercuric pyrophosphate, $\text{Hg}_2\text{P}_2\text{O}_7$.

Sol. in acids; insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$, after being heated to 100° . Sol. in $\text{NaCl} + \text{Aq}$; quickly decomp. by $\text{NaOH} + \text{Aq}$, and $\text{Na}_2\text{HPO}_4 + \text{Aq}$.

Sol. in 6 pts. $\text{NH}_4\text{Cl} + \text{Aq}$. (Trommsdorff.)

Sol. in NH_4NO_3 , $(\text{NH}_4)_2\text{SO}_4$, and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$; also in $\text{KI} + \text{Aq}$.

Mercurous silver orthophosphate, AgHg_2PO_4 .

Sol. in HNO_3 . (Jacobsen, Bull. Soc. 1909, (4) 5. 949.)

Molybdenum phosphate, $\text{Mo}_2(\text{PO}_4)_2$ (?).

Insol. in H_2O . Sol. in $\text{MoCl}_2 + \text{Aq}$.

Molybdenum sodium pyrophosphate, $\text{Na}(\text{MoP}_2\text{O}_7) + 12\text{H}_2\text{O}$.

Ppt. (Rosenheim, B. 1915, 48. 589.)

Nickel dimetaphosphate, NiP_2O_6 .

Insol. in H_2O or dil. acids. Sol. in conc. H_2SO_4 . Not decomp. by boiling alkali carbonates or sulphides + Aq . (Maddrell, A. 61. 58.)

+ $4\text{H}_2\text{O}$. Sol. in cold acids. (Glatzel, Dissert. 1880.)

Nickel tetrametaphosphate, $\text{Ni}_2\text{P}_4\text{O}_{12}$.

Insol. in HCl . Sol. in conc. HNO_3 , and especially sol. in H_2SO_4 on boiling. (Glatzel.)

Nickel tetrametaphosphate, $\text{Ni}_2\text{P}_4\text{O}_{12} + 12\text{H}_2\text{O}$.

Easily sol. in acids. (Glatzel.)

Nickel orthophosphate, $\text{Ni}_3(\text{PO}_4)_2 + 7\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in acids. (Rosenberg, Pogg. 68. 383.)

Sol. in Ni salts + Aq . (Rose, P. 25.)

Insol. in $\text{Na}_2\text{HPO}_4 + \text{Aq}$. (Tupput)

Very sl. sol. in hot $(\text{NH}_4)_2\text{HPO}_4 +$

Insol. in methyl acetate. (Naun 1909, 42. 3790); ethylacetate. (N. B. 1910, 43. 314.)

Nickel pyrophosphate, $\text{Ni}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in mineral acids, + Aq , and $\text{NH}_4\text{OH} + \text{Aq}$. Not ppt. $\text{Ni}_2\text{P}_2\text{O}_7 + \text{Aq}$ by alcohol. (Schwarz, A. 65. 158.)

Nickel potassium dimetaphosphate, $\text{NiK}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$.

Sol. in 130 pts. H_2O . (Glatzel.)

Nickel potassium orthophosphate, $\text{NiK}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in dil. acids. (Rosenberg, A. 65. 1729.)

3NiO , $3\text{K}_2\text{O}$, $2\text{P}_2\text{O}_5$. As above.

Nickel sodium triphosphate, $\text{Na}_3\text{NiP}_3\text{O}_{10} + 12\text{H}_2\text{O}$.

Very sol. in H_2O ; decomp. in Aq . (Schwarz, Z. anorg. 1895, 9. 261.)

Nickel sodium metaphosphate, 3NiNaPO_3 .

Insol. in H_2O and dil. acids. Sol. in H_2SO_4 . (Maddrell, A. 61. 56.)

$\text{NiNa}_4(\text{PO}_3)_3 + 8\text{H}_2\text{O}$. Easily sol. (Lindbom.)

Nickel sodium dimetaphosphate, $\text{NiNa}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$.

Sl. sol. in H_2O . Moderately sol. (Glatzel, Dissert. 1880.)

Nickel sodium trimetaphosphate, $\text{NiNa}_3(\text{P}_3\text{O}_9) + 9\text{H}_2\text{O}$.

1 l. H_2O dissolves 60.6 g. at 20° . (Mann, J. pr. 1892, (2) 45. 426.)

$\text{Na}_3\text{NiP}_3\text{O}_{10} + 12\text{H}_2\text{O}$. Insol. and comp. by H_2O . Sol. in acids. (Schwarz, Z. anorg. 1895, 9. 261.)

Nickel sodium octometaphosphate, $\text{Na}_8\text{Ni}_2(\text{PO}_3)_8$.

(Tammann, J. pr. 1892, (2) 45. 461)

Nickel sodium orthophosphate, $\text{NiNa}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$.

Ppt. (Debray, C. R. 59. 40.)

NiO , $2\text{Na}_2\text{O}$, P_2O_5 . Insol. in H_2O . Sol. in dil. acids. (Ouvrard.)

sodium pyrophosphate,
 $\text{Na}_2\text{P}_2\text{O}_7$.
l. in H_2O . Moderately sol. in acids.
oth.)

m phosphate (?).
sol. in H_2O ; sol. in $\text{HNO}_3 + \text{Aq}$. (Ber-
)

ium orthophosphate (?).

lorus phosphate, $4\text{P}_4\text{O}_{10}$, $3\text{P}_2\text{O}_5$ (?).
omp. spontaneously. Sol. in H_2O and
l when fresh; insol. in ether. (le Verrier,
167; Reinitzer, B. 14. 1884.)

im phosphate, PtP_2O_7 .
l. in H_2O , acids and alkalies. Decomp.
ing with potassium carbonate. (Bar-
N. 1895, 71. 256.)

ium monometaphosphate, KPO_3 .
rly insol. in H_2O ; sol. in weak acids,
1 acetic acid. (Maddrell, A. 61. 62.)
l. in H_2O and weak acids. (Fleitmann,
78. 250.)
l. in liquid NH_3 . (Franklin, Am. Ch.
3, 20. 829.)

ium dimetaphosphate, $\text{K}_2\text{P}_2\text{O}_6 + \text{H}_2\text{O}$.
in 1.2 pts. cold H_2O , but not more in
 O . (Fleitmann, Pogg. 78. 250.)

ium trimetaphosphate, $\text{K}_3\text{P}_3\text{O}_9$.
r sol. in cold H_2O before it is fused.
om, Acta Lund. 1873. 14.)

ium orthophosphate, K_3PO_4 .
deliquescent. Very sol. in H_2O .
am, Pogg. 32. 47.)
r sl. sol. in cold, easily in hot H_2O .
icq.)

Solubility in $\text{H}_3\text{PO}_4 + \text{Aq}$ at 25° .

In 1000 g. of the solution, mols.	
K	PO_4
9.14	3.13
8.84	3.22
8.42	3.44
7.52	3.78
6.90	4.15
6.88	4.12

s and Schreiner, Z. phys. Ch. 1910, 75.
103.)

l. in alcohol.

ium hydrogen orthophosphate,
 K_2HPO_4 , $\text{KH}_2\text{PO}_4 + \text{H}_2\text{O}$.
r unstable; very sol. in H_2O with de-
Identical with the substance de-

scribed as dipotassium phosphate by Ber-
zelius. (Staudenmaier, Z. anorg. 1894, 5.
389.)
 $3\text{K}_2\text{HPO}_4$, $\text{KH}_2\text{PO}_4 + 2\text{H}_2\text{O}$. Very unstable;
very sol. in H_2O with decomp. (Stauden-
maier.)

Potassium hydrogen orthophosphate,
 K_2HPO_4 .
Deliquescent. Very sol. in H_2O and alcohol.

Solubility in $\text{H}_3\text{PO}_4 + \text{Aq}$ at 25° .

In 1000 g. of the solution, mols.	
K	PO_4
6.80	4.08
6.80	4.05
6.76	3.96
6.50	3.81
6.16	3.61
5.24	3.25
4.42	2.94

(D'Ans and Schreiner, Z. phys. Ch. 1910, 75.
103.)

Potassium dihydrogen phosphate, KH_2PO_4 .
Deliquescent. Easily sol. in H_2O . (Vau-
quelin, A. ch. 74. 96.)
1 l. sat. aq. solution at 7° contains 249.9 g.
 KH_2PO_4 . (Muthmann and Kuntze, Z. Kryst.
Min. 1894, 23. 308.)

Solubility in $\text{H}_3\text{PO}_4 + \text{Aq}$ at 25° .

In 1000 g. of the solution, mols.	
K	PO_4
2.90	2.36
1.70	1.71
1.60	1.67
1.48	1.46
1.78	3.15
2.18	4.65
2.54	6.32
2.66	6.76
2.98	8.03
3.32	8.80

(D'Ans and Schreiner, Z. phys. Ch. 1910, 75.
103.)

Sp. gr. of $\text{KH}_2\text{PO}_4 + \text{Aq}$ at 18° containing:
5 10 15% KH_2PO_4 .
1.0341 1.0691 1.1092
(Kohlrausch, W. Ann. 1879. 1.)

Sol. in 20% $\text{KC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Strömeyer.)
For solubility in H_2O , see K_2HPO_4 , H_3PO_4 .
Insol. in alcohol.

Potassium orthophosphate, acid, KH_2PO_4 , H_2PO_4 .

Solubility in H_2O at t° .

t°	$\text{KH}_2\text{PO}_4, \text{H}_2\text{PO}_4$ %	Solid phase
— 0.6	3.337	Ice
— 2.5	12.13	"
— 6.7	29.43	"
— 9.2	36.98	"
— 13	44	Ice + KH_2PO_4
0 (?)	45.8	KH_2PO_4
+10.9	50.3	"
65.2	68.44	"
78	72.43	"
87.5	77.6	"
105.5	85.9	"
120	92.1	" + $\text{KH}_2\text{PO}_4, \text{H}_2\text{PO}_4$
135	96.1	$\text{KH}_2\text{PO}_4, \text{H}_2\text{PO}_4$
139	100	"

(Parravano and Mieli, Gazz. ch. it. 38. II, 536.)

Solubility in anhydrous H_2PO_4 at t° .

t°	% $\text{KH}_2\text{PO}_4, \text{H}_2\text{PO}_4$
38.5	18.17
84	58.42
110	77.53
126.5	92.26

(Parravano and Mieli.)

Potassium pyrophosphate, $\text{K}_2\text{P}_2\text{O}_7 + 3\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O .

Precipitated from aqueous solution by alcohol. (Schwarzenberg, A. 65. 136.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Potassium hydrogen pyrophosphate, $\text{K}_2\text{H}_2\text{P}_2\text{O}_7$.

Very deliquescent, and sol. in H_2O . Insol. in alcohol. (Schwarzenberg.)

Potassium silver metaphosphate, $\text{K}_2\text{Ag}_4(\text{PO}_3)_6 + \text{H}_2\text{O}$.

(Tammann, J. pr. 1892, (2) 45. 417.)

Potassium sodium dimetaphosphate, $\text{KNaP}_2\text{O}_6 + \text{H}_2\text{O}$.

Sol. in 24 pts. H_2O . (Fleitmann, Pogg. 78. 339.)

Potassium sodium phosphate, $\text{KNaHPO}_4 + 7\text{H}_2\text{O}$.

Not efflorescent. Sol. in H_2O .

Tripotassium trisodium hexahydrogen phosphate, $\text{H}_6\text{Na}_3\text{K}_3(\text{PO}_4)_4 + 22\text{H}_2\text{O}$.

Sol. in H_2O . (Filhol and Senderens, C. R. 93. 388.)

Potassium sodium pyrophosphate, $\text{K}_2\text{Na}_2\text{P}_2\text{O}_7 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Schwarzenberg, A. 6

Potassium strontium dimetaphosphate, $\text{K}_2\text{Sr}(\text{P}_2\text{O}_6)_2 + 4\text{H}_2\text{O}$.

As the KBa comp. (Glatzel, Dimer

Potassium strontium hexametaphosphate, $\text{K}_2\text{Sr}_2\text{P}_6\text{O}_{18}$.

(Tammann, J. pr. 1892, (2) 45. 43

Potassium strontium orthophosphate, KSrPO_4 .

Insol. in H_2O ; sol. in dil. acids. (G. A. ch. (6) 8. 193.)

Potassium strontium pyrophosphate, $\text{K}_2\text{SrP}_2\text{O}_7$.

Insol. in H_2O ; sol. in dil. acids. (C. R. 106. 1599.)

Potassium thorium phosphate, $\text{K}_2\text{ThP}_2\text{O}_7$.

Insol. in HCl , HNO_3 , or aq. (Troost and Ouvrard, C. R. 102. 14) K_2O , ThO_2 , P_2O_5 . Insol. in H_2O , HNO_3 + Aq. (Troost and Ouvrard 6 K_2O , 3 ThO_2 , 4 P_2O_5 . Sol. (Troost and Ouvrard.)

Potassium tin (stannic) phosphate, $\text{K}_2\text{SnP}_2\text{O}_7$.

(Ouvrard, C. R. 111. 177.)

K_2O , 2 SnO_2 , P_2O_5 . (Ouvrard.)

Potassium titanium phosphate, $\text{K}_2\text{TiP}_2\text{O}_7$.

(Ouvrard, C. R. 111. 177.)

K_2O , 2 TiO_2 , P_2O_5 . (Ouvrard.)

Potassium uranous phosphate, $\text{K}_2\text{UO}_2\text{P}_2\text{O}_7$.

Practically insol. in conc. HNO_3 even when the acids are boiling. by $\text{HF} + \text{HNO}_3$. (Colani, A. ch. I 133.)

Potassium uranous metaphosphate, K_2O , P_2O_5 .

Easily sol. in HNO_3 . Sol. in (Colani.)

Potassium uranous pyrophosphate, $6\text{K}_2\text{O}$, 4 P_2O_5 .

Sol. in acids. (Colani.)

Potassium uranyl phosphate, K_2UO_4 .

(Ouvrard, C. R. 110. 1333.)

2 K_2O , UO_3 , P_2O_5 . (Ouvrard.)

K_2O , 2 UO_3 , P_2O_5 . (Ouvrard.)

iranyl orthophosphate,
 $\text{PO}_4 + 3\text{H}_2\text{O}$.

mp. (Lienau, Dissert. 1898.)

anadium phosphate.

ovanadate, potassium.

trium phosphate, $3\text{K}_2\text{O}$, Y_2O_3 ,

$2\text{P}_2\text{O}_5$.

O_3 , $6\text{P}_2\text{O}_5$. (Duboin, C. R. 107.

nc tetrametaphosphate,

$\text{O}_3)_4 + 6\text{H}_2\text{O}$.

pts. H_2O . (Glatzel, Dissert.

nc phosphate, KZnPO_4 .

O_2 . Sol. in dil. acids. (Ouvrard, 29.)

As above.

ronium phosphate, K_2O , 4ZrO_2 ,

ids or aqua regia. (Troost and R. 102. 1422.)

P_2O_5 . Insol. in H_2O , HNO_3 , regia. Sol. in hot conc. H_2SO_4 . (Ouvrard.)

osphate selenate.

phosphate, potassium.

ydrogen phosphate sulphate,
 KHSO_4 .

by H_2O and alcohol. (Jacque-

osphate, basic, $4\text{Rh}_2\text{O}_3$, $3\text{P}_2\text{O}_5$ +

O_2 or acids. (Claus.)

$\text{O}_3 + 6\text{H}_2\text{O} = \text{RhPO}_4 + 3\text{H}_2\text{O}$. Sol. (Claus.)

taphosphate, RbPO_3 .

. (von Berg, B. 1901, 34. 4183.)

hophosphate, $\text{Rb}_3\text{PO}_4 + 4\text{H}_2\text{O}$.

ic; sol. in H_2O ; pptd. by alcohol. (von Berg, B. 1901, 34. 4183.)

drogen orthophosphate,

$\text{H}_4 + \text{H}_2\text{O}$.

); insol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$; sol. (von Berg.)

hydrogen orthophosphate,

H_4 .
 H_2O ; pptd. by alcohol. (von

Rubidium pyrophosphate, $\text{Rb}_4\text{P}_2\text{O}_7$.

Hydroscopic; sol. in H_2O . (von Berg.)

Samarium anhydrometaphosphate, Sm_2O_3 ,
 $5\text{P}_2\text{O}_5$.

Insol. in H_2O or $\text{HNO}_3 + \text{Aq}$. (Cleve.)

Samarium orthophosphate, SmPO_4 .

Scarcely attacked by boiling $\text{HNO}_3 + \text{Aq}$. (Cleve.)

+ $2\text{H}_2\text{O}$.

Samarium pyrophosphate, SmHP_2O_7 +
 $1\frac{1}{2}\text{H}_2\text{O}$.

(Cleve.)

Silicon phosphate.

See Silicophosphoric acid.

Silver metaphosphate, Ag_2O , $2\text{P}_2\text{O}_5 + \text{H}_2\text{O}$.

(Kroll, Z. anorg. 1912, 76. 408.)

Silver dimetaphosphate, $\text{Ag}_2\text{P}_2\text{O}_6$.

Very sl. sol. in H_2O . (Fleitmann, Pogg. 78. 253.)

Sol. in cold aniline metaphosphate + Aq . (Nicholson.)

Very sl. sol. in H_2O . (Warschauer, Dissert. 1903.)

+ H_2O . Very sol. in H_2O . (Langheld, B. 1912, 45. 3760.)

Silver trimetaphosphate, $\text{Ag}_3\text{P}_3\text{O}_9$.

Sol. in 60 pts. cold H_2O . Can be crystallized from conc. $\text{HNO}_3 + \text{Aq}$. (Fleitmann and Henneberg.)

+ H_2O . (Lindbom.)

Silver hexametaphosphate, $\text{Ag}_6\text{P}_6\text{O}_{18}$.

Insol. in H_2O . Sol. in HNO_3 or $\text{NH}_4\text{OH} + \text{Aq}$, and in a large excess of sodium hexametaphosphate + Aq . (Rose.)

Easily decomp. by $\text{Na}_2\text{S} + \text{Aq}$.

Decomp. gradually by hot H_2O into $\text{Ag}_3\text{P}_3\text{O}_9$.

When freshly pptd., easily sol. in H_2O . Easily sol. in dil. acids. (Lüder, Z. anorg. 5. 15.)

Silver orthophosphate, Ag_3PO_4 .

Very sl. sol. in H_2O . 1 l. H_2O dissolves 6.5×10^{-3} g. Ag_3PO_4 at 19.46° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

Sol. in H_3PO_4 , HNO_3 , or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, in NH_4OH or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Less easily in ammonium nitrate or succinate, and incompletely in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Lassaigne, J. Pharm. (3) 16. 289.)

Insol. in $\text{Na}_2\text{HPO}_4 + \text{Aq}$. (Stromeyer.)

Not pptd. in presence of Na citrate. (Spiller.)

If 1 mol. Ag_3PO_4 is boiled with 1 mol. Na_2CO_3 , 44% of it is decomp. (Malaguti.)

Readily sol. in soluble hyposulphites + Aq with decomp. (Herschel.)

Insol. in Ag salts + Aq . (Rose.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Silver hydrogen orthophosphate, Ag_3HPO_4 .

Decomp. by H_2O or alcohol into H_3PO_4 and Ag_3PO_4 . (Joly, C. R. 103. 1071.)

Sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$; insol. in ether. (Schwarzenberg, A. 65. 162.)

Silver pyrophosphate, $\text{Ag}_4\text{P}_2\text{O}_7$.

Insol. in hot or cold H_2O . Sol. in cold $\text{HNO}_3 + \text{Aq}$ without decomp. Decomp. by hot HNO_3 or H_2SO_4 into orthophosphate. Decomp. by $\text{HCl} + \text{Aq}$ into AgCl and H_3PO_4 . Insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ without decomp. (Stromeyer, Schw. J. 58. 126.)

Insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. Very sl. sol. in $\text{AgNO}_3 + \text{Aq}$. (Schwarzenberg, A. 65. 161.)

Not completely insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Rose.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Silver hydrogen pyrophosphate, $\text{Ag}_2\text{H}_2\text{P}_2\text{O}_7$.

Decomp. by H_2O into $\text{Ag}_4\text{P}_2\text{O}_7$. (Hurtzig and Geuther, A. 111. 160.)

Decomp. by cold H_2O . (Cavalier, C. R. 1904, 139. 285.)

Silver hydrogen pyrophosphate metaphosphate, $2\text{Ag}_2\text{HP}_2\text{O}_7, \text{HPO}_3$.

Decomp. by H_2O . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (H. and G.)

Silver tetraphosphate, $6\text{Ag}_2\text{O}, 4\text{P}_2\text{O}_5 = \text{Ag}_6\text{P}_4\text{O}_{12}$.

Insol. in, but gradually decomp. by boiling H_2O . (Berzelius.)

Sol. in large excess of the corresponding Na salt + Aq.

Silver dekaphosphate, $\text{Ag}_{12}\text{P}_{10}\text{O}_{31}$.

Easily sol. in sodium dekaphosphate + Aq. (Fleitmann and Henneberg, A. 65. 330.)

Silver ultraphosphate, $\text{Ag}_2\text{O}, 3\text{P}_2\text{O}_5$.

(Kroll, Z. anorg. 1912, 76. 407.)

Silver sodium dimetaphosphate, AgNaP_2O_6 .

Sol. in H_2O . (Fleitmann and Henneberg, Pogg. 65. 310.)

Silver sodium pyrophosphate, $6\text{Ag}_4\text{P}_2\text{O}_7, \text{Na}_4\text{P}_2\text{O}_7 + 4\text{H}_2\text{O}$.

Not completely sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Baer, Pogg. 75. 152.)

Easily sol. in H_2O . (Stromeyer.)

$\text{Ag}_2\text{NaP}_2\text{O}_6 + \frac{1}{2}\text{H}_2\text{O}$. Ppt. (Stange, Z. anorg. 1896, 12. 460.)

Silver uranyl phosphate, $2\text{Ag}_2\text{O}, 6\text{UO}_3, 3\text{P}_2\text{O}_5 + 30\text{H}_2\text{O}$.

(Blinkoff, Dissert. 1900.)

Silver phosphate ammonia, Ag_3PO_4 . (Widmann, B. 17. 2284.)

Sodium triphosphate, $\text{Na}_3\text{P}_3\text{O}_{10}$.

Very sol. in H_2O ; decomp. easily solution at 100° . (Schwarz, Z. anorg. 9. 253.)

Sodium monometaphosphate, NaPO_3 .

Insol. in H_2O . Sol. in dil. and conc. (Maddrell, A. 61. 63.)

Insol. in acids. (Graham.)

Gradually decomp. by alkalis.

Sodium dimetaphosphate, $\text{Na}_2\text{P}_2\text{O}_6$.

Deliquescent. Sol. in 7.2 pts. of H_2O . Very sol. in conc. $\text{HCl} + \text{Aq}$. $\text{NaOH} + \text{Aq}$. Insol. in strong, very dilute alcohol. (Fleitmann, Pogg. 71.

Sodium trimetaphosphate, $\text{Na}_3\text{P}_3\text{O}_{10}$.

Sol. in 4.5 pts. cold H_2O . Insol. very sl. sol. in dil. alcohol. (Fleitmann, Henneberg, A. 65. 307.)

Decomp. by boiling H_2O . (Lind.

Sodium tetrametaphosphate, $\text{Na}_4\text{P}_4\text{O}_{12}$.

Sol. in H_2O ; cryst. with about 4H₂O. Sol. in alcohol than in H_2O . (Fleitmann, Pogg. 78. 854.)

Sodium hexametaphosphate, $\text{Na}_6\text{P}_6\text{O}_{18}$.

Deliquescent. Very sol. in H_2O . Sol. in alcohol. (Graham, Pogg. 32. 56.)

Sodium orthophosphate, Na_3PO_4 .

Not deliquescent in dry air.

100 pts. H_2O dissolve 19.6 pts. crys. (Graham.)

100 pts. H_2O dissolve 28.3 pts. $12\text{H}_2\text{O}$ at 15° . (Schiff.)

Solubility in $\text{H}_3\text{PO}_4 + \text{Aq}$ at

In 1000 g. of the solution. mo

Na	P
4.28	0.0
3.24	0.1
2.24	0.7
2.73	1.0

(D'Ans and Schreiner, Z. phys. Cl. 101.)

Sp. gr. of $\text{Na}_3\text{PO}_4 + \text{Aq}$ at
 $\% = \% \text{Na}_3\text{PO}_4 + 12\text{H}_2\text{O}$

%	Sp. gr.	%	Sp. gr.	%
1	1.0043	9	1.0399	17
2	1.0086	10	1.0455	18
3	1.0130	11	1.0492	19
4	1.0174	12	1.0539	20
5	1.0218	13	1.0586	21
6	1.0263	14	1.0633	22
7	1.0308	15	1.0681	23
8	1.0353	16	1.0729	24

(Schiff, calculated by Gerlach, Z. anorg. 1900.)

sol. in CS_2 . (Arctowski, Z. anorg. 1894, 7.)

sol. in methyl acetate. (Naumann, B. 42. 3790.)

(OH_2O . (Rammelsberg.)

could not be obtained. (Hall, J. pr. 94.

(H_2O . (Hall.)

alts in crystal water at 76.6° . (Graham.)

am hydrogen phosphate, Na_2HPO_4 .

l. in H_2O with evolution of heat.

100 pts. H_2O dissolve at t° .

Pts. Na_2HPO_4	t°	Pts. Na_2HPO_4	t°	Pts. Na_2HPO_4
1.55	40	30.88	80	81.29
4.10	50	43.31	90	95.02
11.08	60	55.29	100	108.30
19.95	70	68.72	106.2	114.43

(Poggiale, J. Pharm. (3) 44. 273.)

pts. H_2O at 13° dissolve 3.4 pts. Na_2HPO_4 (M. Ph. Viertelj. 7. 244); at 15° , 5.9 pts. (Neese); 6.3 pts. (Mulder); at 16° , 8.4 pts. (Möller, J. l. 52); at 20° , 6.8 pts. (Neese, Russ. Z. Pharm. 1. at 25° , 12.5 pts. (ibid.).

Solubility in 100 pts. H_2O at t° .

Pts. Na_2HPO_4	t°	Pts. Na_2HPO_4	t°	Pts. Na_2HPO_4
2.5	35	40.8	69	94.8
2.6	36	43.6	70	95.0
2.6	37	49.5	71	95.1
2.7	38	55.5	72	95.2
2.7	39	60.6	73	95.4
2.8	40	63.9	74	95.6
3.0	41	65.2	75	95.8
3.2	42	68.6	76	96.0
3.4	43	70.8	77	96.1
3.6	44	72.9	78	96.3
3.9	45	74.8	79	96.5
4.2	46	76.5	80	96.6
4.5	47	78.2	81	96.8
4.9	48	79.7	82	96.9
5.3	49	81.2	83	97.0
5.8	50	82.5	84	97.1
6.3	51	83.7	85	97.2
6.9	52	84.8	86	97.4
7.6	53	85.8	87	97.5
8.4	54	86.7	88	97.6
9.3	55	87.7	89	97.7
10.3	56	88.6	90	97.8
11.4	57	89.4	91	97.9
12.6	58	90.2	92	98.0
14.0	59	91.0	93	98.1
15.4	60	91.6	94	98.2
16.9	61	92.2	95	98.4
18.5	62	92.7	96	98.5
20.2	63	93.1	97	98.6
22.0	64	93.5	98	98.7
24.1	65	93.8	99	98.8
26.4	66	94.1	105	82.5
29.1	67	94.4	105.57	80.7
32.1	68	94.6	106.4	79.2
NA. 1

older, Schell. Verhandl. 1864. 103.)

Solubility in H_2O at t° .

t°	G. Na_2HPO_4 in 100 g. H_2O
10.26	3.55
25.15	12.02
40.29	54.88
60.23	83.00
99.77	102.15

Three breaks in the curve: at 36.45° , transition from dodecahydrate to heptahydrate; at 48° , transition from heptahydrate to the dihydrate; at 95.2° , transition from dihydrate to the monohydrate. (Shiomi, C. C. 1909, II. 106.)

Solubility of Na_2HPO_4 in H_2O at t° .

t°	G. Na_2HPO_4 in 100 g. H_2O	Solid phase
-0.43	1.42	Ice
-0.24	0.70	"
-0.5*		
+0.05	1.67	$\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$
20.0	7.68	"
25.0	12.0	"
32.0	25.7	"
34.0	33.8	"
35.2*		
39.2	51.8	$\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$
45.0	67.3	"
48.3*		
50.0	80.2	$\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$
60.0		"
82.9		"
90.0	92.4	"
95*	101.0	"
96.2	104.6	Na_2HPO_4
105.0	102.3	"
120	99.2	"

(Menzies and Humphery, Int. Cong. App. Chem. 1912, 2. 177.)

* Transition points.

Solubility in H_2O at t° .

t°	100 g. H_2O dissolve g. Na_2HPO_4	Solid phase
0°	2.51	$\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$
25	12.47	"
35.4*	46.11	$\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$
40.3	54.80	"
48.35*	79.00	$\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$
59.7	91.3	
71	95.1	
91	98.15	

(D'Ans and Schreiner, Z. phys. Ch. 1911, 75. 99.)

* Transition points.

The composition of the hydrates formed by this salt at different dilutions is calculated.

from determinations of the lowering of the fr-pt. produced by the salt and of the conductivity and sp. gr. of its aqueous solutions. (Jones, Am. Ch. J. 1905, 34. 318.)

Na₂HPO₄+Aq saturated at 15° has 1.0469 sp. gr. (Michel and Krafft); saturated at 16°, 1.0511 (Stolba).

Sp. gr. of Na₂HPO₄+Aq at 19°.

% Na ₂ HPO ₄ +12H ₂ O	Sp. gr.	% Na ₂ HPO ₄ +12H ₂ O	Sp. gr.	% Na ₂ HPO ₄ +12H ₂ O	Sp. gr.
1	1.0041	5	1.0208	9	1.0376
2	1.0083	6	1.0250	10	1.0418
3	1.0125	7	1.0292	11	1.0460
4	1.0166	8	1.0332	12	1.0503

(Schiff, A. 110. 70.)

Saturated solution freezes at -0.45° (Rüdorff, Pogg. 122. 337), and boils at 105° (Griffiths), 105-106.4° (Mulder), 106.5° (Legrand).

Sat. Na₂HPO₄+Aq boils at 105.5° (Griffiths); at 106.5°, and contains 113.2 pts. Na₂HPO₄ to 100 pts. H₂O (Legrand); forms a crust at 106.4°, and contains 108.8 pts. Na₂HPO₄ to 100 pts. H₂O; highest temp. observed, 106.8°. (Gerlach, Z. anal. 26. 427.)

B.-pt. of Na₂HPO₄+Aq containing pts. Na₂HPO₄ to 100 pts. H₂O. G=according to Gerlach (Z. anal. 26. 450); L=according to Legrand (A. ch. (2) 59. 426.)

B.-pt.	G	L	B.-pt.	G	L
100.5°	8.6	11.0	104°	68.4	76.4
101	17.2	21.0	104.5	76.9	84.2
101.5	25.8	31.0	105	85.3	91.5
102	34.4	40.8	105.5	93.7	98.4
102.5	42.9	50.3	106	102.1	105.0
103	51.4	59.4	106.5	110.5	111.4
103.5	59.9	68.1	106.6	...	112.6

+H₂O. Transition point, 95.2°. (Shiomi.)
+2H₂O. Transition point, 48.35°. (D'Ans and Schreiner); 48° (Shiomi); 48.3° (Menzies and Humphery).

+7H₂O. Not efflorescent. Sol. in H₂O with absorption of heat.

Sol. in 8 pts. H₂O at 23°. (Neese, J. B. 1863. 181.)

Transition point, 35.4° (D'Ans and Schreiner); 36.45° (Shiomi); 35.2° (Menzies and Humphery).

Solubility in H₂PO₄+Aq at 25°.

In 1000 g. of the solution, mol.	
Na	PO ₄
6.31	4.63
6.76	4.88
7.31	5.55

(D'Ans and Schreiner, Z. phys. Ch. 1910, 101.)

See also above.

+12H₂O. Efflorescent. Sol. in H₂O with absorption of heat.

14 pts. Na₂HPO₄+12H₂O mixed with 1 pts. H₂O at 10.8° lower the temperature (Rüdorff, B. 2. 68.)

Sol. in 8.48 pts. H₂O at 17°, or 100 pts. H₂O in 11.8 pts. at 17°, and solution has sp. gr. = 1.042. (Schiff.)

Sol. in 4 pts. cold, and 2 pts. boiling H₂O. (Pogg.)

Sol. in 4 pts. H₂O at 18.75°. (Abt.)

100 pts. H₂O dissolve 12.735 pts. Na₂HPO₄+12H₂O (Michel and Krafft.)

100 pts. H₂O dissolve 6.5 pts. Na₂HPO₄+12H₂O at 0°; 27.5 pts. at 30°. (Tilden, C. Soc. 45. 409.)

Solubility in H₂PO₄+Aq at 25°.

In 1000 g. of the solution, mol.	
Na	PO ₄
2.62	1.09
1.56	0.78
2.38	1.60
3.18	2.24
4.65	3.55
5.63	3.87

(D'Ans and Schreiner, Z. phys. Ch. 1910, 101.)

See also above.

Melts in crystal water below 100° easily forms supersaturated solutions. (Lussac.)

Melts in crystal H₂O at 34.6° (Persson), 40-41° (Mulder).

Melts in crystal H₂O at 35°. (Tilden, C. Soc. 45. 409.)

Supersaturated solutions are brought to crystallization by addition of a crystal of Na₂HPO₄+12H₂O or an isomorphous substance as Na₂HAsO₄+12H₂O. (Tilden, C. Soc. 35. 200.)

Insol. in alcohol.

Sodium dihydrogen phosphate, NaH₂PO₄·H₂O.

Very sol. in H₂O. Insol. in alcohol. (Graham.)

+2H₂O. Unchanged on air. Very sol. in H₂O, and solubility increases rapidly with temperature. (Joly and Dufet, C. R. 1391.)

pts. H_2O dissolve 59.9 pts. at 0° ; 84.6 at 18° . (Joly and Dufet.)

Solubility of NaH_2PO_4 in H_2O at t° .
of anhydrous NaH_2PO_4 in 100 g. H_2O .
Solid phase; $\text{NaH}_2\text{PO}_4 + 2\text{H}_2\text{O}$.

	G. NaH_2PO_4	t°	G. NaH_2PO_4
1	57.86	28.0	101.71
0	59.08	30.0	106.45
0	61.47	31.0	108.93
0	63.82	33.0	114.31
0	69.87	34.0	117.14
0	76.72	35.0	120.44
0	85.21	37.0	126.76
0	94.63	40.2	138.16
0	96.73	40.55	110.83
0	99.20

40.8° ; transition point.

Solid phase; $\text{NaH}_2\text{PO}_4 + \text{H}_2\text{O}$.

	142.55	52	163.84
	143.83	55	170.85
	148.20	56	173.23
	158.61	57	175.81

57.4° ; transition point.

Solid phase; NaH_2PO_4 .

	177.24	69.	190.24
	179.33	80.	207.29
	181.35	90.	225.31
	184.99	99.1	246.56

Madzu, Chem. Soc. 1912, 33. 359.)

Solubility in $\text{H}_3\text{PO}_4 + \text{Aq}$ at 25° .

In 1000 g. of the solution, mols.

Na	PO_4
6.19	4.68
6.01	4.67
5.12	4.36
4.81	4.22
4.36	4.08
4.06	4.03
4.19	4.38
4.32	4.96
4.65	5.89
4.88	6.40

and Schreiner, Z. phys. Ch. 1910, 75. 101.)

Sodium trihydrogen phosphate,
 $\text{Na}_3\text{H}_3(\text{PO}_4)_2$.

hygroscopic. Sol. in H_2O in all proportions. (Joulié, C. R. 1902, 134. 604.)
 $\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O . (Filhol and others, C. R. 93. 388.)

$+7\text{H}_2\text{O}$. (Salzer, Arch. Pharm. 1894, 232. 365.)
 $+7\frac{1}{2}\text{H}_2\text{O}$. (Joulié.)

Sodium orthophosphate acid, NaH_2PO_4 , H_3PO_4 .

Hygroscopic. Decomp. by alcohol. (Staudenmaier, Z. anorg. 1894, 5. 395.)

Solubility of NaH_2PO_4 , H_3PO_4 in H_2O at t° .

t°	% NaH_2PO_4 , H_3PO_4	Solid phase
- 5.7	20.77	Ice
- 7.9	26.92	"
- 11.4	34.15	"
- 38	56.66	"
- 34	80.46	NaH_2PO_4
+ 41	81.82	"
51.7	83.68	"
79.7	87.48	"
85	88.65	"
101.7	91.47	" + NaH_2PO_4 , H_3PO_4
104.5	92.67	NaH_2PO_4 , H_3PO_4
110	95.79	"
119	97.99	"
126.5	100	"

(Parravano and Mieli, Gazz. ch. it. 38, II. 536.)

Solubility in anhydrous H_3PO_4 .

t°	% NaH_2PO_4 , H_3PO_4
98.5	52.72
111	69.59
119	77.55
122	81.71
123	87.20

(Parravano and Mieli.)

$+ \text{H}_2\text{O}$. Very deliquescent. (Salzer, Arch. Pharm. 1894, 232. 369.)

Sodium pyrophosphate, $\text{Na}_4\text{P}_2\text{O}_7$, and $+10\text{H}_2\text{O}$.

Less sol. in H_2O than sodium hydrogen orthophosphate. (Clark, Ed. J. Sci. 7. 298.)

100 pts. H_2O dissolve (a) pts. $\text{Na}_4\text{P}_2\text{O}_7$, (b) pts. $\text{Na}_4\text{P}_2\text{O}_7 + 10\text{H}_2\text{O}$ at:

	0°	10°	20°	30°	40°	50°
a.	3.16	3.95	6.23	9.95	13.50	17.45
b.	5.41	6.81	10.92	18.11	24.97	33.25

	60°	70°	80°	90°	100°
a.	21.83	25.62	30.04	35.11	40.26
b.	44.07	52.11	63.40	77.47	93.11

(Poggiale.)

Sol. in H_2SO_4 . (Walden, Z. anorg. 1902, 29. 384.)

Crystallizes unchanged from $\text{NH}_4\text{Cl} + \text{Aq}$ (Winkler), or conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Uelsmann.)

Decomp. into orthophosphate by heating with H_2SO_4 , HCl , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{H}_3\text{PO}_4 + \text{Aq}$. Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in alcohol. Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Sodium hydrogen pyrophosphate, $\text{Na}_2\text{H}_2\text{P}_2\text{O}_7$.

Decomp. by H_2O . Sol. in H_2O containing $\text{HC}_2\text{H}_3\text{O}_2$ without decomp. (Bayer, J. pr. 106. 501.)

Sl. sol. in alcohol. Much more sol. in H_2O than NaH_2PO_4 .

+ $6\text{H}_2\text{O}$. (Rammelsberg, B. A. B. 1883. 21.)

100 g. sat. solution contain 14.95 g. $\text{Na}_2\text{H}_2\text{P}_2\text{O}_7$ at 18° . (Giran, A. ch. 1902, (7) 30. 249.)

$\text{NaH}_2\text{P}_2\text{O}_7$. Very hygroscopic. (Salzer, Arch. Pharm. 1894, 232. 369.)

100 g. sat. solution contain 62.7 g. at 18° . (Giran, A. ch. 1902, (7) 30. 249.)

$\text{Na}_2\text{HP}_2\text{O}_7 + \text{H}_2\text{O}$. Sol. in 3 pts. H_2O . (Salzer, Arch. Pharm. 1894, 232. 366.)

+ $6\text{H}_2\text{O}$. 100 g. sat. solution contain 28.17 g. $\text{Na}_2\text{HP}_2\text{O}_7$ at 18° . (Giran.)

Sodium tetraphosphate, $\text{Na}_4\text{P}_4\text{O}_{13}$.

Slowly sol. in 2 pts. cold H_2O . Easily decomp.

+ $18\text{H}_2\text{O}$. (Uelsmann.)

Sodium hydrogen tetraphosphate,

$\text{Na}_4\text{H}_2\text{P}_4\text{O}_{13}$.

Sol. in H_2O .

Sodium dekaphosphate, $\text{Na}_{12}\text{P}_{10}\text{O}_{31}$.

Sol. in H_2O . (Fleitmann and Henneberg, A. 65. 333.)

Sodium strontium dimetaphosphate,

$\text{Na}_2\text{Sr}(\text{P}_2\text{O}_5)_2 + 4\text{H}_2\text{O}$.

As the NaBa comp. (Glatzel, Dissert. 1880.)

Sodium strontium trimetaphosphate,

$\text{NaSrP}_3\text{O}_9 + 3\text{H}_2\text{O}$.

Easily sol. in H_2O and acids. (Fleitmann, A. 65. 315.)

Sodium strontium orthophosphate, NaSrPO_4 , + H_2O .

Scarcely sol. in H_2O ; sol. in acids.

+ $9\text{H}_2\text{O}$. (Joly, C. R. 104. 905.)

Sodium strontium pyrophosphate (?).

Sl. sol. in H_2O . Insol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Baer, Pogg. 75. 166.)

Easily sol. in $\text{HCl} + \text{Aq}$, or $\text{HNO}_3 + \text{Aq}$.

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Sodium thallium pyrophosphate,
 $\text{Na}_4[\text{Tl}(\text{P}_2\text{O}_7)_2] + 6\text{H}_2\text{O} = \text{Na}(\text{TlP})_2\text{O}_7 + 6\text{H}_2\text{O}$.

Decomp. by H_2O . (Rosenheim, I 48. 588.)

Sodium thorium orthophosphate,
 $\text{NaTh}_2(\text{PO}_4)_4$.

Insol. in acids. (Wallroth, Bull. 39. 316.)

Sodium thorium phosphate, Na_2O , $3\text{P}_2\text{O}_5$.

Insol. in HNO_3 , HCl , or aqua regia. and Ouvrard, C. R. 105. 30.)

$5\text{Na}_2\text{O}$, 2ThO_2 , $3\text{P}_2\text{O}_5$. Sol. in HN (T. and O.)

Na_2O , ThO_2 , P_2O_5 . (T. and O.)

Sodium thorium pyrophosphate,
 $\text{ThP}_2\text{O}_7 + 2\text{H}_2\text{O}$.

(Cleve.)

Sodium tin (stannic) phosphate, Na_2Sn

(Ouvrard, C. R. 111. 177.)

$\text{Na}_2\text{Sn}(\text{PO}_4)_2$. (Wunder, J. pr. (2)

$6\text{Na}_2\text{O}$, 3SnO_2 , $4\text{P}_2\text{O}_5$. (Ouvrard.)

Sodium titanium phosphate, $\text{NaTi}_2\text{P}_2\text{O}_7$

Insol. in acids. (Rose, J. B. 1857.

$6\text{Na}_2\text{O}$, TiO_2 , $4\text{P}_2\text{O}_5$. (Ouvrard, C 177.)

Sodium uranium phosphate, $\text{UO}_2 \cdot \text{Na}_2\text{P}_2\text{O}_7$

Easily attacked by acids. (Colan 1907, (8) 12. 137.)

Sodium uranium metaphosphate, 4UC
 $3\text{P}_2\text{O}_5$.

Insol. in boiling HNO_3 . (Colani.)

Sodium uranium pyrophosphate, 3I
 $6\text{Na}_2\text{O}$, $4\text{P}_2\text{O}_5$.

Sol. in acids. (Colani.)

Sodium uranyl phosphate, Na_2O , UO_2

(Ouvrard, C. R. 110. 1333.)

$2\text{Na}_2\text{O}$, UO_2 , P_2O_5 . (Ouvrard.)

Na_2O , 5UO_2 , $2\text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$. Insol decomp. by acetic acid. (Werthe 312.)

Sodium uranyl pyrophosphate.

Very sol. in H_2O . (Persoz, A. ch 322.)

Sodium ytterbium pyrophosphate, $\text{Na}_2\text{YbP}_2\text{O}_7$

Easily sol. in the strong acids. (W

Sodium yttrium pyrophosphate, Na_2Y

Sol. in H_2O . (Stromeyer.)

Insol. in H_2O . Easily sol. in stro (Wallroth.)

zinc triphosphate,
 $4\text{ZnO}, 3\text{P}_2\text{O}_5 + 19\text{H}_2\text{O}$.
 H_2O . (Schwarz, Z. anorg. 1895, 9.)

zinc trimetaphosphate, $\text{Na}_2\text{O}, 2\text{ZnO}$,
 $3\text{P}_2\text{O}_5$.
 sol. in H_2O . (Fleitmann and Henne-
 35. 304.)

zinc tetrametaphosphate,
 $\text{Zn}(\text{PO}_3)_4 + 6\text{H}_2\text{O}$.
 comp. (Glatzel, Dissert. 1880.)

zinc octometaphosphate,
 $\text{Zn}_8(\text{PO}_3)_8$
 in acids.
 conc. H_2SO_4 . (Tammann, J. pr.
 45. 420.)

zinc orthophosphate, NaZnPO_4 .
 ltly sol. in H_2O or acetic acid. Easily
 l. mineral acids. (Scheffer, A. 145.
 , $\text{ZnO}, \text{P}_2\text{O}_5$. Insol. in H_2O ; sol. in
 . (Ouvrard, C. R. 106. 1796.)

zinc pyrophosphate, $\text{Na}_2\text{ZnP}_2\text{O}_7$.
 in H_2O ; sol. in dil. acids. (Wall-
 $\text{P}_2\text{O}_7, \text{Zn}_2\text{P}_2\text{O}_7 + 24\text{H}_2\text{O}$. Very efflores-
 ahl.)
 $\text{P}_2\text{O}_7, \text{Zn}_2\text{P}_2\text{O}_7 + 2\frac{1}{2}, 3, 3\frac{1}{2}$, and $8\text{H}_2\text{O}$.
 H_2O ; sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Pahl,
 . F. 30, 7. 35.)
 $\text{P}_2\text{O}_7, 5\text{Zn}_2\text{P}_2\text{O}_7 + 20\text{H}_2\text{O}$. Insol. in
 ahl.)
 $\text{P}_2\text{O}_7, 4\text{Zn}_2\text{P}_2\text{O}_7 + 12\text{H}_2\text{O}$. Sl. sol. in
 ahl.)

zirconium phosphate, $\text{Na}_2\text{O}, 4\text{ZrO}_2$,
 $3\text{P}_2\text{O}_5 = \text{NaZr}_2(\text{PO}_4)_3$.
 in acids or aqua regia. (Troost and
 C. R. 105. 30.)
 , $3\text{ZrO}_2, 4\text{P}_2\text{O}_5$. Sol. in acids. (T.
 , $\text{ZrO}_2, 2\text{P}_2\text{O}_5$. Sol. in acids. (T. and

phosphate fluoride, $\text{Na}_3\text{PO}_4, \text{NaF} +$
 H_2O .
 s. H_2O dissolve, at 25° , 12 pts. salt
 1 solution of 1.0329 sp. gr.; at 70° ,
 salt and form solution of 1.1091 sp.
 gleb, A. 97. 95.)
 $\text{O}_4, \text{NaF} + 19\text{H}_2\text{O}$, and $22 \text{H}_2\text{O}$. Sol.
 (Baumgarten, J. B. 1865. 219.)

phosphate stannate,
 $\text{SnPO}_4, \text{Na}_2\text{O}, \text{SnO}_2 + 4\text{SH}_2\text{O}$.
 tl, B. 1907, 40. 2132.)

Sodium phosphate titanate,
 $\text{Na}_2\text{O}, \text{TiO}_2, \text{P}_2\text{O}_5 + 3\text{H}_2\text{O}$.
 Hygroscopic. (Mazzuchelli and Pantan-
 elli, C. C. 1909, II. 420.)

Sodium phosphate vanadate.
 See Phosphovanadate, sodium.

Strontium monometaphosphate, $\text{Sr}(\text{PO}_3)_2$.
 Insol. in H_2O and acids. Not decomp. by
 alkali carbonates + Aq. (Maddrell, A. 61. 61.)

Strontium hexametaphosphate.
 Nearly insol. in H_2O ; easily sol. in acids.
 (Lüder, Z. anorg. 5. 15.)

Strontium orthophosphate, basic, $\text{Sr}(\text{OH})_2$,
 $\text{Sr}_3(\text{PO}_4)_2$.
 (Woyczynski, Z. anorg. 1894, 6. 311.)

Strontium orthophosphate, $\text{Sr}_3(\text{PO}_4)_2$.
 Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. (Erlen-
 meyer, J. B. 1857. 145.)

Strontium hydrogen phosphate, SrHPO_4 .
 Insol. in H_2O . Sol. in H_3PO_4 , HCl , or
 $\text{HNO}_3 + \text{Aq}$. (Vauquelin.) Easily sol. in
 cold ammonium nitrate, chloride, or suc-
 cinate + Aq, but is partly precipitated by a
 little $\text{NH}_4\text{OH} + \text{Aq}$. (Brett.)
 Sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$. (Fuchs, 1834.)
 Sol. in Na citrate + Aq. (Spiller.)
 Partly decomp. by boiling Na_2CO_3 , and
 $\text{K}_2\text{CO}_3 + \text{Aq}$. (Dulong.)
 $\text{SrH}_4(\text{PO}_4)_2 + 2\text{H}_2\text{O}$. Decomp. by treating
 with H_2O , leaving 4.29% SrHPO_4 . (Barthe.)

Strontium phosphate, acid, $\text{H}_2\text{O}, 2\text{SrO}$,
 $3\text{P}_2\text{O}_5 + x\text{H}_2\text{O}$.
 Entirely sol. in H_2O . (Barthe, C. R. 114..
 1267.)

Strontium pyrophosphate, $\text{Sr}_2\text{P}_2\text{O}_7 + \text{H}_2\text{O}$.
 Somewhat sol. in H_2O . Easily sol. in HCl
 or $\text{HNO}_3 + \text{Aq}$. Insol. in $\text{HC}_2\text{H}_3\text{O}_2$ or
 $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Schwarzenberg, A. 65. 144.)
 $+ 2\frac{1}{2}\text{H}_2\text{O}$. (Knorre and Oppelt, B. 21.
 773.)

Strontium hydrogen pyrophosphate,
 $\text{SrH}_2\text{P}_2\text{O}_7, 2\text{Sr}_2\text{P}_2\text{O}_7 + 6\text{H}_2\text{O}$.
 Ppt. (Knorre and Oppelt, B. 21. 772.)
 $\text{SrH}_2\text{P}_2\text{O}_7, 3\text{Sr}_2\text{P}_2\text{O}_7 + \text{H}_2\text{O}$, and $+ 2\text{H}_2\text{O}$.
 (Knorre and Oppelt.)
 $\text{Sr}_3\text{H}_2(\text{P}_2\text{O}_7)_5 + 8\text{H}_2\text{O}$, and $+ 12\text{H}_2\text{O}$.
 Ppt. (Pahl, Gm. - K. 2, 2. 172.)
 $\text{Sr}_{10}\text{H}_2(\text{P}_2\text{O}_7)_{10} + 5\text{H}_2\text{O}, + 18\text{H}_2\text{O}$ and
 $+ 20\text{H}_2\text{O}$.
 Insol. in $\text{Sr}(\text{NO}_3)_2 + \text{Aq}$ or $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$.
 (Pahl, Gm. - K. 2, 2. 171.)

Strontium thorium phosphate,
 $\text{Th}_2\text{O}, \text{SrO}, \text{P}_2\text{O}_5$.
 (Colani, C. R. 1909, 149. 209.)

Strontium uranium metaphosphate,
 $\text{UO}_2, \text{SrO}, \text{P}_2\text{O}_5$.
 (Colani, A. ch. 1907, (8) 12. 141.)

Strontium uranyl phosphate,
 $\text{SrO}, 4\text{UO}_2, 2\text{P}_2\text{O}_5 + 21\text{H}_2\text{O}$.
 (Blinkoff, Dissert. 1900.)
 $2\text{SrO}, 5\text{UO}_2, 2\text{P}_2\text{O}_5 + 24\text{H}_2\text{O}$. As Ba comp.
 (Blinkoff.)

Strontium phosphate chloride, $3\text{Sr}_3(\text{PO}_4)_2, \text{SrCl}_2$.
Strontium apatite. Insol. in H_2O . (Deville and Caron.)

Tellurium phosphate (?).
 Insol. in H_2O . (Berzelius.)

Thallous metaphosphate, TlPO_3 .
 Two modifications:
 a. Difficultly sol. in H_2O .
 β. Extremely easily sol. in H_2O . (Lamy.)

Thallous orthophosphate, Tl_3PO_4 .
 1 pt. is sol. in 201.2 pts. H_2O at 15° , and 149 pts. boiling H_2O ; sol. in $\text{HNO}_3 + \text{Aq}$. (Crookes.) Sl. sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Very easily sol. in solutions of NH_4 salts. (Carstanjen.) Insol. in alcohol. (Lamy.)

Thallous hydrogen phosphate, Tl_2HPO_4 .
Anhydrous. Much less sol. in H_2O than the hydrous salt, but easily sol. in a solution of the hydrous salt. (Lamy.)
 $+ \frac{1}{2}\text{H}_2\text{O}$. Easily sol. in H_2O . Insol. in alcohol. (Lamy.)
 Composition is $\text{HTl}_2\text{PO}_4, 2\text{H}_2\text{TlPO}_4$. (Rammelsberg, W. Ann. 16. 694.)

Thallous dihydrogen phosphate, TiH_2PO_4 .
 Very easily sol. in H_2O . Insol. in alcohol. (Rammelsberg, B. 3. 278.)

Trithallous trihydrogen phosphate, $\text{Tl}_3\text{HPO}_4, 2\text{TiH}_2\text{PO}_4$.
 True composition of Tl_3HPO_4 of Lamy. (Rammelsberg.)

Thallous pyrophosphate, $\text{Tl}_4\text{P}_2\text{O}_7$.
 Sol. in 2.5 pts. H_2O with slight decomposition. (Lamy.)
 $+ 2\text{H}_2\text{O}$. More sol. in H_2O than the above salt, with partial decomp. (Lamy.)

Thallous hydrogen pyrophosphate, $\text{H}_2\text{Tl}_2\text{P}_2\text{O}_7 + \text{H}_2\text{O}$.
 Very sol. in H_2O . (Lamy.)

Thallic phosphate, basic, $2\text{Tl}_2\text{O}_3, \text{P}_2\text{O}_5 + 5\text{H}_2\text{O}$.
 Insol. in H_2O .

Thallic phosphate, basic, $\text{Tl}_2\text{P}_2\text{O}_7 + 12\text{H}_2\text{O}$. (Rammelsberg, W. Ann. 16. 694.)
 $\text{Tl}_4\text{P}_4\text{O}_{19} + 12\text{H}_2\text{O}$. (R.)

Thallic phosphate, $\text{TlPO}_4 + 2\text{H}_2\text{O}$.
 Completely insol. in H_2O . Sol. in HNO_3 , and dil. $\text{HCl} + \text{Aq}$. (Willm.)

Thorium metaphosphate, $\text{Th}(\text{PO}_3)_4$.
 Insol. in H_2O . (Troost, C. R. 101.)

Thorium metaphosphate, $\text{ThO}_2, 2\text{P}_2\text{O}_5$.
 Insol. in acids. (Johnson, B. 22.)

Thorium orthophosphate, $\text{Th}_3(\text{PO}_4)_4$.
 Insol. in H_2O and phosphoric acid (Cleve); also acetic acid. (Cleve.)
 Sol. in HCl , and $\text{HNO}_3 + \text{Aq}$. (C.)

Thorium hydrogen phosphate, ThH_2PO_4 .
 H_2O .
 Precipitate.

Thorium pyrophosphate, $\text{ThP}_2\text{O}_7 + 2\text{H}_2\text{O}$.
 Precipitate. Insol. in H_2O . Sol. in excess of pyrophosphoric acid or sodium phosphate + Aq . (Cleve.)

Thorium phosphate bromide.
 See Bromophosphate, thorium.

Thorium phosphate chloride.
 See Chlorophosphate, thorium.

Tin (stannous) phosphate, $5\text{SnO}, 4\text{H}_2\text{O}$.
 Insol. in H_2O . (Lassen, A. 114. 1.)
 $\text{Sn}_2(\text{PO}_4)_2$. Insol. in H_2O . Sol. in acids. (Kühn.)
 Insol. in NH_4Cl or $\text{NH}_4\text{NO}_3 + \text{Aq}$.
 $\text{KOH} + \text{Aq}$.

Tin (stannic) phosphate, $2\text{SnO}_2, 10\text{H}_2\text{O}$.
 Insol. in H_2O or $\text{HNO}_3 + \text{Aq}$. (Repr. 54. 261.)
Anhydrous. Insol. in acids. (Hautefeuille and Margottet, C. R. 102. 1017.)

Tin (stannic) phosphate, SnP_2O_7 .
 Insol. in acids. (Hautefeuille and Margottet, C. R. 102. 1017.)

Tin (stannous) phosphate chloride, $\text{P}_2\text{O}_5, \text{SnCl}_2 + \text{H}_2\text{O}$.
 Not decomp. by hot H_2O . (Lassen, 114. 113.)

Titanium phosphate, $\text{Ti}_2\text{P}_2\text{O}_7 = 2\text{TiO}, \text{P}_2\text{O}_5$.
 Insol. in acids. (Hautefeuille and Margottet, C. R. 102. 1017.)
 (Ouvrard, C. R. 111. 177.)
 $+ 3\text{H}_2\text{O}$. Ppt. Insol. in H_2O .

(Knop.) Is $\text{NaTi}_2(\text{PO}_4)_3$.
1871. 324.)
Sol. in HCl , HNO_3 , H_2SO_4 .
 PO_4 .
 H , NH_4OH , $(\text{NH}_4)_2\text{CO}_3$ and
-Aq.
l. in acetic acid. (Faber, Z.
b. 288.)

phosphate, $\text{U}(\text{PO}_3)_4$.
 HNO_3 , HCl , or H_2SO_4 , even
conc. (Colani, A. ch. 1907, (8)

osphate, $\text{U}_2(\text{PO}_3)_4$.
and acids. (Hautefeuille and
R. 96. 849.)

phosphate, $\text{U}_3(\text{PO}_4)_4$.
attacked by acids than the pyro-
phosphates, especially by HNO_3 .
. 1907, (8) 12. 123.)

gen orthophosphate, $\text{UHPO}_4 +$
 H_2O . Insol. in dil., sl. sol. in
aq. Decomp. by $\text{KOH} + \text{Aq}$,
 $\text{H} + \text{Aq}$. (Rammelsberg, Pogg.

osphate, UP_2O_7 .
).
in acids. (Colani.)

hate, 2UO_2 , P_2O_5 .
acked by boiling HNO_3 .
- $5\text{H}_2\text{O}$. Insol. in H_2SO_4 and
f medium concentration. Sol.
. $\text{HCl} + \text{Aq}$. (Aloy, Dissert.

osphate, $\text{UO}_2(\text{PO}_3)_3$.
rg, B. A. B. 1872. 447.)
Insol. in acids. (Johnsson,

osphate, $\text{UO}_2\text{HPO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$.
).

Insol. in H_2O . Sol. in 67,000
+Aq, 50,000 pts. $\text{NH}_4\text{C}_2\text{H}_3\text{O}_2 +$
0 pts. of a mixture of the above
Sol. in K_2CO_3 or $\text{Na}_2\text{CO}_3 +$
, C. N. 27. 199.)

gen phosphate,
 $(\text{UO}_2)_3 + 3\text{H}_2\text{O}$.
r H_2O . Sol. in $\text{H}_3\text{PO}_4 + \text{Aq}$.
. 43. 322.)

Uranyl pyrophosphate, $(\text{UO}_2)_2\text{P}_2\text{O}_7 + 5\text{H}_2\text{O}$.
Efflorescent. Insol. in H_2O . Sol. in
 $\text{HNO}_3 + \text{Aq}$, and $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. Insol. in
 $\text{Na}_2\text{HPO}_4 + \text{Aq}$. Insol. in alcohol or ether.
(Girard, C. R. 34. 22.)
+ $4\text{H}_2\text{O}$. (Casteing, Bull. Soc. (2) 34. 20.)

Uranyl tetraphosphate (?), $\text{UO}_2\text{P}_4\text{O}_{11}$.
(Johnsson, B. 22. 978.)

Uranous orthophosphate chloride, $\text{U}_3(\text{PO}_4)_4$,
 UCl_4 .
Sl. sol. in $\text{HCl} + \text{Aq}$. Sol. in HNO_3 and
 $\text{HNO}_3 + \text{HCl}$. (Colani, A. ch. 1907, (8) 12.
127.)

Uranous hydrogen orthophosphate chloride,
 $\text{U}(\text{HPO}_4)_2$, UCl_4 .
Very sl. sol. in H_2O . (Aloy, Dissert. 1901.)

Vanadium phosphate, $(\text{VO}_2)\text{H}_2\text{PO}_4 + 4\frac{1}{2}\text{H}_2\text{O}$.
Sol. in H_2O .
See Phosphovanadic acid.

Vanadium pyrophosphate,
 $\text{V}_4(\text{P}_2\text{O}_7)_3 + 30\text{H}_2\text{O}$.
Insol. in H_2O . (Rosenheim, B. 1915, 48.
590.)

Divanadyl phosphate.
Very deliquescent, and sol. in H_2O . Insol.
in alcohol. (Berzelius.)

Ytterbium metaphosphate, $\text{Yb}(\text{PO}_3)_3$.
Insol. in H_2O . (Cleve, Z. anorg. 1902, 32.
149.)

Ytterbium orthophosphate, $\text{YbPO}_4 + 4\frac{1}{2}\text{H}_2\text{O}$.
Ppt. (Cleve.)

Ytterbium phosphate, Yb_2O_3 , $2\text{P}_2\text{O}_5 + 5\text{H}_2\text{O}$.
Sol. in H_2O . (Cleve.)

Yttrium metaphosphate, $\text{Y}(\text{PO}_3)_3$.
Insol. in H_2O or acids. (Cleve.)

Yttrium orthophosphate, YPO_4 .
Anhydrous. Insol. in H_2O or acids after
ignition.
Min. Xenotime. Insol. in conc. acids. Sl.
sol. in much conc. $\text{HCl} + \text{Aq}$, but easily sol.
therein when first heated with a little $\text{HCl} +$
 Aq . (Wartha, A. 139. 237.)

Yttrium hydrogen orthophosphate, $\text{Y}_2(\text{HPO}_4)_3$.
Decomp. by boiling with H_2O into insol.
 YPO_4 and sol. acid salt.

Yttrium pyrophosphate, $\text{YHP}_2\text{O}_7 + 3\frac{1}{2}\text{H}_2\text{O}$.
Difficultly sol. in acids. Decomp. by
 H_2SO_4 . Sol. in $\text{Na}_4\text{P}_2\text{O}_7 + \text{Aq}$. (Cleve.)
 $2\text{Y}_2\text{O}_3$, $3\text{P}_2\text{O}_5$. Insol. in acids. (Johnsson,
B. 22. 976.)

Zinc metaphosphate.

Sol. in H_2O . (Berzelius.)

Zinc dimetaphosphate, $Zn_2P_2O_6$.

Sol. only in boiling H_2SO_4 . (Fleitmann, Pogg. 78. 350.)

Not decomp. by boiling Na_2S or $(NH_4)_2S + Aq$.

+ $4H_2O$. Insol. in H_2O , but decomp. by boiling therewith. (Fleitmann, Pogg. 78. 258.)

Sol. in 4 pts. H_2O . Conc. H_2SO_4 decomp. it easily; other acids act slightly. (Glatzel, Dissert. 1880.)

Difficultly decomp. by boiling acids.

Zinc trimetaphosphate, $Zn_3(PO_3)_3 + 9H_2O$.

1 l. H_2O dissolves 0.1 g. at 20° . (Tammann, J. pr. 1892, (2) 45. 426.)

Zinc tetrametaphosphate, $Zn_4(PO_3)_4 + 10H_2O$.

Sol. in 55 pts. H_2O . Decomp. by acids only on boiling. (Glatzel, Dissert. 1880.)

Somewhat sol. in $HNO_3 + Aq$. Sol. in boiling H_2SO_4 . (Glatzel.)

Zinc orthophosphate, $Zn_3(PO_4)_2 + 4H_2O$.

Insol. in H_2O . Easily sol. in acids, NH_4OH , $(NH_4)_2CO_3$, $(NH_4)_2SO_4$, or $NH_4NO_3 + Aq$. (Heintz, A. 143. 356.)

Sol. in $NH_4Cl + Aq$. (Fuchs.)

Easily sol. in Zn salts + Aq . (Rose.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 830.)

Min. *Hopeite*.

+ $6H_2O$. (Reynoso.)

Zinc hydrogen phosphate, $ZnHPO_4 + H_2O$.

Insol. in H_2O ; sol. in $H_3PO_4 + Aq$. (Graham.)

Zinc tetrahydrogen phosphate, $ZnH_4(PO_4)_2 + 2H_2O$.

Nearly insol. in H_2O , but decomp. thereby into H_3PO_4 and $10ZnO$, $4P_2O_5 + 10H_2O$. (Demel, B. 12. 1171.)

Zinc phosphate, $10ZnO$, $4P_2O_5 + 10H_2O$.

Insol. in H_2O . (Demel, B. 12. 1171.)

Zinc pyrophosphate, $Zn_2P_2O_7 + \frac{1}{2}H_2O$.

Ppt. Sol. in $H_2SO_4 + Aq$. Sol. in acids, $KOH + Aq$, $NH_4OH + Aq$. (Schwarzenberg, A. 65. 151.)

Sol. in $Na_4P_2O_7 + Aq$ (Gladstone), and in $ZnSO_4 + Aq$. (Rose.)

Insol. in acetic acid. (Knorre, Z. anorg. 1900, 24. 389.)

+ $5H_2O$. Insol. in H_2O . (Pahl, J. B. 1873. 229.)

Zinc hydrogen pyrophosphate.

Sol. in H_2O . (Pahl, Sv. V. A. F. 30, 7. 45.)

Zinc metaphosphate ammonia.

Ppt. (Bette.)

Zinc orthophosphate ammonia, $2Zn(3NH_3 + 8H_2O)$.

(Rother, A. 143. 356.)

$6ZnO$, $3P_2O_5$, $8NH_3 + 4H_2O$. (Sch A. 145. 517.)

Zinc pyrophosphate ammonia, $3(4NH_3 + 9H_2O)$.

Ppt. Insol. in H_2O . (Bette.)

Zirconium orthophosphate, $5ZrO_2, 8H_2O$.

Somewhat sol. in acids. (Hermann 97. 321.)

Insol. in acids. (Paykull, Bull. S. 65.)

$2ZrO_2$, P_2O_5 . Not attacked by acids. (Hautefeuille and Margottet, C. 1017.)

Zirconium pyrophosphate, $Zr(PO_3)_2$.
(Knop, A. 159. 36.)**Phosphorovanadicotungstic acid****Ammonium phosphorovanadicotungstic acid, $14(NH_4)_2O$, $2P_2O_5$, $7V_2O_5$, $78H_2O$.**

Sol. in H_2O . Insol. in alcohol, ether, and benzene. (Rogers, J. Am. Ch. 1903, 25. 305.)

Phosphorimidamide, PN_2H_3 .

(Joannis, C. R. 1904, 139. 365.)

Phosphorimide, $P_2(NH)_2$.

Very sol. in ammoniacal solution. (Hugot, C. R. 1905, 141. 1236.)

Phosphornitryl, PON .

See Phosphoryl nitride.

Phosphorosomolybdic acid, $P_2O_5, 24MoO_3 + 63H_2O$.

(Rosenheim and Pinsker, Z. anorg. 70. 77.)

Ammonium phosphorosomolybdate, $2(NH_4)_2O$, $2H_3PO_4$, $12MoO_3 + 12H_2O$.

Insol. in cold, slightly sol. in acids. (Gibbs, Am. Ch. J. 5. 361.)

Phosphorosophosphomolybdic acid**Ammonium phosphorosophosphomolybdate, $9(NH_4)_2O$, $2H_3PO_4$, $3P_2O_5$, $12MoO_3 + 3SH_2O$.**

Nearly insol. in H_2O . (Gibbs.)

sophosphotungstic acid.

phosphorosophosphotungstate,
 $2\text{H}_3\text{PO}_3, \text{P}_2\text{O}_5, 24\text{WO}_3 + 13\text{H}_2\text{O}$.
 Insol. in cold boiling H_2O . (Gibbs, Am. Ch.

sotungstic acid.

phosphorosotungstate, $6(\text{NH}_4)_2\text{O}$,
 $22\text{WO}_3 + 25\text{H}_2\text{O}$.
 Insol. in cold H_2O .

—, $5\text{K}_2\text{O}, 16\text{H}_3\text{PO}_3, 32\text{WO}_3 +$
 Insol. in hot H_2O .

—, $2\text{Na}_2\text{O}, 8\text{H}_3\text{PO}_3, 22\text{WO}_3 +$
 Insol. in cold, sl. sol. in hot H_2O .
 (Ch. J. 7. 313.)

phosphorus anhydride, P_2O_3 .

phosphorus trioxide.

phosphoric acid, H_3PO_3 .

Element. Very sol. in H_2O .

Other alkali phosphites are sol. in
 of the others are sl. sol. in H_2O ,
 $\text{H}_3\text{PO}_3 + \text{Aq}$; all are insol. in

phosphite, basic, $\text{Al}_2(\text{HPO}_3)_3$.
 (Grützner, Arch. Pharm. 1897, 235.)

Grützner, Arch. Pharm. 1897, 235.

phosphite.

Element. (Rose, Pogg. 9. 39.)
 Insol. in H_2O .

phosphite, $(\text{NH}_4)_2\text{HPO}_3 + \text{H}_2\text{O}$.
 Effervescent, and sol. in H_2O . (Rose,
)
 100 pts. cold, and less hot H_2O . Insol.
 (Berzelius.)
 Insol. in acetone. (Eidmann, C. C. 1899,
 Eidmann, B. 1904, 37. 4328.)

hydrogen phosphite,
 H_2HPO_3 .

Effervescent, and sol. in H_2O . 1 pt.
 dissolves 1.71 pts. salt at 0° ; 1.9 pts. at
 2.60 pts. at 31° . (Amat, C. R.

hydroxylamine phosphite,
 $\text{H}_2\text{OH}\text{HPO}_3$.

Insol. in H_2O and abs. alcohol. (Hofmann,
 1898, 16. 466.)

Ammonium magnesium phosphite,
 $(\text{NH}_4)_2\text{Mg}_3(\text{PHO}_3)_4 + 16\text{H}_2\text{O}$.

Slightly sol. in H_2O . (Rammelsberg, Pogg.
 181. 367.)

Antimonyl phosphite, $(\text{SbO})\text{H}_2\text{PO}_3$.

Very sol. in H_2O containing HCl . (Grütz-
 ner, Arch. Pharm. 1897, 235. 694.)

Barium phosphite, BaHPO_3 .

100 pts. H_2O dissolve 0.25 pt. (Ure.)
 Very slightly sol. in H_2O , and decomp. by
 boiling H_2O . (Dulong.)

Easily sol. in H_2O containing NH_4Cl .
 (Wackenroder, A. 41. 315.)

Sol. in $\text{H}_3\text{PO}_3 + \text{Aq}$ or $\text{HCl} + \text{Aq}$. (Railton.)

Barium hydrogen phosphite, $\text{Ba}_2\text{H}_2(\text{HPO}_3)_2 +$
 $8\text{H}_2\text{O}$.

Easily sol. in H_2O , but decomp. by boiling
 therewith. Insol. in alcohol. (Rammelsberg,
 Pogg. 132. 496.)

Barium dihydrogen phosphite, $\text{BaH}_2(\text{HPO}_3)_2 +$
 $\frac{1}{2}\text{H}_2\text{O}$.

Easily sol. in H_2O . (Rose, Pogg. 9. 215.)
 $+ \text{H}_2\text{O}$. Sol. in H_2O ; decomp. by boiling
 H_2O into a neutral insol., and an acid sol. salt.
 (Wurtz, A. 58. 66.)

$+ 2\text{H}_2\text{O}$. Easily sol. in H_2O . (Rammels-
 berg, Pogg. 132. 496.)

Insol. in alcohol. (Wurtz.)

Bismuth phosphite, $2\text{Bi}_2\text{O}_3, 3\text{P}_2\text{O}_3$.

Insol. in H_2O .
 $\text{Bi}_2(\text{HPO}_3)_3 + 3\text{H}_2\text{O}$. Ppt. (Grützner,
 Arch. Pharm. 1897, 235. 696.)

Decomp. by H_2S . Not decomp. by KOH
 $+ \text{Aq}$. (Vanino, J. pr. 1906, (2) 74. 151.)

Cadmium phosphite, $\text{CdHPO}_3 + 3\text{H}_2\text{O}$.

Ppt. (Rose, Pogg. 9. 41.)

Calcium phosphite, $\text{CaHPO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Sl. sol. in H_2O ; the aqueous solution is de-
 comp. by boiling.

$+ \text{H}_2\text{O}$. Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wacken-
 roder, A. 41. 315.)

Insol. in alcohol.

Calcium hydrogen phosphite, $\text{CaH}_2(\text{HPO}_3)_2 +$
 H_2O .

Sol. in H_2O . Aqueous solution is decomp.
 by alcohol. (Wurtz, A. ch. (3) 7. 212.)

Chromic phosphite.

Precipitate. Almost insol. in H_2O . (Rose,
 Pogg. 9. 40.)

Cobaltous phosphite, $\text{CoPHO}_3 + 2\text{H}_2\text{O}$.

Ppt. Sl. sol. in H_2O . (Rose.)

Cupric phosphite, $\text{CuHPO}_3 + 2\text{H}_2\text{O}$.

Ppt. Insol. in H_2O . (Wurtz, A. ch. (3) 16. 213.)

Didymium phosphite, $\text{Di}_2(\text{HPO}_3)_2$.

Precipitate. (Frerichs and Smith, A. 191. 331.)

Glucinum phosphite.

Precipitate. Insol. in H_2O . (Rose, Pogg. 9. 39.)

Iron (ferrous) phosphite, $\text{FeHPO}_3 + x\text{H}_2\text{O}$.

Ppt. Nearly insol. in H_2O . (Rose, Pogg. 9. 35.)

Iron (ferric) phosphite, basic, $\text{Fe}_2(\text{HPO}_3)_2$, $\text{Fe}_2(\text{OH})_6$.

(Grützner, Arch. Pharm. 1897, 235. 697.)
 $\text{Fe}_4(\text{HPO}_3)_4$, $\text{Fe}(\text{OH})_3 + 5\text{H}_2\text{O}$. Hydroscopic. (Berger, C. R. 1904, 138. 1500.)

Iron (ferric) phosphite, $\text{Fe}_2(\text{HPO}_3)_2 + 9\text{H}_2\text{O}$.

Ppt. Sol. in iron alum + Aq. (Rose.)

Lanthanum phosphite, $\text{La}_2(\text{HPO}_3)_2$.

Precipitate. (Smith.)

Lead phosphite, basic, 4PbO , $\text{P}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Ppt. (Rose, Pogg. 9. 222.)
 3PbO , $\text{P}_2\text{O}_3 + \text{H}_2\text{O}$. Insol. in H_2O . Sol. in warm dil. $\text{H}_3\text{PO}_3 + \text{Aq}$, from which it is pptd. by $\text{NH}_4\text{OH} + \text{Aq}$. (Wurtz, A. ch. (3) 16. 214.)

Lead phosphite, PbHPO_3 ,

Insol. in H_2O . Very sl. sol. in a solution of phosphorous acid; easily sol. in cold $\text{HNO}_3 + \text{Aq}$. (Wurtz.)

Lead hydrogen phosphite, $\text{PbH}_4(\text{PO}_3)_2$.

Decomp. by H_2O . (Amat, C. R. 110. 901.)

Lead pyrophosphite, $\text{PbH}_2\text{P}_2\text{O}_5$.

Gradually decomp. by H_2O into H_3PO_3 and PbHPO_3 . (Amat, C. R. 110. 903.)

Lithium hydrogen phosphite, LiH_2PO_3 .

Very sol. in H_2O . (Amat, A. ch. (6) 24. 309.)

Lithium pyrophosphite, $\text{Li}_2\text{H}_2\text{P}_2\text{O}_5$.

Very sol. in H_2O . (Amat, A. ch. 1891, (6) 24. 352.)

Magnesium phosphite, $\text{MgHPO}_3 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Rose, Pogg. 9. 28.)

Sol. in 400 pts. H_2O . (Berzelius.)
 $+4\text{H}_2\text{O}$.

Magnesium pyrophosphite, $\text{Mg}(\text{H}_2\text{P}_2\text{O}_5)_2$.

Very sol. in H_2O . (Amat, A. ch. 1891, (6) 24. 313.)

Manganous phosphite, $\text{MnHPO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Difficultly sol. in H_2O , easily in MnCl_2 or $\text{MnSO}_4 + \text{Aq}$. (Rose, Pogg. 9. 33.)

Nickel phosphite, $\text{NiHPO}_3 + 3\frac{1}{2}\text{H}_2\text{O}$.

Ppt. Sl. sol. in H_2O .

Potassium phosphite, K_2HPO_3 .

Very deliquescent. Very sol. in H_2O . Insol. in alcohol. (Dulong.)

Potassium hydrogen phosphite, $(\text{KH})\text{HPO}_3$.

1 pt. H_2O dissolves about 1.72 pts. salt at 20° . (Amat, C. R. 106. 1351.)

K_2HPO_3 , $2\text{H}_3\text{PO}_3$. Very sol. in H_2O . (Wurtz, A. 58. 63.)

Sol. in 3 pts. cold, and in less hot H_2O . (Fourcroy and Vauquelin.)

Potassium pyrophosphite, $\text{K}_2\text{H}_2\text{P}_2\text{O}_5$.

Very sol. in H_2O . (Amat, A. ch. (6) 24. 351.)

Sodium phosphite, basic, Na_2HPO_3 , $\text{NaOH} (?)$.

Not obtained in pure state (Zimmerman, B. 7. 290); $= \text{Na}_2\text{PO}_3$ (Wislicenus.)

Does not exist. (Amat.)

Sodium phosphite, $\text{Na}_2\text{HPO}_3 + 5\text{H}_2\text{O}$.

Deliquescent, and very sol. in H_2O . Insol. in alcohol.

Correct formula for Na_2PO_3 of Rose and Dulong.

Sodium hydrogen phosphite, $(\text{NaH})\text{HPO}_3 - 2\frac{1}{2}\text{H}_2\text{O}$.

0.56 pt. salt dissolves in 1 pt. H_2O at 0° ; 0.66 pt. at 10° ; 1.93 pts. at 42° . (Amat, C. R. 106. 1351.)

$\text{Na}_2\text{H}_4(\text{HPO}_3)_2 + \text{H}_2\text{O}$. Deliquescent in moist air. Sol. in 2 pts. cold, and about the same amt. hot H_2O . Sl. sol. in spirit. (Fourcroy and Vauquelin.)

Sodium pyrophosphite, $\text{Na}_2\text{H}_2\text{P}_2\text{O}_5$.

Very sol. in H_2O with gradual decomp. into Na_2HPO_3 . (Amat.)

Strontium phosphite, $\text{SrHPO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.

Difficultly sol. in H_2O . Aqueous solution decomp. on heating into a sol. acid salt and an insol. basic salt.

Strontium hydrogen phosphite, $\text{SrH}_4(\text{PO}_3)_2$.

Very sol. in H_2O . (Amat, A. ch. (6) 24. 312.)

Thallous hydrogen phosphite, TlH_2PO_3 .

Very sol. in H_2O . (Amat, A. ch. (6) 24. 310.)

Thallous pyrophosphite, $\text{Tl}_2\text{H}_2\text{P}_2\text{O}_5$.

Deliquescent. Very sol. in H_2O . (Amat.)

Stannous phosphite, SnHPO_3 .
Ppt. Sol. in $\text{HCl} + \text{Aq.}$ (Rose, Pogg. 9. 45.)

Stannic phosphite, $2\text{SnO}_2, \text{P}_2\text{O}_3$.
Ppt. (Rose, Pogg. 9. 47.)

Titanium phosphite (?)
Precipitate. (Rose, Pogg. 9. 47.)

Uranyl phosphite, $(\text{UO}_2)_2\text{H}_2(\text{HPO}_3)_4 + 12\text{H}_2\text{O}$.
Precipitate. (Rammelsberg. Pogg. 132. 500.)

Zinc phosphite, ZnHPO_3 .
Sol. in H_2O . (Rammelsberg, Pogg. 132. 481.)
+ $2\frac{1}{2}\text{H}_2\text{O}$. More easily sol. in cold than warm H_2O . (Rammelsberg.)

Zinc phosphite, acid, $\text{Zn}_2\text{H}_3\text{P}_3\text{O}_{13}$.
Sol. in H_2O .
+ $2\text{H}_2\text{O}$. Sol. in H_2O . (Rammelsberg, Pogg. 132. 493.)
 $\text{Zn}_2\text{H}_3\text{P}_3\text{O}_{13}$. Sol. in H_2O .
+ $3\text{H}_2\text{O}$. Sol. in H_2O . (Rammelsberg.)
 $\text{Zn}_2\text{H}_3\text{P}_3\text{O}_{14}$. Sol. in H_2O .
+ H_2O . Sol. in H_2O . (Rammelsberg.)

Zirconium phosphite, $\text{Zr}(\text{PO}_3)_2 + \text{H}_2\text{O}$.
Ppt. Nearly insol. in dil. mineral acids. (Hauser, Z. anorg. 1913, 84. 92.)

Phosphorous anhydride, P_2O_3 .
See Phosphorus trioxide.

Phosphorus, P.

(a) *Ordinary white phosphorus*. Insol. in H_2O , but slowly decomp. thereby (G. K.); very sl. sol. in H_2O . (Berzelius and others.)

A pure aqueous solution containing 0.1 g. P in 500 cc. H_2O can be obtained by dissolving 0.1 g. P in CS_2 , mixed with ether and hot alcohol; this solution is poured into 500 cc. boiling H_2O free from air, and the boiling continued with stirring until the alcohol, ether and CS_2 are boiled off. (Bokorny, Ch. Ztg. 1896, 20. 1022.)

100 g. H_2O sat. with P contains 0.0003 g. P. (Stich, C. C. 1903, I. 1291.)

Sol. with decomp. in hot conc. $\text{HNO}_3 + \text{Aq.}$
Decomp. by boiling caustic alkalies + Aq.
Easily sol. in SCl_2 , especially if hot. (Wöhler.)

Sol. in sulphur phosphides.
Largely sol. in PCl_3 .
Easily sol. in PCl_5 .
Sol. in PBr_3 . Sol. in PSCl_3 , easily on warming, separating on cooling. (Serullas, A. ch. 1829, 42. 25.)

Sol. in liquid SO_2 . (Sestini, Bull. Soc. 1868, (2) 10. 226.)

Sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Sol. in S_2Cl_2 , without foaming. (Nicolardot, C. R. 1908, 147. 1304.)

Sol. in PS_2Cl_2 . (Gladstone, A. 1850, 74. 91.)

Sol. in 320 pts. cold alcohol of 0.799 sp. gr., and in 240 pts. of the same when warm. Pptd. from alcoholic solution by H_2O . (Büchner.)

One grain P dissolves in 1 ounce abs. alcohol. (Schacht.)

Sol. in 20 pts. absolute ether at 20° and 240 pts. ordinary ether at 20° . (Bucholz.)

Sol. in 80 pts. absolute ether at 15.5° , and 240 pts. ordinary ether at 15.5° . (Brugnattelli, A. ch. 24. 73.)

Solubility of P_4 in 100 g. ether at t° .

t°	G. phosphorus	Sp. gr.
0	0.4335	...
5	0.62	...
8	0.79	...
10	0.85	...
15	0.9	at 13° 0.7257
18	1.005	...
20	1.04	at 19° 0.7187
23	1.121	...
25	1.39	0.7283
28	1.601	...
30	1.75	...
33	1.8	...
35	1.9984	...

(Christomanos, Z. anorg. 1905, 45. 136.)

Solubility of P_4 in 100 g. benzene at t° .

t°	G. phosphorus	Sp. gr.
0	1.513	...
5	1.99	...
8	2.31	...
10	2.4	...
15	2.7	at 13° 0.8959
18	3.1	...
20	3.21	at 19° 0.8912
23	3.3995	at 22° 0.8875
25	3.7	0.8861
28	4.35	...
30	4.601	...
33	5.0	...
35	5.17	...
40	5.75	...
45	6.105	...
50	6.8	...
55	7.315	...
60	7.9	...
65	8.4	...
70	8.898	...
75	9.4	...
81	10.027	...

(Christomanos.)

Sol. to about 1% in acetic acid. (Vulpus, Arch. Pharm. 1878, 213. 38.)

100 g. 96% acetic acid dissolve 0.105% P. (Stich, Pharm. Ztg. 1903, 48. 343.)

Sol. in 0.05 pt. CS₂ (Böttger); 0.125 pt. (Trommsdorf.)

Alcohol ppts. P from CS₂ solution.

1 pt. CS₂ dissolves 17-18 pts. P. (Vogel, J. B. 1868. 149.)

Solubility in CS₂ at t°.
(g. per 100 g. of solution.)

t°	G. P ₄	t°	G. P ₄
-10	31.40	-2.5	75.00
-7.5	35.85	0.0	81.27
-5	41.95	+5.0	86.30
-3.5	66.14	+10.0	89.80
-3.2	71.72

(Cohen and Inouye, Z. phys. Ch. 1910, 72. 418.)

Very sol. in methylene iodide. (Retgers, Z. anorg. 3. 343.)

Strong vinegar dissolves P. (Beudet.)

Sol. in considerable amount in stearic acid. (Vulpus, Arch. Pharm. (3) 13. 38.)

Sol. in ethyl chloride, benzoyl chloride, stannic chloride, and in liquid cyanogen.

Sl. sol. in ethyl nitrite, and wood-spirit.

Sl. sol. in acetone, with gradual decomposition.

Insol. in nicotine, and coniine.

Sl. sol. in cold, more sol. in hot benzene. (Mansfield.)

Sol. in 14 pts. hot, and less in cold petroleum from Amiano. (Saussure.)

Sl. sol. in "liquid paraffine." (Crismer, B. 17. 649.)

Sl. sol. in warm essential oils, as oil of turpentine, and in the fatty oils.

Sol. in hot oil of copaiba, separating out on cooling.

Sol. in hot oil of caraway, and mandarin oil. (Luca.)

Sl. sol. in cold, more sol. in hot caoutchin, depositing on cooling.

Readily sol. in warm, less in cold styrene.

Sol. in aniline, and quinoline. (Hofmann.)

Sl. sol. in cold creosote.

Somewhat sol. in fusel oil.

Easily sol. in valerianic acid, and amyl valerate.

Sol. in hexyl alcohol, ethylene chloride, allyl sulphocyanide, mercury methyl, chloroform, bromoform, warm chloral, acetic ether, aldehyde, hot cacodyl sulphide, and in cacodyl oxide.

100 g. oil of almonds sat. with P contain 1.25 g. (Stich, C. C. 1903, I. 1291.)

100 g. oleic acid sat. with P contain 1.06 g. (Stich.)

100 g. paraffine sat. with P contain 1. g. (Stich.)

(b) *Amorphous phosphorus*. Insol. in H₂. Insol. in NH₄OH + Aq. (Flückiger.)

Sol. in boiling KOH + Aq.

The statement of Burgess and Chapin (Chem. Soc. 79. 1235) that red P is sol. in aqueous alcoholic alkali is incorrect. B. ordinary crystalline and amorphous red P insol. in aqueous alcoholic alkali. (Micha. A. 1902, 325. 367.)

Insol. in liquid NH₃. (Hugot, A. ch. (7) 21, 31); (Franklin, Am. Ch. J. 1895 828.)

Bright red variety is sol. in liquid N₂ at ord. temp. leaving a black residue. (Böttcher and Lenger, B. 1909, 42. 265)

Red. Amorphous.

Sol. in S₂Cl₂ with foaming. (Nicol C. R. 1908, 147. 1304.)

Solubility of amorphous bright red in PBr₃ is diminished by long heat follows:

	172°	
Initial concentration	0.555	
Final concentration	0.374	
Length of expt. in hours	34	
	198°	218°
	0.592	0.476
	0.416	0.592
	18	17

(Buck, Dissert. 1904.)

Ordinary amorphous P₄ is sol. in A sample prepared by heating br amorphous P with 94.2% P dissolves in heating in PBr₃ as follows:

% P	0.106	0.121	0.
hours	10	20	

A finely pulverized commercial containing 98.0% P:

% P	0.92	0.11
hours	10	20

An ordinary commercial product containing 98% P:

% P	0.056	0.10
hours	10	42

(Buck.)

100 g. PBr₃ dissolve 0.2601 g. br phosphorus at 172°; 0.3634 g. (Schenk, B. 1902, 35. 353.)

Insol. in KOH + Aq.

Conc. H₂SO₄ does not act upon cold, but dissolves easily when hot.

Insol. in dil., easily sol. in conc. Aq with decomposition.

Much more sol. in HNO₃ + Aq than ordinary P. (Personne, C. R. 45. 115.)

Insol. in methylene iodide. (Retgers.)

Appreciably sol. in isobutyl alcohol. (Svedberg.)

Insol. in CS₂, alcohol, ether, ligroine, PCl₃, etc.

Sl. sol. in boiling oil of turpentine.

Other high-boiling liquids, with conversion into ordinary phosphorus.

Insol. in oil of turpentine even at 270°.

(Colson, A. ch. 1908, (8) 14. 554.)

(c) Crystalline. Insol. in, and not attacked

by dil. $\text{HNO}_3 + \text{Aq.}$

Sol. in CS_2 .

Phosphorus tribromide, PBr_3 .

Decomposed by H_2O , slowly at 8°, but very rapidly at 25°. (Löwig, Pogg. 14. 485.)

Sol. in liquid H_2S . (Antony and Magri,

Gaz. ch. it. 1905, 35. (1) 206.)

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913,

27.)

Sol. in ether, acetone, CHCl_3 , C_6H_6 and

CS_2 . (Christomanos, Z. anorg. 1904, 41. 287.)

Phosphorus pentabromide, PBr_5 .

Fumes on air, and is violently decomp. by H_2O .

Phosphorus tribromide ruthenium bromide, $\text{Ru}_2\text{P}_3\text{Br}_{18}$.

Decomp. by boiling H_2O .

Slowly sol. in hot alcohol with decomp.

Insol. in benzene, CCl_4 , ligroin and cold

alcohol. (Strecker, B. 1909, 42. 1775.)

Phosphorus thiophosphoryl bromide, PBr_2 , PSBr_2 .

Decomp. by H_2O into PSBr_2 . (Michaelis.)

Phosphorus tribromide ammonia, 3PBr_3 , 5NH_3 .

Slowly but completely sol. with decomp. in

H_2O . (Storer's Dict.)

Phosphorus pentabromide ammonia, PBr_3 , 9NH_3 .

(Besson, C. R. 111. 972.)

Phosphorus monobromotetrachloride, PBrCl_4 .

Decomp. by H_2O . (Prinvault, C. R. 74.

868.)

Phosphorus dibromotrichloride, PCl_2Br_2 .

Very unstable. (Michaelis, B. 5. 9.)

Phosphorus tetrabromotrichloride, PCl_2Br_4 .

Decomp. with H_2O . (Geuther.)

Phosphorus heptabromodichloride, PCl_2Br_7 .

Very unstable. (Prinvault, C. R. 74. 868.)

Phosphorus octobromotrichloride, PCl_2Br_8 .

Very easily decomp. (Michaelis, B. 5. 9.)

Phosphorus bromofluoride, PF_2Br_2 .

Decomp. violently with H_2O . (Moissan,

Bull. Soc. (2) 43. 2.)

Phosphorus bromonitride.

See Nitrogen bromophosphide.

Phosphorus dichloride, P_2Cl_4 .

Decomp. by H_2O . (Besson, C. R. 1910, 150. 103.)

Phosphorus trichloride, PCl_3 .

Gradually decomp. by H_2O .

0.11 g. is sol. in 100 ccm. liquid H_2S . (An-

tony, Gazz. ch. it. 1905, 35 (1) 206.)

Acted upon by liquid NH_3 . (Franklin,

Am. Ch. J. 1898, 20. 828.)

Miscible with CS_2 , C_6H_6 , CHCl_3 , and

ether.

Decomp. with alcohol.

Phosphorus pentachloride, PCl_5 .

Very deliquescent, and sol. in H_2O with

violent decomp. and evolution of heat. Sol.

in liquid HCl . Acted upon by liquid NH_3 .

Somewhat sol. without decomp. in CS_2 .

(Schiff, A. 102. 118. (Franklin, Am. Ch. J.

1898, 20. 828.)

Sol. without decomp. in benzoyl chloride.

(Gerhardt.)

Sol. in oil of turpentine with evolution of

heat.

Monophosphorus platinous chloride, PCl_2 , PtCl_2 .

Deliquescent. Sol. in H_2O with formation

of chloroplatinophosphoric acid. Similarly

decomp. by alcohol. Abundantly sol. in hot

benzene, toluene, chloroform, or carbon tetra-

chloride, and crystallizes on cooling. (Schüt-

zenberger, Bull. Soc. (2) 17. 482.)

Diphosphorus platinous chloride, 2PCl_2 , PtCl_2 .

Decomp. by H_2O with formation of chloro-

platinodiphosphoric acid. Similarly decomp.

by alcohol. Sol. without decomp. in PCl_3 ,

CCl_4 , CHCl_3 , C_6H_6 , or C_7H_8 . (Schüt-

zenberger.)

Sol. in propyl alcohol with formation of

the propyl ether of platinochlorophosphor-

ous acid and HCl . (Pomey, C. R. 104. 364.)

Phosphorus diplatinous chloride, PCl_2 , 2PtCl_2 .

Sol. in alcohol, with formation of ether

$(\text{PtCl}_2)_2\text{P}(\text{OC}_2\text{H}_5)_2$. (Cochin, C. R. 86.

1402.)

Phosphorus platonic chloride, PCl_3 , PtCl_4 .

(Schützenberger.)

Phosphorus pentachloride platonic chloride, PCl_5 , PtCl_6 , or $(\text{PCl}_4)_2\text{PtCl}_6$.

Decomp. at once by H_2O . (Baudrimont,

A. ch. (4) 2. 47.)

Phosphorus pentachloride selenium tetrachloride, $2\text{PCl}_5, \text{SeCl}_4$.

Sol. in H_2O with decomp. (Baudrimont, A. ch. (4) 2. 5.)

Phosphorus trichloride ruthenium chloride, $\text{Ru}_2\text{P}_2\text{Cl}_{18}$.

Slowly decomp. by boiling H_2O .

Sol. in benzene and CHCl_3 .

Sl. sol. in CCl_4 . Insol. in ligroin. (Strecker, B. 1909, 42. 1774.)

Phosphorus tellurium chloride, $\text{PCl}_5, 2\text{TeCl}_4$.

Very deliquescent.

Sol. in H_2O . (Metzner, A. ch. 1898, (7) 15. 203.)

Phosphorus pentachloride stannic chloride, $\text{PCl}_5, \text{SnCl}_4$.

Very deliquescent. Sol. in much H_2O with evolution of heat, forming SnCl_4 , HCl , and H_3PO_4 , and soon separates out stannic phosphate. (Casselmann, A. 83. 257.)

Phosphorus trichloride titanium chloride, $\text{PCl}_5, \text{TiCl}_4$.

(Bertrand, Bull. Soc. (2) 33. 565.)

Phosphorus pentachloride titanium chloride, $\text{PCl}_5, \text{TiCl}_4$.

Deliquescent. Decomp. by H_2O and alcohol. Sol. in ether. Sl. sol. in PCl_5 . (Tüttschew, A. 141. 111.)

Completely sol. in dil. acids. (Weber.)

Phosphorus uranium pentachloride, $\text{PCl}_5, \text{UCl}_4$.

Decomp. with H_2O .

Phosphorus pentachloride zirconium chloride, $\text{PCl}_5, \text{ZrCl}_4$.

Decomp. by H_2O with pptn. of Zr phosphate. (Paykull.)

Phosphorus trichloride ammonia, $\text{PCl}_5, 5\text{NH}_3$.

Insol. as such in H_2O , but slowly decomp. by boiling H_2O . More easily sol. with decomp. in acids. Sol. with decomp. by boiling with KOH or $\text{NaOH} + \text{Aq}$. (Berzelius.)

Phosphorus pentachloride ammonia, $\text{PCl}_5, 5\text{NH}_3$.

Properties as $\text{PCl}_5, 5\text{NH}_3$. (Berzelius.)

$\text{PCl}_5, 8\text{NH}_3$. Sl. decomp. on air. (Besson, C. R. 111. 972.)

Phosphorus pentachloride tungsten trioxide, $2\text{PCl}_5, \text{WO}_3(?)$.

(Persoz and Bloch, C. R. 28. 389.)

Phosphorus chlorobromide.

See Phosphorus bromochloride.

Phosphorus chlorofluoride, PCl_2F_2 .

Absorbed by H_2O with decomp. Al by alcohol or ether. (Poulenc, A. ch. 555.)

Phosphorus chloriodide, PCl_2I_2 .

Decomp. by moist air or H_2O . Sol. (Most, B. 13. 2029.)

Phosphorus chloronitride.

See Nitrogen chlorophosphide.

Phosphorus trifluoride, PF_3 .

Decomp. slowly by H_2O . (Moissan, Soc. (2) 43. 2.)

Rapidly absorbed by KOH or NaOH slowly by BaO_2H_2 , and $\text{K}_2\text{CO}_3 + \text{Aq}$ absorbed by absolute alcohol with d (Moissan, C. R. 99. 655.)

Phosphorus pentafluoride, PF_5 .

Fumes on air. (Thorpe, A. 182. 2)

Phosphorus pentafluoride ammonia, 5NH_3 .

(Moissan, C. R. 101. 1490.)

Phosphorus pentafluoride nitrogen peroxide

Decomp. by H_2O . (Tassell, C. R. 11)

Phosphorus fluobromide.

See Phosphorus bromofluoride.

Phosphorus fluochloride.

See Phosphorus chlorofluoride.

Phosphorus subiodide, P_2I .

Sol. in dil. HNO_3 and in alkalis (Boulouch, C. R. 1905, 141. 257.)

Phosphorus diiodide, P_2I_4 .

Decomp. by H_2O . Sol. in CS_2 . (C. winder, A. ch. (3) 30. 242.)

0.09 g. is sol. in 100 ccm. liquid H_2S tony, C. C. 1905, I. 1692.)

Phosphorus triiodide, PI_3 .

Very deliquescent. Decomp. in H_2O and by H_2O . (Corenwinder, A. ch. 242.)

Very sol. in CS_2 .

Phosphorus pentaiodide, $\text{PI}_5(?)$.

(Hampton, C. N. 42. 180.)

Phosphorus iodosulphide.

See Phosphorus sulphotriiodide.

Phosphorus nitride, P_2N_2 .

Very slightly decomp. by long with H_2O .

Completely insol. in any solvent. (B. 1903, 36. 317.)

rus suboxide, P_4O_6 .

aged in dry, gradually oxidized in air. Insol. in H_2O , alcohol, ether, and not acted on by $HCl + Aq$; oxidized by H_2SO_4 . (Marchand, J. pr. 13. 442.) Insol. in H_2O . (le Verrier, A. 27. 167.) Hydrate $P_4O_6 \cdot 2H_2O$, which gives up H_2O when dried.

modifications: (a) decomp. slowly by alkalis, (b) not decomp. by H_2O or HCl . (Reinitzer and Goldschmidt, B. 13. 442.)

oxyphosphuretted hydrogen (?), $P_4O_6 \cdot H_2$. (Franke, J. pr. (2) 35. 341.)

P_4O_{10} .

in all solvents. Decomp. by H_2O . Attacked by non-oxidizing acids. Decomposed by dil. alkalis. (Gautier, C. R. 76. 49.)

IO .

in nearly all substances. Not attacked by dilute acids; oxidized by ordinary and conc. H_2SO_4 at 200° . Attacked by dil. alkaline solutions. Perhaps identical with phosphorus suboxide P_4O_6 . (Gautier, C. R. 76. 49.)

rus oxide, P_2O_3 .

Prepared by heating with H_2O at 100° . (C. R. 1897, 124. 764.)

rus trioxide, P_4O_6 (formerly P_2O_3). Resinous, but very slowly dissolved by H_2O to form H_3PO_3 . Violently decomp. by H_2O or alcohol. Insol. in ether, carbon disulphide, benzene, or chloroform. (Thorpe and Chem. Soc. 57. 545.)

rus tetroxide, P_2O_4 .

Deliquescent. Sol. with evolution of H_2O . (Thorpe and Fulton, Chem. Soc. 833.)

rus pentoxide, P_2O_5 .

Deliquescent. Sol. in H_2O with great evolution of heat, forming H_3PO_4 . Insol. in liquid NH_3 . (Franklin, Am. Ch. Soc. 20. 828.) Insol. in acetone. (Eidmann, C. C. 1899, 10. 100.); (Naumann, B. 1904, 37. 4329.)

rus sulphur oxide, $P_2O_5 \cdot 3SO_2 = (PO_3)_3$ (phosphoryl sulphate) (?). Prepared by H_2O . Sol. in cold, more sol. in H_2O . (Weber, B. 20. 86.)

rus oxy-compounds.

Under Phosphoryl compounds.

Phosphorus oxysulphide.

See Phosphorus sulphoxide.

Phosphorus semiselenide, P_2Se .

Decomp. with H_2O . Insol. in cold, decomp. by boiling $KOH + Aq$. Insol. in, but apparently decomp. by alcohol and ether. Easily sol. in CS_2 . (Hahn, J. pr. 93. 430.)

Phosphorus monoselenide, P_2Se .

Stable in dry, decomp. in moist air and by H_2O . Insol. in alcohol and ether. Decomp. by boiling $KOH + Aq$. CS_2 dissolves out P. (Hahn, J. pr. 93. 430.)

Sl. sol. in CS_2 . (Gore, Phil. Mag. (4) 30. 414.)

Phosphorous sesquiselenide, P_4Se_3 .

Sol. in CCl_4 ; sl. sol. in CS_2 . (Meyer, Z. anorg. 1902, 30. 258.)

Phosphorus triselenide, P_2Se_3 .

Decomp. by boiling H_2O and slowly in moist air. Easily sol. in cold $KOH + Aq$, less easily in $M_2CO_3 + Aq$. Insol. in alcohol, ether, and CS_2 . (Hahn, J. pr. 93. 430.)

Phosphorus pentaselenide, P_2Se_5 .

Slowly decomp. in moist air or by H_2O , easily by $KOH + Aq$ or alcohol. Insol. in CS_2 . Sol. in CCl_4 . (Hahn, J. pr. 93. 430.)

Phosphorus selenides with M_2Se .

See M phosphoselenide, under M.

Phosphorus semisulphide, $P_4S(?)$.

1. *Liquid*. Not decomp. by, and insol. in boiled H_2O . Insol. in alcohol and ether. Sl. sol. in fats and volatile oils; decomp. by alkalis. Dissolves P on warming, with separation on cooling. Sol. in CS_2 .

2. *Red modification*. Not attacked at first by $HNO_3 + Aq$ (sp. gr. 1.22), but after a time is attacked with the greatest violence. Weak acids attack only when hot. (Berzelius, A. 46. 129.)

Existence is doubtful. (Schulze, B. 13. 1862; Isambert, C. R. 96. 1628.)

Phosphorus monosulphide, $P_2S(?)$.

1. *Ordinary*. Same properties as phosphorus semisulphide, 1.

2. *Red modification*. Unchanged by air, H_2O , or alcohol. Decomp. by conc. $KOH + Aq$, not by dilute. Sl. sol. in $NH_4OH + Aq$. (Berzelius, A. 46. 129.)

Existence is doubtful. (Schulze; Isambert.)

Does not exist. (Helff, Z. phys. Ch. 12. 206.)

Phosphorus sesquisulphide, P_4S_3 .

Not attacked by cold, slowly by hot H_2O . Cold $KOH + Aq$ dissolves with decomp.

Oxidised by HNO_3 and aqua regia. Sol. in alcohol and ether with decomp. Sol. in CS_2 (100 pts. CS_2 dissolve 80 pts. P_4S_3), PCl_3 , and PSCl_3 , and in K_2S or $\text{Na}_2\text{S} + \text{Aq.}$ (Lemoine, Bull. Soc. (2) 1. 407.)

Very sol. in CS_2 . (Rebs, A. 246. 367.)

Decomp. by dil. and conc. $\text{KOH} + \text{Aq.}$

1 pt. P_4S_3 is sol. in 9 pts. CS_2 at -20° ; in 3.7 pts. CS_2 at 0° ; in 1 pt. CS_2 at 17° ; in 40 pts. benzene at 17° ; in 9 pts. benzene at 80° ; in 32 pts. toluene at 17° ; in 6.5 pts. toluene at 111° . (Stock, B. 1910, 43. 156.)

Phosphorus trisulphide, P_2S_3 .

Decomp. by water. (Kekulé, A. 90. 310.)

Sol. in $\text{M}_2\text{CO}_3 + \text{Aq}$ with separation of S. Easily sol. in KOH , NaOH , $\text{NH}_4\text{OH} + \text{Aq.}$ (Berselius, A. 46. 129.)

Sol. in alcohol and ether. (Lemoine.)

Correct formula is P_2S_3 . (Isambert, C. R. 103. 1386.)

Extremely sl. sol. in CS_2 . (Rebs, A. 246. 368.)

Existence doubtful. (Heff, Z. phys. Ch. 12. 210.)

Phosphorus sulphide, P_2S_7 .

Sl. sol. in CS_2 . (Mai, A. 265. 192.)

Slowly decomp. by cold, rapidly by hot H_2O .

Sol. in cold alkalies.

1 pt. is sol. in 3500 pts. CS_2 at 17° ; in 20,000 pts. at 0° . (Stock, B. 1910, 43. 416.)

Phosphorus disulphide, P_2S_2 (formerly P_2S_4).

Almost insol. in CS_2 . (Heff.)

Phosphorus pentasulphide, P_2S_5 .

Very deliquescent. Decomp. by H_2O . Very sol. in KOH , NaOH , $\text{NH}_4\text{OH} + \text{Aq.}$ Sol. in $\text{M}_2\text{CO}_3 + \text{Aq}$ with separation of S at low temp. Decomposes alcohol, acetic acid, etc. (Kekulé, A. 106. 331.)

Sol. in CS_2 . (Isambert, C. R. 103. 1386.)

Not very sol. in CS_2 . (Rebs, A. 246. 367.)

Mpt., 200° ; bpt., $513-515^\circ$ at 760 mm.

Decomp. by H_2O .

Easily sol. in warm $\text{NaOH} + \text{Aq.}$

1 pt. is sol. in 450 pts. CS_2 at room temp.; in 550 pts. at 0° ; in 1200 pts. at -20° . (Stock, B. 1910, 43. 1225.)

Ordinary form.

Sol. in 195 pts. boiling CS_2 .

New form.

Sol. in 32 pts. CS_2 . (Stock, B. 1905, 38. 2722.)

Phosphorus persulphide, P_2S_{12} (?).

Decomp. by H_2O , alkalies, etc. Consists of S, and mechanically united P. (Ramme, B. 12. 941.)

Phosphorus sulphides with M_2S .

See M Phosphosulphide, under M.

Phosphorus zinc sulphide, ZnP_2S_3 .

Sol. in $\text{HCl} + \text{Aq}$ with separation of P. (Berselius, A. 46. 150.)

Phosphorus trisulphide ammonia, $\text{P}_2\text{S}_3 \cdot 2$

Decomp. by H_2O . (Bineau.)

Phosphorus pentasulphide ammonia, $\text{P}_2\text{S}_5 \cdot 6\text{NH}_3$.

Sol. in liquid NH_3 . (Stock, B. 1905. 314.)

$\text{P}_2\text{S}_5 \cdot 7\text{NH}_3$. (Stock.)

Phosphorus sulphobromide.

See Thiophosphoryl bromide.

Phosphorus sulphochloride.

See Thiophosphoryl chloride.

Phosphorus sulphotiodide, $\text{P}_2\text{S}_4\text{I}_2$.

Sl. attacked by cold, rapidly by hot violently decomp. by fuming HNO_3 . Sl. sol. in CS_2 . Sl. sol. in C_6H_6 or CHCl_3 still less in ether or absolute alcohol. (Lemoine, C. R. 115. 1301.)

$\text{P}_2\text{S}_4\text{I}_2$. Easily sol. in CS_2 . More than $\text{P}_2\text{S}_5\text{I}_2$ and less than PI_3 . (Ouvrard, ch. 1894, (7) 2. 224.)

$\text{P}_2\text{S}_4\text{I}_2$. Easily decomp. (Ouvrard.)

$\text{P}_2\text{S}_5\text{I}_2$. Insol. in H_2O ; sol. in warm sl. sol. in benzene, CHCl_3 , and glacial acid; sol. in toluene and xylene. (Woltz, Ztg. 1907, 31. 640.)

Easily sol. in CS_2 . Sl. sol. in benzene, absolute alcohol and CHCl_3 . (Ouvrard, 1892, 115. 1301.)

Phosphorus sulphoxide, $\text{P}_2\text{O}_5\text{S}_4$.

Deliquescent. Easily sol. in H_2O with comp. Sol. in 2 pts. CS_2 without decomp. Sol. in benzene with decomp. (Thornton, Chem. Soc. 59. 1019.)

$\text{P}_2\text{O}_5\text{S}_4$. Slowly decomp. by H_2O . Slightly attacked by fuming HNO_3 . (B. C. R. 1897, 124. 152.)

$\text{P}_2\text{S}_5\text{O}_4$. Deliquescent; sol. in H_2O decomp.; insol. in most solvents. (Stock, 1913, 46. 1382.)

Phosphoryl triamide, $\text{PO}(\text{NH}_2)_3$.

Insol. in boiling H_2O , $\text{KOH} + \text{Aq.}$, acids. Decomp. by long boiling with $\text{HNO}_3 + \text{Aq.}$ More easily decomp. aqua regia. Easily sol. in warm H_2SO_4 nitrosulphuric acid. (Schiff, A. 101.)

Does not exist. (Gladstone; Mem. 246. 238.)

Phosphoryl bromide, POBr_3 .

Not miscible with H_2O , but gradual comp. in contact with it. Sol. in ether, oil of turpentine (Gladstone, Mag. (3) 35. 345); in CHCl_3 , CS_2 (B. Mont, Bull. Soc. 1861. 118.)

in AsBr_3 . (Walden, Z. anorg. 4.)
 Cl_4 , and in C_6H_6 . (Oddo, Chem. 3. (2) 75.)

bromide sulphide.
 phosphoryl bromide.

bromochloride, POCl_2Br .
 by H_2O . (Menschutkin, A. 139.)

dibromochloride, POClBr_2 .
 by H_2O . (Geuther, Jena Zeit.

chloride, POCl .

rosopic. Sol. in H_2O with de-
 l. in most solvents. Sol. in PCl_3 .
 R. 1897, 125. 772.)

Decomp. by H_2O . Not acted on
 O_2 , P , PH_3 , CS_2 , I , Br , Cl , etc.
 I_4 , C_6H_6 , CS_2 , CHCl_3 and ether.
 z. ch. it. 1899, 29. (2) 318; Chem.
 8 (2) 74.)

boron chloride, POCl_2 , BCl_3 .
 phosphoryl chloride.

stannous chloride, POCl_2 , SnCl_2 .
 ent. Decomp. by H_2O (Cassel-
 l. 242.)

stannic chloride, POCl_2 , SnCl_4 .
 ent. Decomp. by H_2O . (Cassel-

titanium chloride, POCl_2 , TiCl_4 .
 ent, and decomp. by H_2O .
 gg. 132. 453.)

oryl chloride, $\text{P}_2\text{O}_5\text{Cl}_4$.

violently with H_2O . (Geuther
 lis, B. 4. 766.)

in H_2O with decomp.; very un-
 sson, C. R. 1897, 124. 1100.)

oryl chloride, PO_2Cl .
 by H_2O . (Gustavson.)
 exist. (Michaelis.)

fluoride, POF_3 .

l and decomp. at once by H_2O or
 Moissan, C. R. 102. 1245.)

imidoamide, $\text{PN}_2\text{H}_2\text{O} =$
 $\text{H})\text{NH}_2$.

H_2O ; gradually decomp. by boiling
 more rapidly in presence of KOH .
 oiling conc. $\text{HCl} + \text{Aq}$. Insol. in
 np. by hot H_2SO_4 . Moderately
 $+ \text{Aq}$ dissolves without evolution

of gas. Insol. in boiling nitric or nitrosul-
 phuric acid. (Gerhardt, A. ch. (3) 20. 255.)

Insol. in alcohol, oil of turpentine, etc.

Phosphoryl iodide, $\text{P}_2\text{I}_2\text{O}_5$ (?).

Sol. in H_2O , alcohol, and ether. (Burton,
 Am. Ch. J. 3. 280.)

PO_2I_2 . (Burton.)

Phosphoryl nitride, PON .

Insol. in H_2O , acids, or alkalies. (Glad-
 stone, Chem. Soc. 2. 121.)

Phosphoryl chlorosulphide, $\text{P}_2\text{O}_5\text{SCl}_4$.

Slowly decomp. in contact with H_2O .
 (Besson, C. R. 1897, 124. 153.)

Phosphoryl thio-compounds.

See Thiophosphoryl compounds.

Phosphoselenic acid.

See Selenophosphoric acid.

Phosphoselenide, M .

See under M .

Phosphosilicic acid.

See Silicophosphoric acid.

Phosphosilicosovanadicotungstic acid.

Ammonium phosphosilicosovanadicotung-
 state.

Exact formula not known. (E. F. Smith,
 J. Am. Chem. Soc. 1903, 25. 1225.)

Phosphosilicovanadic acid, 3SiO_2 , $2\text{V}_2\text{O}_5$,
 $2\text{P}_2\text{O}_5 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Berzelius.)

Phosphostannosovanadicotungstic acid.

Ammonium phosphostannosovanadicotung-
 state.

Exact formula not known. (E. F. Smith,
 J. Am. Chem. Soc. 1903, 25. 1226.)

Phosphosulphide, M .

See under M .

Phosphosulphuric anhydride, P_2O_5 , 3SO_2 .

Very easily decomp. (Weber, B. 19. 3190.)

Phosphotelluric acid.

Ammonium phosphotellurate, $2(\text{NH}_4)_2\text{O}$,
 P_2O_5 , $\text{TeO}_2 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O . (Weinland, Z. anorg.
 1901, 28. 61.)

$4(\text{NH}_4)_2\text{O}$, $3\text{P}_2\text{O}_5$, $2\text{TeO}_2 + 11\text{H}_2\text{O}$. Sol. in
 H_2O without decomp. (Weinland.)

Potassium phosphotellurate, $1.5K_2O, P_2O_5, TeO_3$.

+17.5 H_2O . Very sol. in H_2O .
+4.5 H_2O . Ppt. (Weinland.)

Rubidium phosphotellurate, $1.5Rb_2O, P_2O_5, TeO_3 + 4.5H_2O$.

Ppt. (Weinland.)

Sodium phosphotellurate, $2Na_2O, P_2O_5, 2TeO_3 + 9H_2O$.

Difficultly sol. in cold H_2O . (Weinland.)

Phosphothorosovanadicotungstic acid.

Ammonium phosphothorosovanadicotungstate.

Exact formula not known. (E. F. Smith, J. Am. Chem. Soc. 1903, 25, 1226.)

Phosphotitanosovanadicotungstic acid.

Ammonium phosphotitanosovanadicotungstate.

Formula not known. (E. F. Smith, J. Am. Chem. Soc. 1903, 25, 1226.)

Phosphotungstic acid, $P_2O_5, 12WO_3 + 42H_2O$.

Not efflorescent. Sol. in H_2O , alcohol, and ether. (Péchar, C. R. 110, 754.)

$P_2O_5, 16WO_3 + 69H_2O$. Very efflorescent. Sol. in H_2O , alcohol, and ether. (Péchar, C. R. 109, 301.)

+ $xH_2O = H_4PW_{10}O_{38} + xH_2O$ (α -phospholutedungstic acid). Known only in aqueous solution. (Kehrmann, B. 20, 1808.)

+48 $H_2O = H_4PW_{10}O_{38} + 16H_2O$ (α -anhydrophospholutedungstic acid). Sol. in its crystal H_2O by warmth of the hand; sol. in less than $\frac{1}{2}$ pt. H_2O . (Kehrmann.)

Correct composition is represented by $H_4PW_{10}O_{38} + 9H_2O$. (Kehrmann, Z. anorg. 1, 422.)

$P_2O_5, 20WO_3 + 8H_2O$. Very efflorescent. (Gibbs, B. 10, 1386.)

+19 $H_2O = H_{11}PW_{10}O_{38} + 8H_2O$. Sol. in H_2O . (Scheibler, B. 5, 801.)

+50, and 62 H_2O . Very efflorescent. (Péchar, C. R. 109, 301.)

3 $H_2O, P_2O_5, 21WO_3 + 30H_2O$. Efflorescent. Sol. in H_2O in nearly every proportion.

$P_2O_5, 22WO_3 + 28H_2O = H_4PW_{10}O_{38} + 18H_2O$. Efflorescent (Scheibler, B. 5, 801.)

Composition is 6 $H_2O, 22WO_3, P_2O_5 + 45H_2O$. (Gibbs.)

$H_4PO_4, 12WO_3 + 18H_2O$, or $P_2O_5, 24WO_3 + 39H_2O$. Sol. in H_2O , alcohol and ether. (Soboleff, Z. anorg. 1896, 12, 18.)

$P_2O_5, 24WO_3 + 40H_2O = 6H_4O, P_2O_5, 24WO_3 + 34H_2O$. Very efflorescent. Sol. in H_2O . (Gibbs.)

+45 H_2O .

Solubility in H_2O at t° .

t°	100 ccm. H_2O dissolve g. of the cryst. acid	Sp. g.
0	16.206	
22	49.718	
43	53.64	
92	86.75	

(Soboleff, Z. anorg. 1896, 12)

Solubility in ether at t°

t°	100 ccm. ether dissolve g. of the cryst.
0	81.19
7.8	85.32
19.2	96.01
24.2	101.34

(Soboleff.)

+53 $H_2O = 6H_4O, P_2O_5, 24WO_3$. Sol. in H_2O . (Gibbs.)

Sol. in ether. If an equal vol. placed above a layer of conc. aq. of acid, oily drops form between layers, which sink to bottom, form layer. The sp. gr. of the latter is crystallized acid dissolved in ether forms an oil of sp. gr. = 2.08 solution is miscible with alcohol with a large quantity of H_2O . (20, 1452.)

+61 H_2O . Sol. in H_2O . (G. Am. Acad. 16, 116.)

Aluminum ammonium phosphotungstate.

See **Aluminicophosphotungstate**.

Ammonium phosphotungstate, 3 $P_2O_5, 7WO_3 + Aq$.

Sl. sol. in cold H_2O without decomposition by hot H_2O . (Kehrmann, 1892, 1, 438.)

2(NH_4) $_2O, P_2O_5, 12WO_3 + 5H_2O$. Cold H_2O . (Péchar, C. R. 110, 754.)

6 NH_4 , $P_2O_5, 16WO_3 + 10H_2O$. Sol. in hot H_2O . (Péchar.)

5(NH_4) $_2O, P_2O_5, 16WO_3 + xH_2O$. (NH_4) $_4PW_{10}O_{38} + xH_2O$. (Ammonopholutedungstate). Sl. sol. in H_2O . (Kehrmann.)

3(NH_4) $_2O, P_2O_5, 16WO_3 + 16H_2O$. (NH_4) $_4PW_{10}O_{38} + 8H_2O$. (Ammonohydrophospholutedungstate). Easily sol. in H_2O . (Kehrmann.)

5 NH_4 , $P_2O_5, 17WO_3 + 16H_2O$. Sl. sol. in cold H_2O . (Kehrmann, 1894, 6, 387.)

3(NH_4) $_2O, P_2O_5, 18WO_3 + 14H_2O$. (pholutedungstate.) (Kehrmann, 1893, 4, 140.)

P_2O_5 , $21WO_3 + xH_2O$. Rather easily in hot H_2O and alcohol. $NH_4Cl + Aq.$ (Kehrmann and 25. 1972.)

$3H_2O$, P_2O_5 , $22WO_3 + 18H_2O$. d H_2O . (Gibbs.)

$3H_2O$, P_2O_5 , $24WO_3 + 26H_2O$. even in hot H_2O . (Gibbs, Proc. 6. 122.)

barium α -anhydropospholuteo-
e, $NH_4BaPW_3O_{11} + xH_2O =$
, $2BaO$, P_2O_5 , $16WO_3 + xH_2O$.
) . (Kehrmann.)

phosphotungstate, $2BaO$, P_2O_5 ,
+ $15H_2O$.

rescent. Sol. in H_2O ; insol. in
Péchar, C. R. 110. 754.)

, $16WO_3 + xH_2O = Ba_3(PW_3O_{11})$
Barium α -anhydropospholuteo-

Not efflorescent. Quite diffi-
 H_2O . (Kehrmann.)

, $16WO_3 + 10H_2O$. Efflorescent.
ch. (6) 22. 240.)

, P_2O_5 , $20WO_3 + 24H_2O$. Sol.
Gibbs, B. 10. 1386.)

, P_2O_5 , $20WO_3 + 46H_2O$. Sol.
Gibbs, Proc. Am. Acad. 16. 126.)

, $22WO_3 + 59H_2O$. Sol. in H_2O ,
pr. (2) 22. 418.)

(Kehrmann, B. 24. 2335.)

, P_2O_5 , $22WO_3 + 39H_2O$. Sol.
out decomp. (Gibbs.)

, $24WO_3 + 59H_2O$. Sol. in H_2O .

, $24WO_3 + 59H_2O$. Sol. in H_2O .

, $24WO_3 + 46H_2O = 3BaO$, $3H_2O$,
+ $43H_2O$. Easily sol. in hot H_2O .

, $24WO_3 + 48H_2O$. Sol. in H_2O .
anorg. 1896, 12. 18.)

Sol. in H_2O . (Sprenger.)

it. Sl. sol. in dil. $BaCl_2 + Aq.$
Z. anorg. 1. 423.)

assium phosphotungstate, $5BaO$,
, P_2O_5 , $22WO_3 + 48H_2O$.

O. (Kehrmann and Freinkel, B.

er phosphotungstate, $4BaO$,
, P_2O_5 , $22WO_3 + 34H_2O$.

sol. in H_2O . (Kehrmann and
25. 1966.)

lium phosphotungstate, $2BaO$,
, P_2O_5 , $24WO_3 + 46H_2O$.

O, forming cloudy liquid, which
Solution in HCl is not cloudy.
and Kraut, A. 249. 380.)

Calcium phosphotungstate, CaO , $5H_2O$,
 $16WO_3$, $P_2O_5 + 3H_2O$.

Readily sol. in H_2O . (Gibbs, Proc. Am.
Acad. 16. 130.)

$2CaO$, P_2O_5 , $12WO_3 + 19H_2O$. Efflores-
cent. Insol. in alcohol. (Péchar, C. R. 110.
754.)

$2CaO$, P_2O_5 , $20WO_3 + 22H_2O$. Efflores-
cent. (Péchar, A. ch. (6) 22. 233.)

Cadmium phosphotungstate, $2CdO$, P_2O_5 ,
 $12WO_3 + 13H_2O$.

Sl. efflorescent. Very sol. in H_2O . (Pé-
char, C. R. 110. 754.)

Cupric phosphotungstate, $3CuO$, $24WO_3$,
 $P_2O_5 + 58H_2O$.

Sol. in H_2O . (Sprenger, J. pr. (2) 22. 418.)

$2CuO$, P_2O_5 , $12WO_3 + 11H_2O$. Very efflores-
cent. (Péchar, C. R. 110. 754.)

$2CuO$, P_2O_5 , $20WO_3 + 13H_2O$. Efflores-
cent. (Péchar, A. ch. (6) 22. 235.)

Lead phosphotungstate, $2PbO$, P_2O_5 , $12WO_3$,
+ $6H_2O$.

Insol. in cold, sol. in boiling H_2O . (Pé-
char, C. R. 110. 754.)

$2PbO$, P_2O_5 , $20WO_3 + 6H_2O$. Sol. in boiling
 H_2O . (Péchar, A. ch. (6) 22. 236.)

Lithium phosphotungstate, Li_2O , P_2O_5 ,
 $12WO_3 + 21H_2O$.

Sol. in H_2O . (Péchar, C. R. 110. 754.)

Magnesium phosphotungstate, $2MgO$, P_2O_5 ,
 $12WO_3$.

Sl. efflorescent. (Péchar, C. R. 110. 754.)

$2MgO$, P_2O_5 , $20WO_3 + 19H_2O$. Sl. efflores-
cent. (Péchar, A. ch. (6) 22. 234.)

Mercurous phosphotungstate.

Insol. in dil. $HNO_3 + Aq.$ (Péchar, C. R.
110. 754.)

Potassium phosphotungstate, K_2O , P_2O_5 ,
 $12WO_3 + 9H_2O$.

Insol. in cold, sl. sol. in hot H_2O . (Péchar,
C. R. 110. 754.)

$5K_2O$, P_2O_5 , $16WO_3 + xH_2O = K_5PW_3O_{11} +$
 xH_2O . (Potassium α -phospholuteotungstate).

Very sl. sol. in cold, more easily in hot H_2O .
Sol. in cold dil. $HNO_3 + Aq.$ (Kehrmann.)

$3K_2O$, P_2O_5 , $16WO_3 + 16H_2O = K_3PW_3O_{11} +$
 $8H_2O$. (Potassium α -anhydropospho-

luteotungstate). Efflorescent. Easily sol.
in H_2O . (Kehrmann.)

$5K_2O$, P_2O_5 , $17WO_3 + 21$ or $22H_2O$. Sl.
sol. in cold H_2O . (Kehrmann, Z. anorg.
1894, 6. 387.)

$3K_2O$, P_2O_5 , $18WO_3 + 28H_2O$. (Duparc
and Pearce, Bull. Soc. Min. 1895, 18. 42.)

K_2O , $5H_2O$, P_2O_5 , $18WO_3 + 14H_2O$. Very
sl. sol. in H_2O . (Gibbs.)

$6K_2O$, P_2O_5 , $18WO_3 + 30H_2O$, and $23H_2O$.

The $23\text{H}_2\text{O}$ salt is more sol. in H_2O than the $30\text{H}_2\text{O}$ salt. (Gibbs.)

$7\text{K}_2\text{O}, \text{H}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 27\text{H}_2\text{O}$. Sol. in H_2O . (Gibbs, B. 10. 1386.)

$\text{K}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 5\text{H}_2\text{O}$. Nearly insol. in H_2O . (Péchar, A. ch. (6) 22. 231.)

$8\text{K}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 18\text{H}_2\text{O}$. Sl. sol. in H_2O . (Gibbs.)

$3\text{K}_2\text{O}, \text{P}_2\text{O}_5, 21\text{WO}_3 + 31\text{H}_2\text{O}$. Easily sol. in cold H_2O or alcohol. Much less sol. in very dil. $\text{HCl} + \text{Aq}$ or $\text{KCl} + \text{Aq}$. Decomp. by boiling H_2O . (Kehrmann and Freinkel, B. 25. 1971.)

$2\text{K}_2\text{O}, 4\text{H}_2\text{O}, \text{P}_2\text{O}_5, 22\text{WO}_3 + 2\text{H}_2\text{O}$. Very sl. sol. in H_2O . (Gibbs.)

$7\text{K}_2\text{O}, \text{P}_2\text{O}_5, 22\text{WO}_3 + 31\text{H}_2\text{O}$. Easily sol. in cold or hot H_2O . Insol. in alcohol. (Kehrmann, B. 25. 1966.)

$3\text{K}_2\text{O}, 3\text{H}_2\text{O}, \text{P}_2\text{O}_5, 24\text{WO}_3 + 8$, and $14\text{H}_2\text{O}$. Sol. in a large amount of H_2O with partial decomp. (Gibbs, Proc. Am. Acad. 16. 120.)

Practically insol. in H_2O . Easily sol. in NH_4OH , alkalies, or alkali carbonates + Aq . (Kehrmann, B. 24. 2329.)

$6\text{K}_2\text{O}, \text{P}_2\text{O}_5, 24\text{WO}_3 + 18\text{H}_2\text{O}$. Sol. in H_2O . (Gibbs, Proc. Am. Acad. 15. 1.)

Potassium lead α -phospholuteotungstate.

Sl. sol. in H_2O . (Kehrmann.)

Silver phosphotungstate, $\text{Ag}_2\text{O}, \text{P}_2\text{O}_5, 12\text{WO}_3 + 8\text{H}_2\text{O}$.

Ppt. Insol. in H_2O . (Péchar, C. R. 110. 754.)

$5\text{Ag}_2\text{O}, \text{P}_2\text{O}_5, 16\text{WO}_3 + x\text{H}_2\text{O} = \text{Ag}_5\text{PW}_6\text{O}_{31} + x\text{H}_2\text{O}$ (Silver α -phospholuteotungstate). Ppt. (Kehrmann.)

$3\text{Ag}_2\text{O}, \text{P}_2\text{O}_5, 16\text{WO}_3 + 16\text{H}_2\text{O} = \text{Ag}_3\text{PW}_6\text{O}_{28} + 8\text{H}_2\text{O}$. (Silver α -anhydrophospholuteotungstate). Easily sol. in H_2O . (Kehrmann.)

$\text{Ag}_2\text{O}, 24\text{WO}_3, \text{P}_2\text{O}_5 + 60\text{H}_2\text{O}$. Insol. in H_2O .

$3\text{Ag}_2\text{O}, 24\text{WO}_3, \text{P}_2\text{O}_5 + 58\text{H}_2\text{O}$. Insol. in H_2O . (Sprenger, J. pr. (2) 22. 418.)

Sodium phosphotungstate, $3\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 7\text{WO}_3 + \text{Aq}$.

Sol. in H_2O . (Kehrmann, Z. anorg. 1. 437.)

$5\text{Na}_2\text{O}, 11\text{H}_2\text{O}, 2\text{P}_2\text{O}_5, 12\text{WO}_3 + 26\text{H}_2\text{O} = \text{Na}_5\text{H}_{11}\text{P}_2\text{W}_{12}\text{O}_{11} + 13\text{H}_2\text{O}$ (?). (Scheibler, B. 5. 801.)

$2\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 12\text{WO}_3 + 18\text{H}_2\text{O}$. Sol. in H_2O . Insol. in alcohol. (Péchar, C. R. 110. 754.)

$5\text{Na}_2\text{O}, 14\text{WO}_3, 2\text{P}_2\text{O}_5 + 42\text{H}_2\text{O}$. Easily sol. in H_2O . (Gibbs.)

$\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3, 2\text{H}_2\text{O} + 19\text{H}_2\text{O}$. Sol. in H_2O . (Gibbs, Am. Ch. J. 1895, 17. 183.)

$\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 23\text{H}_2\text{O} = \text{Na}_2\text{O}, 7\text{H}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 16\text{H}_2\text{O}$. Easily sol. in H_2O . (Gibbs.)

+ $25\text{H}_2\text{O}$. Sl. efflorescent; very sol. in H_2O ; insol. in alcohol. (Péchar, A. ch. (6) 22. 227.)

$2\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 10\text{H}_2\text{O}$. Sol. in H_2O ; insol. in alcohol. (Péchar.)

+ $30\text{H}_2\text{O}$. (P.)

$3\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 20\text{WO}_3 + 32\text{H}_2\text{O}$. A. (P.)

$2\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 22\text{WO}_3 + 9\text{H}_2\text{O}$. Ver. in H_2O . (Gibbs.)

$3\text{Na}_2\text{O}, \text{P}_2\text{O}_5, 24\text{WO}_3 + 22\text{H}_2\text{O}$. Sol. (Brandhorst and Kraut, A. 249. 379.)

+ $30\text{H}_2\text{O}$. Sol. in H_2O . (Sob. anorg. 1896, 12. 18.)

+ $42\text{H}_2\text{O}$.

Solubility in H_2O at t° .

t°	100 ccm. H_2O d. of the crys.
0	22.04
22	59.65
93	98.184

(Soboleff, Z. anorg. 1896, 12. 3)

$2\text{Na}_2\text{O}, 4\text{H}_2\text{O}, 24\text{WO}_3, \text{P}_2\text{O}_5$. Readily sol. in H_2O . (Gibbs, Proc. A. 16. 118.)

Sp. gr. at 20° of solutions of $2\text{Na}_2\text{P}_2\text{O}_5, 24\text{WO}_3 + 23\text{H}_2\text{O}$ containing:

10.22	20.94	31.13%
1.085	1.190	1.316

42.61	52.92	64.11%
1.496	1.702	2.001

or, by calculation, $a = \text{sp. gr. if } \%$ is hydrated salt, $b = \text{sp. gr. if } \%$ is anhydrous

	5	10	15	20	
a	1.040	1.064	1.131	1.181	1
b	1.044	1.092	1.143	1.199	1

	30	35	40	45	
a	1.299	1.370	1.449	1.538	1
b	1.333	1.414	1.507	1.613	1

	55	60	64% salt
a	1.754	1.884	1.998
b	1.872

(Brandhorst and Kraut, A. 249.)

Strontium phosphotungstate, $2\text{SrO}, 12\text{WO}_3 + 17\text{H}_2\text{O}$.

Sol. in H_2O . Insol. in alcohol. C. R. 110. 754.)

Thallium phosphotungstate, $\text{Tl}_2\text{O}, 12\text{WO}_3 + 4\text{H}_2\text{O}$.

Ppt. (Péchar, C. R. 110. 754.)

Zinc phosphotungstate, $2\text{ZnO}, \text{P}_2\text{O}_5, 7\text{H}_2\text{O}$.

Efflorescent. (Péchar, C. R. 110. 754.)

Monometaphosphotungstic acid.

Ammonium monometaphosphotungstate, $(\text{NH}_4)_2\text{O}, 2\text{NH}_4\text{PO}_3, 18\text{WO}_3 + 1$

Sl. sol. in cold H_2O .

onometaphosphotungstate,
 $\text{KPO}_3, 24\text{WO}_3 + 20\text{H}_2\text{O}$.

l. in H_2O . (Gibbs, Am. Ch. J. 7.

iosphotungstic acid.

sodium orthometaphosphotung-
 $\text{K}_2\text{O}, 4\text{Na}_2\text{O}, 6\text{NaPO}_3, 6\text{K}_3\text{PO}_4,$
 $+42\text{H}_2\text{O}$.

H_2O . (Gibbs, Am. Ch. J. 7. 319.)

otungstic acid.

manganous sodium pyrophos-
state, $5(\text{NH}_4)_2\text{O}, 6\text{MnO}, 2\text{Na}_2\text{O},$
 $8\text{WO}_3 + 48\text{H}_2\text{O}$.

in cold and in hot H_2O . (Gibbs,
 1895, 17. 90.)

sodium pyrophosphotungstate,
 $\text{P}_2\text{O}_7, 3\text{Na}_4\text{P}_2\text{O}_7, 2(\text{NH}_4)_2\text{O},$
 $+31\text{H}_2\text{O}$.

sol. in cold H_2O or $\text{NH}_4\text{OH} + \text{Aq.}$
 ge amount of hot H_2O .

sodium pyrophosphotungstate,
 $3\text{MnO}, \text{P}_2\text{O}_5, 14\text{WO}_3 + 36\text{H}_2\text{O}$.

it in dry air. Sol. in H_2O and
 st. therefrom. (Gibbs.)

pyrophosphotungstate, $9\text{K}_4\text{P}_2\text{O}_7,$
 $+49\text{H}_2\text{O}$.

sl. in cold H_2O .

$3\text{H}_4\text{P}_2\text{O}_7, 22\text{WO}_3, \text{K}_2\text{O}, \text{H}_2\text{O} +$
 sol. in cold. Sol. in much boiling
 is, Am. Ch. J. 7. 392.)

nadic acid, $\text{P}_2\text{O}_5, \text{V}_2\text{O}_5, 2\text{H}_2\text{O} +$

O .

on is vanadium phosphate
 $+4\frac{1}{2}\text{H}_2\text{O}$. (Friedheim, B. 23.

e only "acid" which exists. (F.)
 $+14\text{H}_2\text{O}$. Sol. in H_2O ; can be
 i dil. $\text{H}_3\text{PO}_4 + \text{Aq.}$ (Ditte, C. R.

$\text{O}_5 + 9\text{H}_2\text{O}$. Sol. in H_2O . (Ditte.)
 O_5 . (Berzelius.)

$\text{O}_5, 6\text{V}_2\text{O}_5 + 34\text{H}_2\text{O}$. Sol. in H_2O .
 much H_2O into—

$\text{O}_5, 20\text{V}_2\text{O}_5 + 53\text{H}_2\text{O}$. Sol. in H_2O .
 Ch. J. 7. 209.)

phosphovanadate, $(\text{NH}_4)_2\text{O},$
 $\text{O}_5 + \text{H}_2\text{O}$.

cold H_2O . (Gibbs, Am. Ch. J.

Composition is $(\text{VO}_2)(\text{NH}_4)\text{HPO}_4$,
 iedheim.)

$\text{P}_2\text{O}_5, 2\text{V}_2\text{O}_5 + 7\text{H}_2\text{O}$. Easily sol.
 Gibbs.) Sl. sol. in H_2O . (Fried-

heim.) Composition is $(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5,$
 $+2(\text{VO}_2)\text{H}_2\text{PO}_4 + 5\text{H}_2\text{O}$. (Friedheim.)

$5(\text{NH}_4)_2\text{O}, 2\text{P}_2\text{O}_5, 3\text{V}_2\text{O}_5 + 24\text{H}_2\text{O}$. Easily
 sol. in H_2O . (Ditte, C. R. 102. 1019.) Could
 not be obtained. (Friedheim.)

$5(\text{NH}_4)_2\text{O}, 4\text{P}_2\text{O}_5, 2\text{V}_2\text{O}_5 + 24\text{H}_2\text{O}$. As
 above. (Ditte.) Could not be obtained.
 (Friedheim.)

$7(\text{NH}_4)_2\text{O}, \text{P}_2\text{O}_5, 12\text{V}_2\text{O}_5 + 26\text{H}_2\text{O}$. Easily
 sol. in H_2O . Composition is $2(\text{NH}_4)_2\text{HPO}_4 +$
 $5(\text{NH}_4)_2\text{O}, 12\text{V}_2\text{O}_5 + 25\text{H}_2\text{O}$. (Friedheim.)

Potassium phosphovanadate, $\text{K}_2\text{O}, \text{P}_2\text{O}_5,$
 $2\text{V}_2\text{O}_5 + 7\text{H}_2\text{O}$.

Sl. sol. in H_2O ; decomp. thereby to $7\text{K}_2\text{O},$
 $12\text{V}_2\text{O}_5, \text{P}_2\text{O}_5 + 26\text{H}_2\text{O}$.

Composition is $\text{K}_2\text{O}, \text{V}_2\text{O}_5 + 2(\text{VO}_2)\text{H}_2\text{PO}_4$
 $+ 5\text{H}_2\text{O}$. (Friedheim.)

$3\text{K}_2\text{O}, 4\text{P}_2\text{O}_5, 6\text{V}_2\text{O}_5 + 21\text{H}_2\text{O}$. Sl. sol. in
 H_2O . (Gibbs.)

$7\text{K}_2\text{O}, \text{P}_2\text{O}_5, 12\text{V}_2\text{O}_5 + 26\text{H}_2\text{O}$. Easily sol.
 in H_2O . Composition is $2\text{K}_2\text{HPO}_4 + 5\text{K}_2\text{O},$
 $12\text{V}_2\text{O}_5 + 25\text{H}_2\text{O}$. (Friedheim.)

$2\text{K}_2\text{O}, \text{P}_2\text{O}_5, \text{V}_2\text{O}_5$.

$3\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 2\text{V}_2\text{O}_5 + 5\text{H}_2\text{O}$.

$13\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 22\text{V}_2\text{O}_5 + 58\text{H}_2\text{O}$.

$15\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 25\text{V}_2\text{O}_5 + 76\text{H}_2\text{O}$.

(Friedheim, Z. anorg. 1894, 5. 446.)

$16\text{K}_2\text{O}, 2\text{P}_2\text{O}_5, 27\text{V}_2\text{O}_5 + 57\text{H}_2\text{O}$.

$6\text{K}_2\text{O}, \text{P}_2\text{O}_5, 11\text{V}_2\text{O}_5 + 33\text{H}_2\text{O}$.

$7\text{K}_2\text{O}, \text{P}_2\text{O}_5, 13\text{V}_2\text{O}_5 + 38\text{H}_2\text{O}$.

$4\text{K}_2\text{O}, \text{P}_2\text{O}_5, 3\text{V}_2\text{O}_5 + 3\text{H}_2\text{O}$.

(Friedheim, Z. anorg. 1894, 5. 459-465.)

Silver phosphovanadate, $2\text{Ag}_2\text{O}, \text{P}_2\text{O}_5, \text{V}_2\text{O}_5 +$
 $5\text{H}_2\text{O}$.

Sl. sol. in cold or hot H_2O . (Gibbs.)

Phosphovanadicotungstic acid.

Ammonium phosphovanadicotungstate,
 $(\text{NH}_4)_2\text{O}, \text{P}_2\text{O}_5, \text{V}_2\text{O}_5, \text{WO}_3 + x\text{H}_2\text{O}$.

Ppt. (Smith, J. Am. Chem. Soc. 1902, 24.
 577.)

$15(\text{NH}_4)_2\text{O}, 2\text{P}_2\text{O}_5, 6\text{V}_2\text{O}_5, 44\text{WO}_3 +$
 $106\text{H}_2\text{O}$. Sol. in H_2O . Insol. in alcohol,
 ether or benzene. (Rogers, J. Am. Chem.
 Soc. 1903, 25. 303.)

Phosphovanadicovanadiotungstic acid.

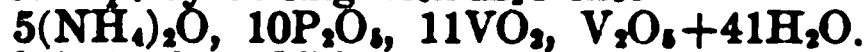
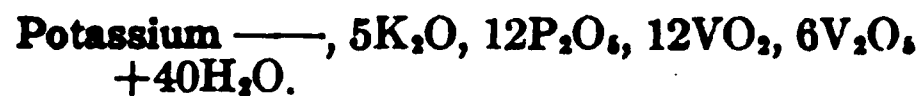
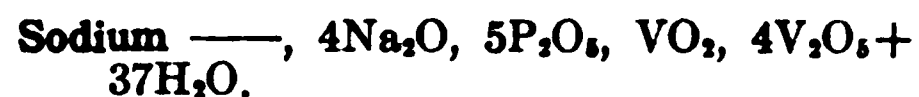
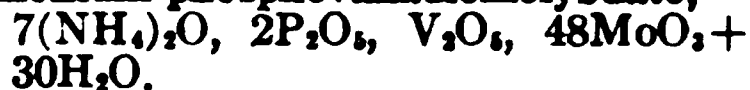
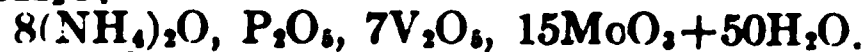
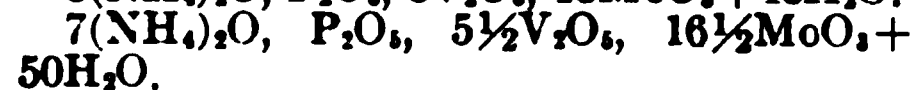
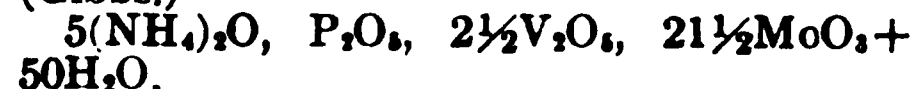
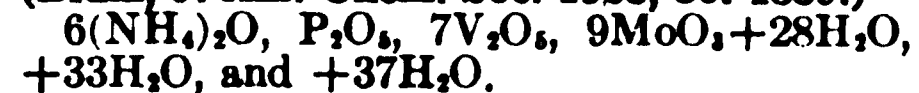
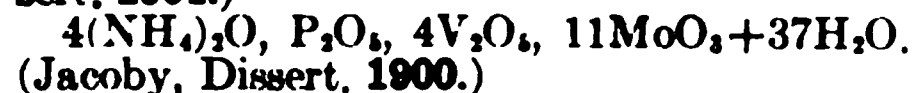
Ammonium phosphovanadicovanadiotung-
state, $14(\text{NH}_4)_2\text{O}, 2\text{P}_2\text{O}_5, 3\text{V}_2\text{O}_5, 7\text{V}_2\text{O}_5,$
 $27\text{WO}_3 + 66\text{H}_2\text{O}$.

Sparingly sol. in cold H_2O . Sol. in hot H_2O .
 (Rogers, J. Am. Chem. Soc. 1903, 25. 309.)

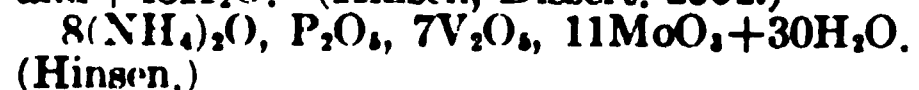
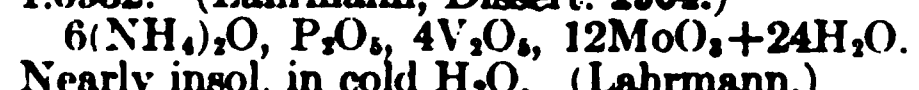
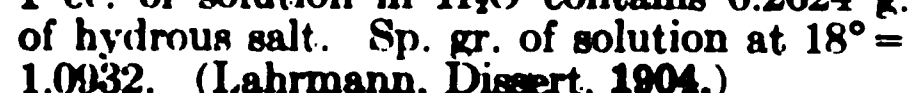
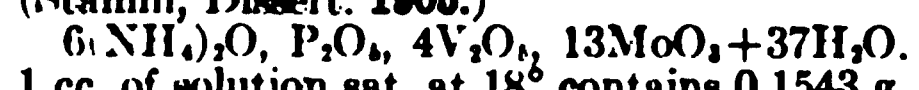
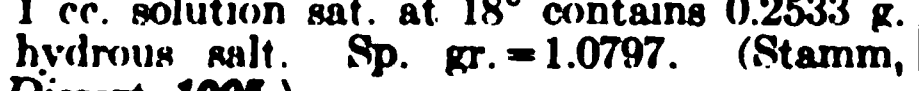
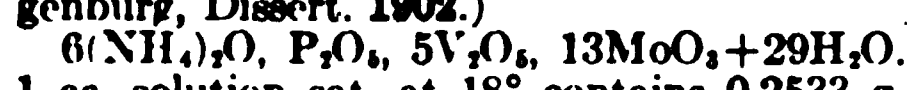
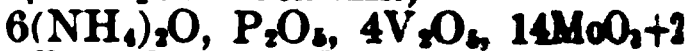
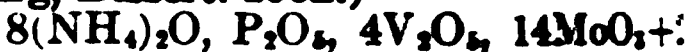
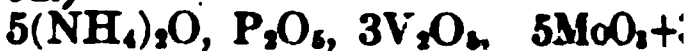
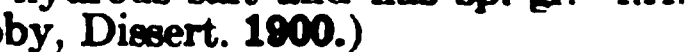
Phosphovanadicozirconosotungstic acid.

Ammonium phosphovanadicozirconosotung-
state.

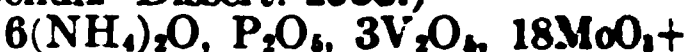
Exact formula not known. (E. F. Smith,
 J. Am. Chem. Soc. 1903, 25. 1226.)

Phosphovanadicovanadic acid.**Ammonium phosphovanadicovanadate,**Sol. in H_2O . (Gibbs, Am. Ch. J. 7. 209.)Decomp. by boiling with H_2O into—Sol. in H_2O . (Gibbs.)Decomp. by hot H_2O into—Sol. in H_2O . (Gibbs.)Insol. in H_2O . (Gibbs.)**Phosphovanadiomolybdic acid.****Ammonium phosphovanadiomolybdate,**Sl. sol. in cold, somewhat more in hot H_2O with partial decomp. (Gibbs, Am. Ch. J. 5. 391.)Easily sol. in hot H_2O without decomp. (Gibbs.)All above compounds are sol. in H_2O . (Blum, J. Am. Chem. Soc. 1908, 30. 1859.)Can be recryst. from H_2O . (Hinsen, Dissert. 1904.)

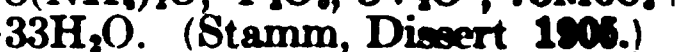
(Jacoby, Dissert. 1900.)

and $+ 43\text{H}_2\text{O}$. (Hinsen, Dissert. 1904.)1 cc. of solution in H_2O contains 0.2624 g. of hydrous salt. Sp. gr. of solution at $18^\circ = 1.0932$. (Lahrman, Dissert. 1904.)Nearly insol. in cold H_2O . (Lahrman.)1 cc. of solution sat. at 18° contains 0.1543 g. hydrous salt and has sp. gr. = 1.0900. (Toggenburg, Dissert. 1902.)1 cc. solution sat. at 18° contains 0.2533 g. hydrous salt. Sp. gr. = 1.0797. (Stamm, Dissert. 1905.)Easily sol. in H_2O with decomp. (Toggenburg, Dissert. 1902.)Decomp. by cold H_2O . (Lahrman, Dissert. 1904.)1 cc. of solution sat. at 18° contains 1.14 g. hydrous salt and has sp. gr. = 1.14. (Toggenburg, Dissert. 1900.)Extraordinarily easily sol. in H_2O . (Toggenburg, Dissert. 1902.)

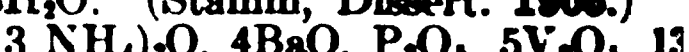
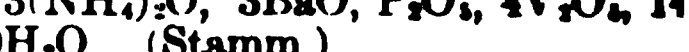
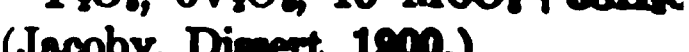
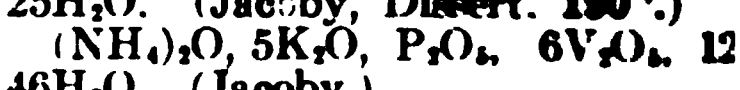
(Schulz, Dissert. 1905.)



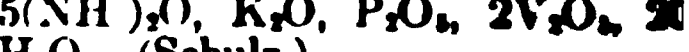
(Schulz.)

 $+ 33\text{H}_2\text{O}$. (Stamm, Dissert. 1905.)

(Hinsen, Dissert. 1904.)

 $43\text{H}_2\text{O}$. Sl. sol. in H_2O . Decomp. on heating. (Toggenburg, Dissert. 1902.) $49\text{H}_2\text{O}$. Less sol. in H_2O than NE. (Jacoby, Dissert. 1900.) $37\text{H}_2\text{O}$. Sol. in much hot H_2O with decomp. (Toggenburg, Dissert. 1902.) $46\text{H}_2\text{O}$. (Stamm, Dissert. 1905.) $40\text{H}_2\text{O}$. (Stamm.) $39\text{H}_2\text{O}$. (Stamm.) $46\text{H}_2\text{O}$. (Schulz, Dissert. 1905.)

(Jacoby, Dissert. 1900.)

 $25\text{H}_2\text{O}$. (Jacoby, Dissert. 1900.) $46\text{H}_2\text{O}$. (Jacoby.) $+ 25\text{H}_2\text{O}; + 29\text{H}_2\text{O}; + 30\text{H}_2\text{O}$. Sl. sol. in cold, more easily in hot H_2O . (Stamm, Dissert. 1905.) $31\text{H}_2\text{O}$. (Stamm.) $36\text{H}_2\text{O}$. (Jacoby, Dissert. 1900.) $43\text{H}_2\text{O}$. (Schulz, Dissert. 1905.) $52\text{H}_2\text{O}$. (Schulz.)

(Schulz, Dissert. 1905.)

sphovanadiomolybdate, $7K_2O$, $3O_2$, $9MoO_3 + 25H_2O$.

(Vèzes, 1904.)

tert. $2V_2O_5$, $20MoO_3 + 53H_2O$. (Vèzes, 1905.)

diotungstic acid.

phosphovanadiotungstate, $3P_2O_5$, V_2O_5 , $60WO_3 +$

in cold, sl. sol. in hot H_2O . $HPO_4 + Aq$, and in $NH_4OH +$

P_2O_5 , $3V_2O_5$, $16WO_3 + 37H_2O$. H_2O . (Gibbs, Am. Ch. J. 5.

$2P_2O_5$, $8V_2O_5$, $34WO_3 + 86H_2O$. Cold and hot H_2O .

alcohol, ether, CS_2 , benzene and (Rogers, J. Am. Chem. Soc.

$18BaO$, $3P_2O_5$, $2V_2O_5$, $60WO_3$

in hot H_2O with decomp. (J. 5. 391.)

$- 3K_2O$, P_2O_5 , V_2O_5 , $7WO_3 +$

$4V_2O_5$, $18WO_3 + 23H_2O$. Sol. in hot H_2O with decomp. into preceding salt. (J. 5. 391.)

Ag_2O , $2P_2O_5$, $8V_2O_5$, $33WO_3 +$

sl. in H_2O .

sol. in H_2O containing a few (Rogers, J. Am. Chem. Soc.

phosphovanadicotungstic acid.

phosphovanadiovanadicotungstate, P_2O_5 , V_2O_5 , VO_2 , $60WO_3 +$

old, easily sol. in hot H_2O . (J. 5. 391.)

1 hydrogen.

1 phosphide.

ous acid.

bromonitrite, $K_2Pt(NO_2)_4Br_2$.

in H_2O . (Blomstrand, J. pr.

40 pts. cold, and 20 pts. boil- in alcohol. Sl. sol. in KBr or (Vèzes, A. ch. (6) 29. 198.)

rs . Sol. in about 5 pts. warm ap. (Vèzes.)

$K_2Pt(NO_2)_4Br_4$. Sol. in less than 5 pts. H_2O with decomp. (Vèzes.)

Platichloronitrous acid.

Potassium platichloronitrite, $K_2Pt(NO_2)_4Cl_2$.

Rather sl. sol. in H_2O . (Blomstrand J. pr. (2) 3. 214.)

Sol. in 40 pts. cold, and 20 pts. boiling H_2O . Insol. in alcohol. Sl. sol. in KCl or $KNO_3 + Aq$. (Vèzes, A. ch. (6) 29. 183.)

$K_2Pt(NO_2)_4Cl_2$. Very sol. in H_2O . (Vèzes.)

$K_2Pt(NO_2)_4Cl_2 + H_2O$. Sol. in H_2O with decomp. (Vèzes.)

Platiodonitrous acid.

Potassium platiodonitrite, $K_2Pt(NO_2)_4I_4$.

Sl. sol. in cold, more easily in hot H_2O ; decomp. by boiling. (Vèze, A. ch. (6) 29. 207.)

$K_2Pt(NO_2)_4I_4$. As above. (Vèzes.)

Platin-

See also **Platino-**, **plato-**, **p'at-**, and **platos-**.

Platindiamine compounds.

See **Chloro-**, **bromo-**, **hydroxylo-**, **iodo-**, **nitrate-**, **nitrito-**, **sulphato-**, etc., **platindiamine compounds**.

Platintriamine carbonate,

$Pt(NH_3)_3(CO_3)_2$.

Ppt. Sol. in $NaOH + Aq$. (Geddes, J. pr. (2) 26. 257.)

— **chloride** $Pt(NH_3)_3Cl_4$.

Sol. in hot H_2O (Gerdes.)

— **chloroplatinate**, $Pt(NH_3)_3Cl_4$, $PtCl_4 + 2H_2O$.

Very sl. sol. in H_2O . (Gerdes.)

— **nitrate**, $Pt(NH_3)_3(NO_3)_4$.

Easily sol. in H_2O ; sl. sol. in $HNO_3 + Aq$. (Gerdes.)

— **sulphate**, $Pt(NH_3)_3(SO_4)_2 + H_2O$.

Nearly insol. in H_2O . (Gerdes.)

Tetraplatinamine iodide, $Pt_4(NH_3)_8I_{10}$.

(Blomstrand, B. 16. 1469.)

Octoplatinamine iodide, $Pt_8(NH_3)_{16}I_{18}$.

(Blomstrand.)

Platinic acid.

Barium platinate, basic (?), $3BaO$, $2PtO_2$.

Insol. in $HC_2H_3O_2 + Aq$; easily sol. in $HCl + Aq$. (Rousseau.)

Barium platinate, BaPtO₃.

(Rousseau, C. R. 109. 144.)

+H₂O. Insol. in dil. HNO₃+Aq; sol. in warm HCl+Aq. (Topsoë, B. 3. 464.)+4H₂O. Very sl. sol. in H₂O, BaO₂H₂, or NaOH+Aq. Easily sol. in dil. acids, except HC₂H₃O₂, in which it is insol. in the cold, but decomp. on heating. (Topsoë, l. c.)Composition is 3BaPtO₃, BaCl₂, PtCl₂O + 4H₂O (?). (Johannsen, A. 155. 204.)**Calcium platinate chloride (?), 2Ca₂Pt₂O₅Cl₂ + 7H₂O (?).**

"Herschel's precipitate."

Easily sol. in HCl+Aq, and in HNO₃+Aq, if freshly pptd. (Herschel.)Very sol. in HNO₃+Aq. (Weiss and Döbereiner, A. 14. 252.)Composition is CaPtO₃, PtCl₂O, CaO + 7H₂O (?). (Johannsen, A. 155. 204.)**Potassium platinate.**Sol. in H₂O. (Berzelius.)K₂O, PtO₃+3H₂O. Very sol. in H₂O. (Blondel, A. ch. 1905, (8) 6. 90.)K₂Pt(OH)₆. Sol. in H₂O; insol. in alcohol. (Bellucci, Z. anorg. 1905, 44. 173.)**Sodium platinate, Na₂O, 3PtO₃+6H₂O.**Dil. acids dissolve out Na₂O and leave PtO₃. Sol. in HNO₃+Aq. (Döbereiner, Pogg. 28. 180.)Na₂O, PtO₃+3H₂O. Sol. in H₂O. (Blondel.)**Metaplatinic acid, 5PtO₃, 5H₂O.**Insol. in H₂O. (Blondel, A. ch. 1905, (8) 6. 103.)**Sodium metaplatinate, Na₂O, 5PtO₃+9H₂O.**Insol. in H₂O. (Blondel.)**Platinimolybdic acid, 4H₂O, PtO₃, 10MoO₃. (Gibbs.)****Ammonium platinimolybdate,**8MoO₃, 2PtO₃, 3(NH₄)₂O + 12H₂O.4MoO₃, 2PtO₃, 2(NH₄)₂O + 19H₂O.Sol. in hot H₂O. (Gibbs, Am. Ch. J. 1895, 17. 80-82.)**Potassium platinimolybdate,**60MoO₃, PtO₃, 10K₂O + 40H₂O.Sol. in hot H₂O. (Gibbs.)**Silver platinimolybdate.****Sodium platinimolybdate, 4Na₂O, PtO₃, 10 MoO₃+29H₂O.**Sol. in H₂O. (Gibbs, Sill. Am. J. (3) 14. 61.)**Platinitungstic acid.****Ammonium platinitungstate, 4(NH₄)₂WO₆+12H₂O.**Sol. in H₂O. (Gibbs, B. 10. 1384.)**Potassium platinitungstate, 4K₂O 10WO₆+9H₂O.**Sol. in H₂O. (Gibbs.)**Sodium platinitungstate, 4Na₂O 10WO₆+25H₂O.**Sol. in H₂O. (Gibbs.)5Na₂O, 7WO₆, 2PtO₃+35H₂O.H₂O. (Gibbs.)Is double salt 3Na₂O, 7WO₆+2N (Rosenheim, B. 24. 2397.)10WO₆, PtO₃, 4Na₂O + 23H₂O.10WO₆, PtO₃, 6Na₂O + 28H₂O.20WO₆, PtO₃, 9Na₂O + 58H₂O.30WO₆, 2PtO₃, 15Na₂O + 89H₂O.30WO₆, PtO₃, 12Na₂O + 72H₂O.All are sol. in boiling H₂O. (G. Ch. J. 1895, 17. 74-80.)**Platino-**

See also Plato—.

Platinochlorophosphoric acid.

See Chloroplatinophosphoric acid.

Platinocyanhydric acid, H₂Pt(CN)Deliquescent. Very sol. in H₂ and ether.**Ammonium platinocyanide, (NH₄)₂Pt(CN)₆+3H₂O.**Very sol. in H₂O.+2H₂O. Sol. in 1 pt. H₂O, and easily in alcohol. +H₂O.**Ammonium hydroxylamine platino-**NH₄(NH₄O)Pt(CN)₆+3½H₂O.Sol. in H₂O. (Scholz, M. Ch. 1**Ammonium magnesium platinocyanide, (NH₄)₂Mg[Pt(CN)₆]+6H₂O.****Barium platinocyanide, BaPt(CN)**Sol. in 33 pts. H₂O at 16°, and i at 100°. Sol. in alcohol.**Barium potassium platinocyanide, BaK₂[Pt(CN)₆].**Sol. in H₂O.**Barium rubidium platinocyanide, BaRb₂[Pt(CN)₆].**Sol. in H₂O.

platinocyanide, $\text{CdPt}(\text{CN})_4$.
Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Martius, A.)
 $\text{CN})_4, 2\text{NH}_3 + \text{H}_2\text{O.}$ (M.)

platinocyanide, $\text{CaPt}(\text{CN})_4 + 5\text{H}_2\text{O.}$
Sol. in $\text{H}_2\text{O.}$

potassium platinocyanide,
 $\text{K}_2[\text{Pt}(\text{CN})_4]_2$.
 $\text{H}_2\text{O.}$

platinocyanide, $\text{Ce}_2[\text{Pt}(\text{CN})_4]_2 +$
 $\text{H}_2\text{O.}$

s platinocyanide ammonia,
 $\text{Pt}(\text{CN})_4, 2\text{NH}_3$.
in H_2O , but sol. in hot $\text{NH}_4\text{OH} + \text{Aq.}$

platinocyanide, $\text{CuPt}(\text{CN})_4 + x\text{H}_2\text{O.}$

platinocyanide ammonia, $\text{CuPt}(\text{CN})_4$,
 $[\text{Pt}(\text{CN})_4]_2 + \text{H}_2\text{O.}$
 $\text{CN})_4, 4\text{NH}_3$. Sol. in H_2O , alcohol,
etc.

m platinocyanide, $\text{Di}_2[\text{Pt}(\text{CN})_4]_2 +$
 $\text{H}_2\text{O.}$
scent in dry air. Sol. in $\text{H}_2\text{O.}$

um platinocyanide, $\text{Dy}_2[\text{Pt}(\text{CN})_4]_2 +$
 $\text{H}_2\text{O.}$
sol. in $\text{H}_2\text{O.}$ (Jantsch, B. 1911, 44.)

platinocyanide, $\text{Er}_2[\text{Pt}(\text{CN})_4]_2 +$
 $\text{H}_2\text{O.}$
 $\text{H}_2\text{O.}$ (Cleve.)

um platinocyanide,
 $[\text{Pt}(\text{CN})_4]_2, 3\text{Pt}(\text{CN})_2 + 18\text{H}_2\text{O.}$
in H_2O ; decomp. in the air. (Bene-
anorg. 1900, 22. 405.)

n platinocyanide, $\text{GlPt}(\text{CN})_4$.
ynski, Dissert. 1871.)

amine platinocyanide,
 $\text{I}_4\text{O}_7\text{Pt}(\text{CN})_4 + 2\text{H}_2\text{O.}$
efflorescent. Very sol. in $\text{H}_2\text{O.}$ (Scholz.)

amine lithium platinocyanide,
 $\text{I}_4\text{O}_7\text{LiPt}(\text{CN})_4 + 3\text{H}_2\text{O.}$
in $\text{H}_2\text{O.}$

platinocyanide,
 $[\text{Pt}(\text{CN})_4]_2 + 2\text{H}_2\text{O.}$
microscopic; sol. in $\text{H}_2\text{O.}$ (Renz, B. 1901,
1.)

Lanthanum platinocyanide, $\text{La}_2[\text{Pt}(\text{CN})_4]_2$
 $+ 18\text{H}_2\text{O.}$
Easily sol. in $\text{H}_2\text{O.}$ (Cleve.)

Magnesium platinocyanide, $\text{MgPt}(\text{CN})_4 +$
 $2\text{H}_2\text{O.}$

Solubility in $\text{H}_2\text{O.}$

100 g. of the sat. solution contain at:

96.4° 100°

44.33 43.96 g. $\text{MgPt}(\text{CN})_4$

(Buxhoevden, Z. anorg. 1897, 15. 325.)

$+ 4\text{H}_2\text{O.}$

Solubility in $\text{H}_2\text{O.}$

100 g. of the sat. solution contain at:

42.2° 46.3° 48.7° 55°

40.21 39.79 40.75 40.02 g. $\text{MgPt}(\text{CN})_4$,

58.1° 69° 77.8° 87.4°

42.01 43.48 44.88 45.52 g. $\text{MgPt}(\text{CN})_4$,

90°

45.59

93°

45.04 g. $\text{MgPt}(\text{CN})_4$.

(Buxhoevden.)

$+ 7\text{H}_2\text{O.}$ Sol. in 3.4 pts. H_2O at 16°.
Easily sol. in alcohol and ether.

Solubility in $\text{H}_2\text{O.}$

100 g. of the sat. solution contain at:

—4.12° +0.5° 5.5° 18.0°

24.9 26.33 28.07 31.23 g. $\text{MgPt}(\text{CN})_4$,

36.6° 45.0° 46.2°

38.36 41.32 41.96 g. $\text{MgPt}(\text{CN})_4$.

(Buxhoevden.)

Magnesium potassium platinocyanide,
 $\text{MgK}_2[\text{Pt}(\text{CN})_4]_2 + 7\text{H}_2\text{O.}$

Sol. in $\text{H}_2\text{O.}$

Mercuric platinocyanide, $\text{HgPt}(\text{CN})_4$.
Ppt.

Mercuric platinocyanide nitrate, $5\text{HgPt}(\text{CN})_4$,
 $\text{Hg}(\text{NO}_3)_2 + 10\text{H}_2\text{O.}$

Ppt.

Nickel platinocyanide ammonia, $\text{NiPt}(\text{CN})_4$,
 $2\text{NH}_3 + \text{H}_2\text{O.}$

Potassium platinocyanide, $\text{K}_2\text{Pt}(\text{CN})_4 +$
 $3\text{H}_2\text{O.}$

Extremely efflorescent. Sl. sol. in cold,
easily in hot $\text{H}_2\text{O.}$ (Willm, B. 19. 950.)
Sol. in alcohol and ether.

Potassium sodium platinocyanide, $\text{K}_2\text{Pt}(\text{CN})_4$,
 $\text{Na}_2\text{Pt}(\text{CN})_4 + 6\text{H}_2\text{O.}$

Sol. in $\text{H}_2\text{O.}$ (Willm, B. 19. 950.)

Praseodymium platinocyanide,
 $2\text{Pr}(\text{CN})_3, 3\text{Pt}(\text{CN})_2$.

Sol. in $\text{H}_2\text{O.}$ (Von Scheele, Z. anorg. 1898,
18. 355.)

Samarium platinocyanide, $\text{Sm}_2[\text{Pt}(\text{CN})_4]_2 + 18\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve.)

Scandium platinocyanide, $\text{Sc}_2[\text{Pt}(\text{CN})_4]_2 + 21\text{H}_2\text{O}$.

Sol. in H_2O and insol. in alcohol; when boiled in alcohol it is dehydrated. (Crookes, Phil. Trans. 1910, 210. A, 368.)

+21 H_2O . (Orlow, Ch. Z. 1912, 36. 1407.)

Silver platinocyanide, $\text{Ag}_2\text{Pt}(\text{CN})_4$.

Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Silver platinocyanide ammonia, $\text{Ag}_2\text{Pt}(\text{CN})_4 \cdot 2\text{NH}_3$.

Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Silver platinocyanide bromide.

See Bromoplatinocyanide, silver.

Silver platinocyanide chloride.

See Chloroplatinocyanide, silver.

Silver platinocyanide iodide.

See Iodoplatinocyanide, silver.

Sodium platinocyanide, $\text{Na}_2\text{Pt}(\text{CN})_4 + 3\text{H}_2\text{O}$.

Easily sol. in H_2O . (Willm, Z. anorg. 4. 298.)

Sol. in alcohol.

Strontium platinocyanide, $\text{SrPt}(\text{CN})_4 + 5\text{H}_2\text{O}$.

Sol. in H_2O .

Thallous platinocyanide, $\text{Tl}_2\text{Pt}(\text{CN})_4$.

Nearly insol. in cold, sl. sol. in hot H_2O . (Friswell, Chem. Soc. 24. 461.)

Thallous platinocyanide carbonate, $2\text{Tl}_2\text{Pt}(\text{CN})_4 \cdot \text{Tl}_2\text{CO}_3$.

Nearly insol. in cold H_2O . (F.)

Thorium platinocyanide, $\text{Th}[\text{Pt}(\text{CN})_4]_2 + 16\text{H}_2\text{O}$.

Somewhat difficultly sol. in cold, easily in hot H_2O . (Cleve, Sv. V. A. H. Bih. 2. No. 6.)

Uranyl platinocyanide, $(\text{UO}_2)\text{Pt}(\text{CN})_4 + x\text{H}_2\text{O}$.

Sol. in H_2O . (Levy, Chem. Soc. 1908, 93. 1459.)

Ytterbium platinocyanide, $2\text{Yb}(\text{CN})_3 \cdot 3\text{Pt}(\text{CN})_4 + 18\text{H}_2\text{O}$.

Easily sol. in H_2O . (Cleve, Z. anorg. 1902, 32. 139.)

Yttrium platinocyanide, $\text{Y}_2[\text{Pt}(\text{CN})_4]_2 + 21\text{H}_2\text{O}$.

Easily sol. in H_2O . Insol. in absolute alcohol. (Cleve and Höglund.)

Zinc platinocyanide ammonia, $\text{ZnPt}(\text{CN})_4 \cdot 2\text{NH}_3 + \text{H}_2\text{O}$.

Platinonitrous acid.

See Platonitrous acid.

Platinoplatinicyanhydric acid, $\text{HPt}(\text{CN})_4 + x\text{H}_2\text{O}$.

Sol. in H_2O . (Levy, Chem. Soc. 1912, 101. 1093.)

Platinoselenocyanhydric acid.

Potassium platinoselenocyanide, $\text{K}_2\text{Pt}(\text{SeCN})_4$.

Sol. in H_2O and alcohol. (Clarke and Daley, B. 1878, 11. 1325.)

Platinoselenostannic acid.

See under Selenostannate, platinum.

Platinosulphocyanhydric acid, $\text{H}_2\text{Pt}(\text{SCN})_4$.

Known only in aqueous solution.

Potassium platinosulphocyanide, $\text{K}_2\text{Pt}(\text{SCN})_4$.

Permanent. Sol. in 2.5 pts. H_2O at 15°, and more readily at higher temp. Very sol. in warm alcohol.

Silver —, $\text{Ag}_2\text{Pt}(\text{SCN})_4$.

Insol. in H_2O . Sol. in $\text{KSCN} + \text{Aq}$, and partly sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Platinosulphocyanhydric acid, $\text{H}_2\text{Pt}(\text{SCN})_4$.

Known only in aqueous, and alcoholic solutions.

Ammonium platinosulphocyanide, $(\text{NH}_4)_2\text{Pt}(\text{SCN})_4$.

Sol. in H_2O and alcohol.

Barium —, $\text{BaPt}(\text{SCN})_4$.

Sol. in H_2O and alcohol.

Ferrous —, $\text{FePt}(\text{SCN})_4$.

Insol. in H_2O or alcohol. Not attacked by dil. H_2SO_4 , HCl , or $\text{HNO}_3 + \text{Aq}$.

Lead —, $\text{PbPt}(\text{SCN})_4$.

Sl. sol. in cold, decomp. by hot H_2O . Sol. in alcohol.

$\text{PbPt}(\text{SCN})_4$, PbO . Insol. in H_2O or alcohol. Sol. in acetic or nitric acids.

Mercurous —, $\text{Hg}_2\text{Pt}(\text{SCN})_4$.

Ppt. Insol. in H_2O .

platinosulphocyanide,
 $\text{Pt}(\text{SCN})_4$.

2 pts H_2O at 60° . Much more
sol. in H_2O , and still more easily in
1.

(Miolati and Bellucci, *Gazz.*
30, II. 592.)

-, $\text{Ag}_2\text{Pt}(\text{SCN})_4$.

H_2O or $\text{K}_2\text{Pt}(\text{SCN})_4 + \text{Aq.}$ Sol.
 $\text{OH} + \text{Aq}$ and in $\text{KCNS} + \text{Aq.}$

—, $\text{Na Pt}(\text{SCN})_4$.

I_2O and alcohol.

phostannic acid.

7 Sulphostannate, platinum.

phurous acid.**osulphurous acid.**

Pt.

acked by H_2O , H_2SO_4 , HCl , or
1. Slowly sol. in aqua regia, or a
 HBr and HNO_3 , but much less
Au.

uted Pt is remarkably sol. in
n presence of air. (Wilm, B. 1881,

foil is attacked by fuming HCl
ence of light, but not in the dark.
C. R. 1904, **138**. 1297.)

l + Aq dissolves 10–15% Pt from
lack. (Wöhler, B. 1903, **36**. 3482.)
 NO_2 oxidizes Pt black; Pt sponge,
; sheet Pt, slightly. (Wöhler,
11.)

essence of Hg is more or less sol.
 NO_2 . (Tarugi, *Gazz. ch. it.* 1903,

als are attacked by evaporating
rein. (Jaunek and Meyer, Z.
3, **83**. 71.)

in conc. H_2SO_4 containing small
of nitrogen oxides. (Scheurer-
R **86**. 1082.)

., pptd. by formic acid, is easily
ling H_2SO_4 . (Deville and Stas,
.)

et Pt is attacked by boiling H_2SO_4
 K_2SO_4 ; 1 sq. cm. loses 0.01 g.
and velocity of the reaction is not
by addition of HNO_3 . Pt black
ly dissolved under the above con-
50 hours. (Delépine, C. R. 1905,

data on solubility of Pt in H_2SO_4
by Delépine. (C. R. 1906, **142**.

SO_4 dissolves 0.04 g. Pt from com-
at 250 – 280° in 28 hours. (Con-
Chem. Ind. 1903, **22**. 465.)

Quenessen. (Bull. Soc. 1906,
.)

0.0038 g. is dissolved by 10 cc. of boiling
 H_2SO_4 . (McCoy, Eighth Inter. Cong. App.
Chem. **1912**. 2)

$\text{HCl} + \text{HNO}_3$, so long as they are sufficiently
dil. or the temperature is so low that they
cannot react on each other, have no action on
Pt. Addition of Cl does not bring about re-
action, but a few drops of KNO_3 or $\text{N}_2\text{O}_5 + \text{Aq}$
bring about an immediate reaction. (Millon.)

Slowly sol in $\text{HI} + \text{Aq.}$ (Deville, C. R. **42**.
896.)

Conc. H_3PO_4 attacks Pt when heated in
presence of air, but not in its absence. (Hütt-
ner, Z. anorg. 1908, **59**. 216.)

Pt dissolves easily in most acids when they
contain H_2O_2 . (Fairley, B. 1875, **8**. 1600.)

Slowly sol. in boiling $\text{FeCl}_3 + \text{Aq}$ (Saint-
Pierre, C. R. **54**. 1077.)

FeCl_3 in acid solution is without influence
on Pt. (Marie, C. R. 1908, **146**. 476.)

Pt is completely insol. in $\text{KCN} + \text{Aq.}$
(Rösler, Z. Chem. **1886**. 175.)

Pt is attacked by boiling conc. $\text{KCN} + \text{Aq.}$
(Deville and Debray, C. R. **82**. 241.)

Solubility of Pt in 10% $\text{KCN} + \text{Aq}$ is very
small at ord. temp. (1.4 mg. in 8 days) but
is considerably greater in boiling conc. KCN
+ Aq. (71.5 mg. in 5 hours). (Glaser, Z.
Elektrochem. 1903, **9**. 15.)

Pt foil is dissolved in boiling $\text{KCN} + \text{Aq}$
(0.030 g for 1 cc. in 1 hour). Insol. in cold
 $\text{KCN} + \text{Aq.}$ (Brochet and Petit, C. R. 1904,
C. R. **138**. 1255.)

Sol in $\text{RbCl}_3 + \text{Aq.}$ (Erdmann, Arch.
Pharm 1894, **232**. 30)

Insol. in liquid NH_3 . (Gore, Am. Ch. J.
1898, **20**. 828.)

Platinum ammonium compounds.

Data published since the first edition of this
work have not been included in this edition.

See —

Platosamine comps., $\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ R} \\ \text{NH}_3 \text{ R} \end{smallmatrix}$.

Platosemidiamine comps.,
 $\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ NH}_3 \text{ R} \\ \text{R} \end{smallmatrix}$.

Platomonodiamine comps.,
 $\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ NH}_3 \text{ R} \\ \text{NH}_3 \text{ R} \end{smallmatrix}$.

Platodiamine comps., $\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ NH}_3 \text{ R} \\ \text{NH}_3 \text{ NH}_3 \text{ R} \end{smallmatrix}$.

Platososemidiamine comps., $\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ R} \\ \text{R} \end{smallmatrix}$.

Diplatodiamine comps.,
 $\text{Pt} - \text{NH}_3 \text{ NH}_3 \text{ R}$

$\text{Pt} - \text{NH}_3 \text{ NH}_3 \text{ R}$
Bromoplatinamine comps.,

$\text{Br}_2\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ R} \\ \text{NH}_3 \text{ R} \end{smallmatrix}$.

Chloroplatinamine comps.,
 $\text{Cl}_2\text{Pt} < \begin{smallmatrix} \text{NH}_3 \text{ R} \\ \text{NH}_3 \text{ R} \end{smallmatrix}$.

Chloronitratoplatinamine comps.,
 $\text{Cl}(\text{NO}_3)\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Iodoplatinamine comps., $\text{I}_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Hydroxyloptatinamine comps.,
 $(\text{OH})_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Nitratoplatinamine comps.,
 $(\text{NO}_3)_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Sulphatoplatinamine comps.,
 $\text{SO}_4\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Bromoplatinsemidiamine comps.,
 $\text{Br}_2\text{PtNH}_2\text{.NH}_2\text{.R}$

Bromonitritoplatinsemidiamine comps.,
 $\text{Br}_2(\text{NO}_2)\text{PtNH}_2\text{.NH}_2\text{.R}$

Chloroplatinsemidiamine comps.,
 $\text{Cl}_2\text{PtNH}_2\text{.NH}_2\text{.R}$

Chlorohydroxylonitritosemidiamine comps.,
 $\text{Cl}(\text{OH})(\text{NO}_2)\text{PtNH}_2\text{.NH}_2\text{.R}$

Chloronitritoplatinsemidiamine comps.,
 $\text{Cl}_2(\text{NO}_2)\text{PtNH}_2\text{.NH}_2\text{.R}$

Iodoplatinsemidiamine comps.,
 $\text{I}_2\text{PtNH}_2\text{.NH}_2\text{.R}$

Hydroxylosemidiamine comps.,
 $(\text{OH})_2\text{PtNH}_2\text{.NH}_2\text{.R}$

Bromoplatinmonodiamine comps.,
 $\text{Br}_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Bromohydroxyloptatinmonodiamine comps.,
 $\text{Br}(\text{OH})\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Chloroplatinmonodiamine comps.,
 $\text{Cl}_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Iodonitratoplatinmonodiamine comps.,
 $\text{I}(\text{NO}_3)\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Hydroxyloptatinmonodiamine comps.,
 $(\text{OH})_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Bromoplatindiamine comps.,
 $\text{Br}_2\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.NH}_2\text{.R} \end{smallmatrix}$

Bromocarbonatoplatindiamine comps.,
 $\text{CO}_3 > [\text{Pt}(\text{NH}_2)\text{R}]_2$
 Br_2

Bromochloroplatindiamine comps.,
 $\text{BrClPt}(\text{NH}_2)_4\text{R}_2$

Bromohydroxyloptatindiamine comps.,
 $\text{Br}(\text{OH})\text{Pt}(\text{NH}_2)_4\text{R}_2$

Bromonitratoplatindiamine comps.,
 $\text{Br}(\text{NO}_3)\text{Pt}(\text{NH}_2)_4\text{R}_2$

Bromosulphatoplatindiamine comps.,
 $\text{Br}_2(\text{SO}_4)[\text{Pt}(\text{NH}_2)_4\text{R}_2]_2$

Carbonatochloroplatindiamine comps.,
 $(\text{CO}_3)\text{Cl}_2[\text{Pt}(\text{NH}_2)_4\text{R}_2]_2$

Carbonatoplatindiamine comps.,
 $(\text{CO}_3)(\text{NO}_3)_2[\text{Pt}(\text{NH}_2)_4\text{R}_2]_2$

Chloroplatindiamine comps.,
 $\text{Cl}_2\text{Pt}(\text{NH}_2)_4\text{R}_2$

Chlorohydroxyloptatindiamine comps.,
 $\text{Cl}(\text{OH})(\text{NH}_2)_4\text{R}_2$

Chloriodoplatindiamine comps.,
 $\text{ClIPt}(\text{NH}_2)_4\text{R}_2$

Chloronitratoplatindiamine comps.,
 $\text{Cl}(\text{NO}_3)\text{Pt}(\text{NH}_2)_4\text{R}_2$

Hydroxyloptatindiamine comps.,
 $(\text{OH})_2\text{Pt}(\text{NH}_2)_4\text{R}_2$

Hydroxylonitratodiamine comps.,
 $(\text{OH})(\text{NO}_3)\text{Pt}(\text{NH}_2)_4\text{R}_2$

Hydroxylosulphatodiamine comps.,
 $(\text{OH})_2\text{SO}_4[\text{Pt}(\text{NH}_2)_4\text{R}_2]_2$

Iodoplatindiamine comps., $\text{I}_2\text{Pt}(\text{NH}_2)_4\text{R}_2$

Iodonitritoplatindiamine comps.,
 $\text{I}(\text{NO}_2)\text{Pt}(\text{NH}_2)_4\text{R}_2$

Nitratoplatindiamine comps.,
 $(\text{NO}_3)_2\text{Pt}(\text{NH}_2)_4\text{R}_2$

Nitritoplatindiamine comps.,
 $(\text{NO}_2)_2\text{Pt}(\text{NH}_2)_4\text{R}_2$

Sulphatoplatindiamine comps.,
 $(\text{SO}_4)\text{Pt}(\text{NH}_2)_4\text{R}_2$

Iododiplatinamine comps.,
 $\text{I}-\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$
 $\text{I}-\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.R} \\ \text{NH}_2\text{.R} \end{smallmatrix}$

Bromodiplatindiamine comps.,
 $\text{Br}-\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.NH}_2\text{.R} \end{smallmatrix}$
 $\text{Br}-\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.NH}_2\text{.R} \end{smallmatrix}$

Hydroxyloptatindiamine comps.,
 $(\text{OH})_2\text{Pt}_2(\text{NH}_2)_8\text{R}_4$

Iododiplatindiamine comps.,
 $\text{I}_2\text{Pt}_2(\text{NH}_2)_8\text{R}_4$

Nitratodiplatindiamine comps.,
 $(\text{NO}_3)_2\text{Pt}_2(\text{NH}_2)_8\text{R}_4$

Platintriamine comps.,
 $\text{R}_3\text{Pt} < \begin{smallmatrix} \text{NH}_2\text{.NH}_2\text{.NH}_2\text{.R} \\ \text{NH}_2\text{.NH}_2\text{.NH}_2\text{.R} \end{smallmatrix}$

Tetraplatinamine comps., $\text{Pt}_4(\text{NH}_2)_{12}\text{R}_{12}$

Octoplatinamine comps., $\text{Pt}_8(\text{NH}_2)_{24}\text{R}_{24}$

Platinum antimonide, PtSb_3 .
 (Christofle, 1863.)

Platinum arsenide, Pt_3As_2 .
 (Tivoli, Gazz. ch. it. 14. 487.)
 PtAs_2 . Min. *Sperrylite*. Sl. attacked
 aqua regia. (Wells, Sill. Am. J. (3) 37.)

Platinum arsenic hydroxide (?), PtAsOH
 Insol. in, and slowly decomp. by H_2O
 alcohol. Easily decomp. by $\text{HCl} + \text{Aq}$;
 attacked by $\text{HNO}_3 + \text{Aq}$. Sol. in aqua r
 not attacked by cold conc. H_2SO_4 , but
 comp. on heating. (Tivoli, Gazz. ch. it.
 487.)

m potassium azoimide.

Explodes violently even in aq. solution.
Curtius, J. pr. 1898, (2) 58. 304.)

m boride, Pt_2B_2 .

slowly sol. in aqua regia. (Martius, 79.)

us bromide, $PtBr_2$.

sl. sol. in H_2O . Sol. in $HBr + Aq$. Sl. sol. in $H_2O + Aq$. (Topsoë, J. B. 1868. 274.)

us bromide, $PtBr_4$.

deliquescent; sol. in H_2O . (Meyerblin, B. 13. 404.)

sl. sol. in H_2O . 100 g. $PtBr_4 + Aq$ sat. at 20°C. contain 0.41 g. $PtBr_4$. (Halberstadt, B. 2.)

sl. sol. in $HBr + Aq$; sl. sol. in $HC_2H_3O_2$. Sol. in considerable amount in K or Na acetate + Aq .

sl. sol. in alcohol or ether, also in $CHCl_3$. (Halberstadt.)

us hydrogen bromide.**us bromoplatinic acid.****us bromide carbonyl.****us carbonyl platinous bromide.****us bromide with MBr .****us bromoplatinate, M .****m carbide, PtC_2 .**

aqua regia dissolves out nearly all the platinum. (Lassaigne, J. pr. 20. 209.)

m carbon disulphide, $PtCS_2$.**Platinum sulphocarbide.****m monochloride, $PtCl + xH_2O$.**

sl. sol. in HCl ; mod. sol. in hot dil. HCl without decomp. (Sonstadt, Proc. Soc. 1898, 14. 179.)

us chloride, $PtCl_2$.

sl. sol. in H_2O , conc. H_2SO_4 , or HNO_3 . Sol. in $HCl + Aq$ with exclusion of air. (Berthelot, J. pr. 20. 209.)

sl. sol. in alcohol or ether; sol. in $NH_4OH + H_2O$. (Raewsky, A. ch. (3) 22. 280.) Sol. in H_2O with formation of $PtCl_4$.

sl. sol. in cold conc. $KI + Aq$, but sol. when heated. (Lassaigne, A. ch. (2) 51. 117.)

sl. sol. in liquid NH_3 . (Gore, Am. Ch. J. 1900, 22. 828.)

sl. sol. in acetone. (Eidmann, C. C. 1899, 14.)

m trichloride, $PtCl_3$.

sl. sol. in cold, more sol. in hot H_2O . Partially hydrolyzed by boiling with H_2O .

Insol. in cold conc. HCl . Sol. in hot conc. HCl with decomp.

Sol. in $KI + Aq$. (Wöhler, B. 1909, 42. 3961.)

Platinic chloride, $PtCl_4$.

Not deliquescent. Very sol. in H_2O . (Pulinger, Chem. Soc. 61. 420.)

Sp. gr. of aqueous solution containing:

5	10	15	20	25	% $PtCl_4$
1.046	1.097	1.153	1.214	1.285	
30	35	40	45	50	% $PtCl_4$
1.362	1.450	1.546	1.666	1.785	

(Precht, Z. anal. 18. 512.)

Insol. in conc. H_2SO_4 . (Dumas.)

Sl. sol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 828.)

Sol. in alcohol and ether; sol. in anhydrous acetone. (Zeise, A. 33. 34.)

Insol. in ether. (Willstätter, B. 1903, 36. 1830.)

Sl. sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4328.)

+ H_2O . Sol. in H_2O . (Gutbier and Heinrich, Z. anorg. 1913, 81. 378.)

+ $4H_2O$. Sol. in H_2O . (Pigeon, C. R. 1891, 112. 792.)

+ $5H_2O$. Not deliquescent. Sol. in H_2O or $HCl + Aq$.

Composition is probably $H_2PtCl_4O + 4H_2O$. (Norton, J. pr. 110. 469.)

+ $7H_2O$. Sol. in H_2O . (Pigeon.)

+ $8H_2O$. (Blondel, A. Ch. 1905, (8) 6. 98.)

Platinic thallium chloride, $Tl_2Pt_2Cl_4H_2O_4$.

Ppt.; insol. in H_2O . (Miolati, Z. anorg. 1900, 22. 460.)

Platinous hydrogen chloride.

See Chloroplatinous acid.

Platinic hydrogen chloride.

See Chloroplatinic acid.

Platinous chloride with MCl .

See Chloroplatinite, M .

Platinic chloride with MCl .

See Chloroplatinate, M .

Platinous phosphorus chloride.

See Phosphorus platinous chloride.

Platinic phosphorus chloride.

See Phosphorus platinic chloride.

Platinous chloride carbonyl.

See Carbonyl platinous chloride.

Platinum chloride hydroxylamine,
 $\text{Pt}(\text{NH}_2\text{OH})_2\text{Cl}_2$.

Ppt. Sol. in alcohol and in ether. Decomp. in aq. solution.

$\text{Pt}(\text{NH}_2\text{OH})_2\text{Cl}_2$. Sol. in H_2O . (Uhlenbuth, A. 1900, 311. 124.)

Platinous chloride sulphocarbamide,
 $\text{PtCl}_2 \cdot 4\text{CS}(\text{NH}_2)_2$.

Sl. sol. in H_2O ; very sol. in hot H_2O ; decomp. sl. on boiling. (Kurnakow, J. pr. 1894, (2) 50. 483.)

Platinum chloriodide, PtCl_2I_2 .

Very deliquescent. (Kämmerer, A. 148. 329.)

PtClI_2 . Insol. in H_2O . Sl. sol. in alcohol. Sol. in $\text{KOH} + \text{Aq}$, from which it is pptd. by H_2SO_4 . (Mather, Sil. Am. J. 27. 257.)

Platinum chloronitride, PtNCl .

(Alexander, C. C. 1887. 1254.)

Platinous cyanide with MCN .

See **Platinocyanide, M.**

Platinous fluoride, $\text{PtF}_2(?)$.

Insol. in H_2O . (Moissan, A. ch. (6) 34. 287.)

Platinic fluoride, PtF_4 .

Deliquescent. Sol. in H_2O with immediate decomp. into PtO_4H_4 and HF . (Moissan, C. R. 109. 807.)

Platinous hydroxide, PtO_2H_2 .

Sol. in HCl , HBr , and $\text{H}_2\text{SO}_4 + \text{Aq}$, but not in other oxygen acids. Decomp. by boiling $\text{KOH} + \text{Aq}$. (Thomsen, J. pr. (2) 16. 344.)

When freshly pptd., is insol. in dil. HNO_3 and H_2SO_4 , and in $\text{HC}_2\text{H}_3\text{O}_2$; sol. in conc. HNO_3 and H_2SO_4 . Very sol. in H_2SO_4 and HCl . After drying, is insol. in conc. HNO_3 and H_2SO_4 . (Wöhler, Z. anorg. 1904, 40. 424.)

Platinic hydroxide, $\text{Pt}(\text{OH})_4$.

Easily sol. in dil. acids and in $\text{NaOH} + \text{Aq}$. (Topsoë, J. B. 1870. 386.)

Nearly insol. in acetic acid. (Döbereiner.)

Insol. in all acids except conc. HCl and aqua regia. (Wöhler, Z. anorg. 1904, 40. 438.)

+ H_2O . Ppt. (Prost, Bull. Soc. (2) 44. 256.)

Insol. in $2\text{N}-\text{H}_2\text{SO}_4$ and dil. $\text{HNO}_3 + \text{Aq}$; mod. sol. in conc. HNO_3 , H_2SO_4 , $2\text{N}-\text{HCl}$ and $\text{NaOH} + \text{Aq}$. (Wöhler.)

+ $2\text{H}_2\text{O}$. Easily sol. in dil. acids, even acetic acid, and in $\text{NaOH} + \text{Aq}$. (Topsoë.)

Insol. in acetic acid; sl. sol. in $2\text{N}-\text{H}_2\text{SO}_4$ and HNO_3 ; easily sol. in HCl , and $\text{NaOH} + \text{Aq}$. (Wöhler.)

Platinoplatinic hydroxide, $\text{Pt}_2\text{O}_3 \cdot 9\text{H}_2\text{O}$

Ppt. (Prost, Bull. Soc. (2) 44. 156.)

$\text{Pt}_2\text{O}_{11} \cdot 11\text{H}_2\text{O}$. Ppt. (Prost.)

Platinum hydroxylamine comp.

See—

Platodioxamine comp., $\text{Pt}(\text{NH}_2\text{O})_2$

Platosoxamine comp., $\text{Pt}(\text{NH}_2\text{O})_2$

Platosoxamine-amine comp.,

$\text{Pt}(\text{NH}_2\text{O})_2\text{NH}_2\text{R}_1$.

Platinous iodide, PtI_2 .

Insol. in H_2O , acids, or alcohol. (Lassaigne, A. ch. (2) 51. 113.)

Difficultly sol. in $\text{Na}_2\text{SO}_4 + \text{Aq}$. (Topsoë.) Gradually decomp. by hot $\text{HI} + \text{Aq}$ of 1 sp. gr., also by hot $\text{KI} + \text{Aq}$, PtI_2 being solved out and Pt left behind. Not att. by conc. H_2SO_4 , HCl , or $\text{HNO}_3 + \text{Aq}$, gradually decomp. by KOH or $\text{NaOH} + \text{Aq}$. (Lassaigne.)

Insol. in acetone. (Eidmann, C. C. 111. 1014.)

Platinic iodide, PtI_4 .

Insol. in H_2O . Sol. in NaOH or $\text{NaCl} + \text{Aq}$, H_2SO_4 , or $\text{Na}_2\text{SO}_4 + \text{Aq}$. Sol. in HI or alkali iodides + Aq . Sol. in alcohol, partial decomp. Not attacked by H_2O . (Lassaigne, A. ch. (2) 51. 122.)

Very sol. in liquid NH_3 . (Gore, Am. C. 1898, 20. 828.)

Sol. in alcohol. (Belluci, C. C. 100. 625.)

Platinic iodide with MI .

See **Iodoplatinate, M.**

Platinum nitride chloride, PtNCl .

See **Platinum chloronitride.**

Platinous oxide, PtO .

Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. Insol. in other acids. (Döbereiner, Pogg. 28. 183.)

Sol. in conc. H_2SO_4 ; easily in conc. $\text{HCl} + \text{Aq}$. (Storer's Diet.)

Very sl. sol. in $\text{HCl} + \text{Aq}$. Sl. sol. in regia. (Wöhler, B. 1903, 36. 3482.)

Platinic oxide, PtO_2 .

Insol. in acids, even aqua regia. (Wöhler, 1901.)

Platinum trioxide, PtO_3 .

Unattacked by dil. H_2SO_4 , HNO_3 , or HCl . Sol. with decomp. in dil. and conc. H_2SO_4 or conc. HNO_3 . (Wöhler, 1909, 42. 3329.)

Platinum sesquioxide, $\text{Pt}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Insol. in dil. sol. in conc. H_2SO_4 . Sl. sol. in hot dil. HNO_3 . Sol. in HCl . Sol. in alkali hydroxides + Aq . (Wöhler, B. 1904. 3964.)

+2H₂O. Insol. in HNO₃ and H₂SO₄. Insol. in alkalies + Aq, also cold dil. HCl + Aq. Sol. in aqua regia. (Dudley, Am. Ch. J. 1902, 28. 66.)
+5H₂O. (Delépine, Bull. Soc. 1910, (4) 7. 103.)

Platinoplatinic oxide, Pt₃O₄.

Not attacked by long boiling with HCl, HNO₃, or aqua regia. (Jørgensen, J. pr. (2) 16. 344.)

Does not exist. (Wöhler, Z. anorg. 1904, 40. 450.)

Platinum oxychloride, 3PtO, PtCl₂ (?).

Sol. in HCl, and in KOH + Aq. (Kane, Phil. Trans. 1842. 298.)

PtCl₂(OH)₂ = H₂PtCl₂O₂. (Jørgensen, J. pr. (2) 16. 345.)

5PtO₂, 2HCl + 9H₂O. Insol. in cold H₂O; decomp. on boiling. Slowly sol. in HCl. (Blondel, A. ch, 1905, (8) 6. 100.)

Platinum oxysulphide, PtOS.

See Platinum sulphhydroxide.

Platinum phosphide, PtP₂.

Insol. in HCl + Aq. Sol. in aqua regia. (Schrötter, W. A. B. 1849. 303.)

PtP₂H₂. Insol. in H₂O, and HCl + Aq. (Cavazzi, Gazz. ch. it. 13. 324.)

PtP. Insol. in aqua regia. (Clark and Joslin.)

Pt₂P. Sol. in aqua regia. (Clark and Joslin.)

Pt₃P₂. Partially sol. in aqua regia. (Clark and Joslin, C. N. 48. 385.)

Attacked very slowly by aqua regia. Rapidly sol. in molten alkalies. (Granger, C. N. 1898, 77. 229.)

Completely sol. in aqua regia if the action sufficiently prolonged, though with difficulty. (Granger, C. R. 1898, 123. 1285.)

Platinum diselenide, PtSe₂.

As PtS₂. (Minozzi, Chem. Soc. 1909, 96. (2) 899.)

Platinum triselenide, PtSe₃.

Sl. attacked by hot conc. HNO₃; not attacked by cold conc. HCl + Aq; slowly sol. in aqua regia and Cl₂ + Aq; insol. in CS₂.

Platinum silicide, Pt₂Si.

Sol. in hot aqua regia. (Vigouroux, C. R. 1896, 123. 117.)

Pt₂Si₂. (Colson, C. R. 94. 27.)

Pt₂Si₂. Slowly decomp. by aqua regia. (Guyard, Bull. Soc. (2) 25. 511.)

PtSi. Insol. in HNO₃, H₂SO₄, HF, and HCl. Completely sol. in aqua regia. (Lebeau and Novitsky, C. R. 1907, 145. 241.)

Platinum sulphhydroxide, PtOS + H₂O = PtS(OH)₂.

Decomp. easily into—

$\text{Pt}_2\text{S}_2\text{O}_3\text{H}_2 = \frac{\text{PtS}}{\text{PtS}} \frac{\text{O}}{\text{O}} \frac{\text{OH}}{\text{OH}} = \text{PtOS} + \frac{1}{2}\text{H}_2\text{O}$. H₂O cannot be removed without decomposing the compound. (v. Meyer, J. pr. (2) 15. 1.)

Platinous sulphide, PtS.

Not attacked by boiling acids, aqua regia, or KOH + Aq. (Böttger, J. pr. 2. 274.)

Sol. in large excess of (NH₄)₂S + Aq.

Platinoplatinic sulphide, Pt₂S₃.

Not attacked by HCl or HNO₃ + Aq, and only slowly by aqua regia. (Schneider, Pogg. 138. 607.)

Platinic sulphide, PtS₂.

Anhydrous. Aqua regia attacks sl., other acids not at all. (Davy.)

Hydrated Insol. in HCl + Aq; sl. sol. in boiling HNO₃ + Aq. Sol. in aqua regia. (Fresenius.) Sol. in alkali sulphides, hydrates and carbonates + Aq. (Berzelius.) Very sl. sol. in (NH₄)₂S + Aq. (Claus)

Insol. in NH₄Cl, or NH₄NO₃ + Aq.

1 pt. PtCl₄ in 100 pts. H₂O + 25 pts. HCl is not pptd. by H₂S. (Reinsch.)

Difficultly sol. in alkali sulphhydroxides + Aq, but more easily in presence of SnS, Sb₂S₃, As₂S₃, or SnS₂. (Ribau, C. R. 85. 283.)

Platinum sulphide, Pt₂S₃, or Tetraplatinum sulphoplatinate, 4PtS, PtS₂.

Decomp. on moist air, but not attacked by acids. (Schneider, J. pr. (2) 7. 214.)

Platinum sulphides with M₂S.

See Sulphoplatinate, M.

Platinum sulphocarbide, PtC₂S₂.

Not attacked by hot HCl, HNO₃ + Aq, slightly by aqua regia. (Schützenberger, C. R. 111. 391.)

Platinum telluride, PtTe.

Decomp. by fused oxidizing agents; slowly sol. in conc. HNO₃. (Roessler, Z. anorg. 1897, 15. 407.)

Platinum ditelluride, PtTe₂.

Insol. in boiling conc. KOH + Aq; slowly sol. in boiling conc. HNO₃; decomp. by fused oxidizing agents. (Roessler.)

Plato-.

See also Platino-.

Platoamidosulphonic acid.

Potassium platoamidosulphonate,
 $K_2Pt(NH_2SO_3)_4 + 2H_2O$.

Very sl. sol. in cold H_2O ; sol. in 10 pts. boiling H_2O . (Ramberg and St. Kahlenberg, B. 1912, 45. 1514.)

Sodium —, $Na_2(Pt(NH_2SO_3)_4 + 4H_2O$.

Very sol. in H_2O . (Ramberg and St. Kahlenberg.)

Platodiamine bromide, $Pt[(NH_2)_2Br]_2 + 3H_2O$.

Easily sol. in H_2O . (Cleve.)

— **carbonate, $Pt(N_2H_4)_2CO_3 + H_2O$.**

Sol. in H_2O . (Peyrone, A. 51. 14.)

$Pt(N_2H_4CO_2H)_2$. Sl. sol. in, but decomp. by boiling with H_2O into—

— **sesquicarbonate.**

More sol. than preceding salt. (Reiset, C. R. 11. 711.)

— **chloride, $Pt[(NH_2)_2Cl]_2 + H_2O$.**

"Reiset's first chloride." Sol. in 4 pts. H_2O at 16.5° , and in less hot H_2O . Insol. in alcohol or ether. (Reiset, A. ch. (3) 11. 419.)

As sol. in $NH_4Cl + Aq$ as in H_2O ; insol. in absolute alcohol; sl. sol. in dil. alcohol; very sol. in dil. $HCl + Aq$. (Peyrone, A. ch. (3) 12. 196.)

— **cuprous chloride, $Pt(NH_2)_4Cl_2, Cu_2Cl_2$.**

Sol. in H_2O , and pptd. from H_2O solution by alcohol. (Buckton.)

— **cupric chloride, $Pt(NH_2)_4Cl_2, CuCl_2$.**

Sl. sol. in cold, decomp. by hot H_2O into $Pt(NH_2)_4Cl_2, Cu_2Cl_2$. (Buckton, Chem. Soc. 5. 218.)

Nearly insol. in H_2O ; easily sol. in warm $HCl + Aq$; insol. in alcohol. (Millon and Commaile, C. R. 57. 822.)

Millon and Commaile's salt is $Cu(NH_2)_4Cl_2, PtCl_2$, cuprammonium chloroplatinite.

— **lead chloride, $Pt(NH_2)_4Cl_2, PbCl_2$.**

Sol. in hot, much less in cold H_2O . Insol. in $HCl + Aq$ or alcohol. (Buckton, Chem. Soc. 5. 213.)

— **mercuric chloride, $Pt(NH_2)_4Cl_2, HgCl_2$.**

Easily sol. in hot H_2O , much less in cold. Insol. in $HCl + Aq$. (Buckton.)

— **zinc chloride, $Pt(NH_2)_4Cl_2, ZnCl_2$.**

Easily sol. in hot H_2O . Insol. in alcohol. (Buckton.)

Platodiamine chloroplatinate, $Pt(NH_2)_4PtCl_4$.

Ppt. Insol. in H_2O . (Cossa, Gazz. 17. 1.)

— **chloroplatinite, $Pt(NH_2)_4Cl_2, PtCl$**
 (Magnus' green salt.) Insol. in, and comp. by $H_2O, HCl + Aq$, or alcohol.

Slowly sol. in boiling $NH_4OH + Aq$ conc. NH_4 salts + Aq . (Reiset, A. ch. 427.)

Almost as sol. in $(NH_4)_2CO_3 + Aq$ $NH_4OH + Aq$. Sol. in hot $PtCl_4 + Aq$ set.)

Not decomp. by boiling KOH , dil. $H_2SO_4 + Aq$, but easily by HNO_3 (Gros, A. 27. 245.)

— **chromate, $Pt(NH_2)_4CrO_4$.**

Scarcely sol. in H_2O . (Cleve.)

— **dichromate, $Pt(NH_2)_4Cr_2O_7$.**

Sl. sol. in H_2O . Insol. in alcohol. $KOH + Aq$. (Buckton, Chem. Soc. 1)

— **platinous cyanide, $Pt(NH_2)_4(CN)_2$.**

Sl. sol. in cold, easily in boiling H_2O in KOH, HCl , and dil. $H_2SO_4 + Aq$ decomp., but conc. H_2SO_4 decomposes.

— **potassium ferrocyanide, $Pt(NH_2)_4K_2[Fe(CN)_6] + 3H_2O$.**

— **hydroxide, $Pt[(NH_2)_2OH]_2$.**

"Reiset's first base." Easily sol. Sl. sol. in alcohol.

— **iodide, $Pt[(NH_2)_2I]_2$.**

Sl. sol. in cold, more easily in hot slowly decomp. on boiling. (Reiset.)

— **nitrate, $Pt[(NH_2)_2NO_3]_2$.**

Sol. in about 10 pts. boiling H_2O . but sl. sol. in alcohol. (Peyrone, J. 12. 203.)

— **nitrate sulphate, $[Pt(NH_2)_4Pt(NH_2)_4SO_4]$.**

Very easily sol. in H_2O . (Carlgren, A. F. 47. 310.)

— **nitrite, $Pt[(NH_2)_2NO_2]_2 + 2H_2O$.**

Efflorescent. Very sol. in hot or cold. Insol. in 90% alcohol. (Lang.)

— **platinous nitrite, $Pt[(NH_2)_2NO]Pt(NO_2)_2$.**

Scarcely sol. in cold, somewhat more in hot H_2O . Not attacked by cold. More sol. in $NH_4OH + Aq$ than in H_2O . (Lang.)

phosphate, $\text{Pt}(\text{N}_2\text{H}_4)_2\text{HPO}_4$

difficultly sol. in cold, and very
H₂O. (Cleve.)

ium phosphate,

$(\text{NH}_4)_2\text{PO}_4(\text{NH}_4)_2$, $4\text{NH}_4\text{H}_2\text{PO}_4 +$

sol. in H₂O with decomp. into—
 $(\text{NH}_4)_2\text{PO}_4$, $2\text{NH}_4\text{H}_2\text{PO}_4 + 9\text{H}_2\text{O}$.

sol. in H₂O than the preceding.
(Cleve.)

e, $\text{Pt}(\text{NH}_3)_4\text{SO}_4$.

pts. H₂O at 16.5°; more easily
(Reiset.)

60 pts. boiling H₂O; less in cold
alcohol. (Cleve.)

e, **acid**, $\text{Pt}[(\text{NH}_3)_2\text{SO}_4\text{H}]_2 + \text{H}_2\text{O}$.

by H₂O or alcohol into neutral

SO_4 , $\text{H}_2\text{SO}_4 + \text{H}_2\text{O}$. Sol. in H₂O.

e, $\text{Pt}(\text{NH}_3)_4\text{SO}_3$.

sol. in cold H₂O. (Birnbaum, A.

$\text{SO}_3\text{H}]_2 + 2\text{H}_2\text{O}$. Ppt. Sol. in
Cleve.)

is sulphite,

$(\text{NH}_3)_4\text{SO}_3$, $\text{PtSO}_3 + 2\text{H}_2\text{O}$.

sol. in cold H₂O; sol. in 190 pts.
Easily sol. in warm HCl + Aq

(Peyrone.)

(Carlgren, Sv. V. A. F. 47. 308.)

SO_3 , PtSO_3 , H_2SO_3 . Insol. in
alcohol. Scarcely sol. in hot H₂O.

yanide, $\text{Pt}(\text{NH}_3)_4(\text{CNS})_2 + \text{H}_2\text{O}$.

in H₂O. Solution is decomp. on
boiling, Sv. V. A. H. 10, 9. 7.)

is sulphocyanide,

$(\text{NH}_3)_4(\text{CNS})_2$, $\text{Pt}(\text{CNS})_2$.

sol. in H₂O and alcohol; sol. in dil. HCl +
boiling, Chem. Soc. 13. 122.)

amine chloride, $\text{Pt}(\text{NH}_3)_2\text{Cl}$
 NH_3Cl .

in H₂O. (Cleve.)

amine chloroplatinite

$(\text{NH}_3)_2\text{Cl}$, PtCl_2 .

sol. in cold, but more easily in
boiling. (Cleve.)

$\text{Pt}(\text{NH}_3)_2\text{NO}_2 + \text{H}_2\text{O}$.

in H₂O. (Cleve.)

Platomonodiamine sulphate, $\text{Pt}(\text{NH}_3)_2\text{SO}_4$.

Easily sol. in cold, but much more in hot
H₂O.

Platosemidiamine bromide, $\text{Pt}(\text{NH}_3)_2\text{Br}$
 Br .

Sol. in H₂O. Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
(Cleve.)

— **chloride**, $\text{Pt}(\text{NH}_3)_2\text{Cl}$
 Cl .

(Peyrone's chloride.) Sol. in 387 pts. H₂O
at 0°, and 26 pts. at 100° (Cleve); in 33 pts. at
100°. (Peyrone.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; very sl. sol. in HCl or
 $\text{H}_2\text{SO}_4 + \text{Aq}$; more easily in $\text{HNO}_3 + \text{Aq}$; sol.
in alkali carbonates + Aq. (Peyrone, A. ch.
(3) 12. 193.)

Platosemidiamine chlorosulphurous acid,

$\text{Pt}(\text{NH}_3)_2\text{SO}_3\text{H}$
 Cl .

Easily sol. in H₂O. (Cleve.)

Ammonium platosemidiamine chlorosulphite
platosemidiamine sulphite,

$\text{Pt}(\text{NH}_3)_2\text{SO}_3\text{NH}_4$, $\text{Pt}(\text{NH}_3)_2\text{SO}_3\text{NH}_4 +$
 SO_3NH_4
 $2\text{H}_2\text{O}$.

Easily sol. in H₂O. Insol. in alcohol.
(Cleve.)

Platosemidiamine cyanide,

$\text{Pt}(\text{CN})(\text{NH}_3)_2\text{CN}$.
Easily sol. in H₂O. (Cleve.)

— **platinous cyanide**, $\text{Pt}(\text{CN})(\text{NH}_3)_2\text{CN}$,
 $\text{Pt}(\text{CN})_2(?)$.

Ppt.

— **hydroxide**, $\text{Pt}(\text{NH}_3)_2\text{OH}$
 OH .

Not known.

— **iodide**, $\text{Pt}(\text{NH}_3)_2\text{I}$
 I .

Sl. sol. in boiling H₂O. (Cleve.)

— **nitrate** $\text{Pt}(\text{NH}_3)_2\text{NO}_3$
 NO_3 .

Moderately sol. in H₂O. (Cleve.)

— **nitrite**, $\text{Pt}(\text{NH}_3)_2\text{NO}_2$
 NO_2 .

Very sl. sol. in cold, more easily in hot H₂O.

— **oxalate**, $\text{Pt}(\text{NH}_3)_2\text{C}_2\text{O}_4$.
(Cleve.)

+ 2H₂O. (Cleve.)

— **sulphate**, $\text{Pt}(\text{NH}_3)_2\text{SO}_4$.

Very sl. sol. even in hot H₂O. (Cleve.)

Platosemidiamine sulphocyanide,
 $\text{Pt}(\text{SCN})(\text{NH}_3)_2\text{SCN}.$

Easily sol. in warm H_2O , but solution soon decomposes.

Platosemidiamine sulphurous acid.

Ammonium platosemidiamine sulphite,
 $\text{Pt} \begin{smallmatrix} (\text{NH}_3)_2\text{SO}_3(\text{NH}_4) \\ \text{SO}_3(\text{NH}_4) \end{smallmatrix}, (\text{NH}_4)_2\text{SO}_3.$

Very sol. in H_2O . (Cleve.)

Barium ———, $\text{Pt}(\text{SO}_3)[(\text{NH}_3)_2\text{SO}_3]\text{Ba}$, BaSO_3 .
 Ppt. (Cleve.)

Silver ———. $\text{Pt}(\text{SO}_3\text{Ag})[(\text{NH}_3)_2\text{SO}_3\text{Ag}]$,
 Ag_2SO_3 .
 Ppt. (Cleve.)

Diplatodiamine chloride, $\text{Pt}_2(\text{NH}_3)_4\text{Cl}_2$.
 Insol. in H_2O .

— hydroxide, $\text{Pt}_2(\text{NH}_3)_4(\text{OH})_2 + \text{H}_2\text{O}$.
 Insol. in H_2O .

— nitrate, $\text{Pt}_2(\text{NH}_3)_4(\text{NO}_3)_2$.
 Insol. in H_2O . (Cleve.)

— sulphate, $\text{Pt}_2(\text{NH}_3)_4\text{SO}_4$.
 Insol. in H_2O . (Cleve.)

Platobromonitrous acid.

Potassium platobromonitrite, $\text{K}_2\text{Pt}(\text{NO}_2)_2\text{Br} + 2\text{H}_2\text{O}$.

Sol. in about 3 pts. cold, and 2 pts. boiling H_2O . (Vèzes, A. ch. (6) 29. 194.)

$\text{K}_2\text{Pt}(\text{NO}_2)_2\text{Br}_2 + \text{H}_2\text{O}$. Sol. in 1 pt. cold, and still less hot H_2O . Insol. in alcohol. (Vèzes.)

Platochloronitrous acid.

Potassium chloronitrite, $\text{K}_2\text{Pt}(\text{NO}_2)_2\text{Cl} + 2\text{H}_2\text{O}$.

Sol. in about 3 pts. cold, and 2 pts. boiling H_2O . (Vèzes, A. ch. (6) 29. 178.)

$\text{K}_2\text{Pt}(\text{NO}_2)_2\text{Cl}_2$. Sol. in about 3 pts. cold, and 2 pts. boiling H_2O . (Vèzes.)

Platochlorosulphurous acid.

See Chloroplatosulphurous acid.

Platiodonitrous acid, $\text{H}_2\text{Pt}(\text{NO}_2)_2\text{I}_2$.

Known only in solution. (Nilson, J. pr. (2) 21. 172.)

Aluminum platiodonitrite, $\text{Al}_2[\text{Pt}(\text{NO}_2)_2\text{I}_2] + 27\text{H}_2\text{O}$.

Easily sol. in H_2O . (Nilson.)

Ammonium ———, $(\text{NH}_4)_2\text{Pt}(\text{NO}_2)_2\text{I}_2 + 2\text{H}_2\text{O}$.
 Sol. in H_2O ; decomp. on heating.

Barium platiodonitrite, $\text{BaPt}(\text{NO}_2)_2\text{I}_2 + 4\text{H}_2\text{O}$.

Very sol. in H_2O .

Cadmium ———, $\text{CdPt}(\text{NO}_2)_2\text{I}_2 + 2\text{H}_2\text{O}$.
 Easily sol. in H_2O .

Cæsium ———, $\text{Cs}_2\text{Pt}(\text{NO}_2)_2\text{I}_2 + 2\text{H}_2\text{O}$.
 Easily sol. in H_2O .

Calcium ———, $\text{CaPt}(\text{NO}_2)_2\text{I}_2 + 6\text{H}_2\text{O}$.
 Very easily sol. in H_2O .

Cerium ———, $\text{Ce}_2[\text{Pt}(\text{NO}_2)_2\text{I}_2]_3 + 18\text{H}_2\text{O}$.
 Easily sol. in H_2O .

Cobalt ———, $\text{CoPt}(\text{NO}_2)_2\text{I}_2 + 8\text{H}_2\text{O}$.
 Sol. in H_2O .

Didymium ———, $\text{Di}_2[\text{Pt}(\text{NO}_2)_2\text{I}_2]_3 + 24\text{H}_2\text{O}$.
 Sol. in H_2O .

Erbium ———, $\text{Er}_2[\text{Pt}(\text{NO}_2)_2\text{I}_2]_3 + 18\text{H}_2\text{O}$.
 Sol. in H_2O .

Ferrous ———, $\text{FePt}(\text{NO}_2)_2\text{I}_2 + 8\text{H}_2\text{O}$.
 Sol. in H_2O .

Ferric ———, $\text{Fe}_2[\text{Pt}(\text{NO}_2)_2\text{I}_2]_3 + 6\text{H}_2\text{O}$.
 Sol. in H_2O .

Lanthanum ———, $\text{La}_2[\text{Pt}(\text{NO}_2)_2\text{I}_2]_3 + 18\text{H}_2\text{O}$.
 Sol. in H_2O .

Lead ———, basic, $\text{PbPt}(\text{NO}_2)_2\text{I}_2$, $\text{Pb}(\text{OH})_2$.
 Insol. in H_2O .

Lithium ———, $\text{Li}_2\text{Pt}(\text{NO}_2)_2\text{I}_2 + 6\text{H}_2\text{O}$.
 Very sol. in H_2O .

Magnesium ———, $\text{MgPt}(\text{NO}_2)_2\text{I}_2 + 8\text{H}_2\text{O}$.
 Sol. in H_2O .

Manganese ———, $\text{MnPt}(\text{NO}_2)_2\text{I}_2 + 8\text{H}_2\text{O}$.
 Sol. in H_2O .

Mercurous ———, basic, $2\text{Hg}_2\text{Pt}(\text{NO}_2)_2\text{I}_2 + 9\text{H}_2\text{O}$.
 Insol. in H_2O .

Nickel ———, $\text{NiPt}(\text{NO}_2)_2\text{I}_2 + 8\text{H}_2\text{O}$.
 Sol. in H_2O .

Potassium ———, $\text{K}_2\text{Pt}(\text{NO}_2)_2\text{I}_2 + 2\text{H}_2\text{O}$.
 Sol. in H_2O in all proportions.
 Very sol. in alcohol.

Rubidium ———, $\text{Rb}_2\text{Pt}(\text{NO}_2)_2\text{I}_2 + 2\text{H}_2\text{O}$.
 Sol. in H_2O .

odonitrite, $\text{Ag}_2\text{Pt}(\text{NO}_2)_4 \cdot \text{I}_2 \cdot \text{H}_2\text{O}$.

—, $\text{Na}_2\text{Pt}(\text{NO}_2)_4 \cdot \text{I}_2 + 4\text{H}_2\text{O}$ in H_2O .

—, $\text{SrPt}(\text{NO}_2)_4 \cdot \text{I}_2 + 8\text{H}_2\text{O}$ in H_2O .

—, $\text{Th}_2\text{Pt}(\text{NO}_2)_4 \cdot \text{I}_2 \cdot \text{H}_2\text{O}$.

—, $\text{Y}_2[\text{Pt}(\text{NO}_2)_4 \cdot \text{I}_2]_3 + 27\text{H}_2\text{O}$ in H_2O .

$\text{ZnPt}(\text{NO}_2)_4 \cdot \text{I}_2 + 8\text{H}_2\text{O}$ in H_2O .

nitrosylic acid, $\text{H}_4\text{Pt}_3\text{O}(\text{NO}_2)_8$. pr. (2) 16. 241.)

riplatooctonitrosylate.

Platonitrite, potassium.

acid, $\text{H}_2\text{Pt}(\text{NO}_2)_4$.

O or alcohol. (Lang. J. pr. 83.

“Platotetranitrosylic acid” by

platonitrite, $\text{Al}_2[\text{Pt}(\text{NO}_2)_4]_3 +$

O. $\text{Pt}(\text{NO}_2)_4 \cdot \text{I}_2 + 10\text{H}_2\text{O}$. Sl. sol. in hot H_2O and alcohol. (Nilson. 727.)

platonitrite, $(\text{NH}_4)_2\text{Pt}(\text{NO}_2)_4 +$

very sol. in cold H_2O . (Nilson, B.

nitrite, $\text{BaPt}(\text{NO}_2)_4 + 3\text{H}_2\text{O}$. cold, very sol. in hot H_2O .

nitrite, $\text{CdPt}(\text{NO}_2)_4 + 3\text{H}_2\text{O}$ in H_2O . (Nilson.)

nitrite, $\text{Cs}_2\text{Pt}(\text{NO}_2)_4$. K salt.

nitrite, $\text{CaPt}(\text{NO}_2)_4 + 5\text{H}_2\text{O}$ in H_2O . (Nilson.)

nitrite, $\text{Ce}_2[\text{Pt}(\text{NO}_2)_4]_3 + 18\text{H}_2\text{O}$ in H_2O . (Nilson.)

diplatonitrite, $[\text{Pt}(\text{NO}_2)_4]_2 \cdot \text{O}_2 + 24\text{H}_2\text{O}$ in H_2O . (Nilson.)

Cobalt platonitrite, $\text{CoPt}(\text{NO}_2)_4 + 8\text{H}_2\text{O}$. Easily sol. in H_2O . (Nilson.)

Copper platonitrite, $\text{CuPt}(\text{NO}_2)_4 + 3\text{H}_2\text{O}$. Sol. in H_2O . (Nilson.) $3\text{CuPt}(\text{NO}_2)_4, \text{CuO} + 18\text{H}_2\text{O}$. Decomp. by H_2O . (Nilson.)

Didymium platonitrite, $\text{Di}_2[\text{Pt}(\text{NO}_2)_4]_3 + 18\text{H}_2\text{O}$. Deliquescent; sol. in H_2O .

Erbium platonitrite, $\text{Er}_2[\text{Pt}(\text{NO}_2)_4]_3 + 9$ and $21\text{H}_2\text{O}$. Deliquescent; sol. in H_2O .

Glucinum diplatonitrite, $\text{Gl}^2\text{Pt}(\text{NO}_2)_4 \cdot \text{O} + 9\text{H}_2\text{O}$. Sl. sol. in cold H_2O .

Indium diplatonitrite, $\text{In}(\text{OH})_3[\text{Pt}(\text{NO}_2)_4]_2 \cdot \text{O}_2 + 10\text{H}_2\text{O}$. Sl. sol. in H_2O .

Ferric diplatonitrite, $\text{Fe}_2[\text{Pt}(\text{NO}_2)_4]_3 \cdot \text{O}_2 + 30\text{H}_2\text{O}$. Sl. sol. in cold, easily in hot H_2O .

Lanthanum platonitrite, $\text{La}_2[\text{Pt}(\text{NO}_2)_4]_3 + 18\text{H}_2\text{O}$. Deliquescent; sol. in H_2O .

Lead platonitrite, $\text{PbPt}(\text{NO}_2)_4 + 3\text{H}_2\text{O}$. Sl. sol. in H_2O . (Nilson.)

Lithium platonitrite, $\text{Li}_2\text{Pt}(\text{NO}_2)_4 + 3\text{H}_2\text{O}$. Sl. deliquescent; easily sol. in H_2O .

Magnesium platonitrite, $\text{MgPt}(\text{NO}_2)_4 + 5\text{H}_2\text{O}$. Easily sol. in H_2O .

Manganese platonitrite, $\text{MnPt}(\text{NO}_2)_4 + 8\text{H}_2\text{O}$. Sol. in H_2O .

Mercurous platonitrite, $\text{Hg}_2\text{Pt}(\text{NO}_2)_4, \text{Hg}_2\text{O}$. Nearly insol. in H_2O . (Lang, J. pr. 83. 415.) $+ \text{H}_2\text{O}$. Nearly insol. in H_2O . (Nilson.)

Nickel platonitrite, $\text{NiPt}(\text{NO}_2)_4 + 8\text{H}_2\text{O}$. Easily sol. in H_2O . (Nilson.)

Potassium platonitrite, $\text{K}_2\text{Pt}(\text{NO}_2)_4$. Sol. in 27 pts. H_2O at 15° ; more easily sol. in warm H_2O . (Lang, J. pr. 83. 415.) $+ 2\text{H}_2\text{O}$. Efflorescent. (Lang.) $\text{K}_2\text{H}_4\text{Pt}_3\text{O}(\text{NO}_2)_8 + 3\text{H}_2\text{O}$. Very sl. sol. in cold (0.01 mol. in 1 l. at 16°), but very easily in hot H_2O . (Vèzes, A. ch. (6) 29. 162.) $\text{K}_4\text{Pt}_3\text{O}(\text{NO}_2)_8 + 2\text{H}_2\text{O}$. Sl. sol. in warm H_2O . (Nilson.)

Potassium platonitrite bromide.

See Platibromonitrite and platobromonitrite, potassium.

Potassium platonitrite chloride.

See Plati- and platochloronitrite, potassium.

Potassium plat nitrite hydrogen chloride,

Sol. in H_2O . (Miolati, Att. Linc. Rend. 1896, (5) 5, II. 358.)

Potassium platonitrite iodide.

See Plati- and platiodonitrite, potassium.

Potassium platonitrite nitrogen dioxide,

Violently decomp. by H_2O . (Miolati, Atti Linc. Rend. 1896, (5) 5, II, 356.)

Rubidium platonitrite, $Rb_2(Pt)(NO_2)_4$, and $+2H_2O$.

Very slowly sol. in cold, more easily in warm H_2O . (Nilson.)

Silver platonitrite, $Ag_2Pt(NO_2)_4$.

Very sl. sol. in cold, easily in hot H_2O .

Silver diplatonitrite, $Ag_2Pt_2(NO_2)_4O$.

Insol. in H_2O . (Nilson.)

Sodium platonitrite, $Na_2Pt(NO_2)_4$.

Easily sol. in H_2O .

Strontium platonitrite, $SrPt(NO_2)_4 + 3H_2O$.

Somewhat sl. sol. in cold H_2O , but easily sol. in warm H_2O .

Thallium platonitrite, $Tl_2Pt(NO_2)_4$.

Very sl. sol. in H_2O . (Nilson.)

Yttrium platonitrite, $Y_2[Pt(NO_2)_4]_3 + 9$, or $21H_2O$.

Sol. in H_2O .

Zinc platonitrite, $ZnPt(NO_2)_4 + 8H_2O$.

Sol. in H_2O .

Platodioxamine chloride,

Easily sol. in H_2O . (Alexander, A. 246. 239.)

— chloroplatinite, $Pt(NH_2O.NH_2OCl)_2$, $PtCl_2$.

Sol. in warm $HCl + Aq$. Insol. in cold H_2O or alcohol; very sl. sol. in hot H_2O . (Alexander.)

— hydroxide, $Pt(NH_2O.NH_2O)_2(OH)_2$.

Insol. in H_2O or alcohol. Easily sol. in HCl or $HNO_3 + Aq$. Difficultly sol. in hot dil. $H_2SO_4 + Aq$. (Alexander.)

Platodioxamine oxalate,

Insol. in cold H_2O , alcohol, or organic (Alexander.)

— phosphate, $Pt_2(NH_2O.NH_2O)_4 \cdot 3H_2O$.

Ppt. (Alexander.)

— sulphate, $Pt(NH_2O.NH_2O)SO_4$.

Sl. sol. in H_2O . (Alexander.)

Platosamine bromide, $Pt(NH_2Br)_2$.

Sl. sol. even in hot H_2O . (Cleve.)

— chloride, $Pt(NH_2Cl)_2$.

"Reiset's second chloride." Sol. in H_2O at 100° . (Peyrone, A. 61. 180.)

Sol. in 130 pts. H_2O at 100° , and 44 at 0° . (Cleve.)

Easily sol. in $NH_4OH + Aq$, HNO_3 , regia, with decomp. Sol. in $KCN + Aq$ evolution of NH_3 . (Cleve.)

— ammonium chloride, $Pt(NH_2Cl)_2 \cdot 2NH_4Cl$.

Sl. sol. in cold, easily in hot H_2O ; in alcohol; sol. in NH_4OH or $(NH_4)_2CO_3$ (Grimm, A. 99. 75.)

Platosamine chlorosulphurous acid

Easily sol. in H_2O without decomp. (Cleve.)

Ammonium platosamine chlorosulph

Sol. in H_2O . (Peyrone, A. 61. 180.)

Platosamine cyanide, $Pt(NH_2CN)_2$.

Quite easily sol. in H_2O or NH_4OH (Buckton.)

— hydroxide, $Pt(NH_2OH)_2$.

"Reiset's second base." Very sol. (Odling, B. 3. 685.)

— iodide, $Pt(NH_2I)_2$.

Very sl. sol. in H_2O . Sol. in cold $NH_4OH + Aq$ to form platodiamine iodide. (Cleve.)

— nitrate, $Pt(NH_2NO_3)_2$.

Moderately sol. in hot H_2O . $NH_4OH + Aq$ with combination. (Reiset, ch. (3) 11. 26.)

— nitrite, $Pt(NH_2NO_2)_2$.

Very sl. sol. in cold, easily in hot H_2O . Insol. in alcohol. (Lang.)

platinous nitrite, $\text{Pt}(\text{NH}_2\text{NO}_2)_2$,
 $\text{Pt}(\text{NO})_2$.

d sl. sol. in cold, more easily sol. in

y sl. sol. even in conc. acids; more
)H + Aq than in H_2O . (Lang.)

$\text{Pt}(\text{NH}_3)_2\text{O}$.

H_2O or NH_4OH + Aq. (Reiset.)

$\text{Pt}(\text{NH}_3)_2\text{H}_2(\text{C}_2\text{O}_4)_2 + 2\text{H}_2\text{O}$.
 (Cleve.)

platous sulphate, $\text{Pt}(\text{NH}_3)_2\text{SO}_4 + \text{H}_2\text{O}$.

cold, more easily in hot H_2O .

platous sulphate, $\text{Pt}(\text{NH}_3)_2\text{SO}_4 + \text{H}_2\text{O}$.

in H_2O . (Cleve.)

platous cyanide, $\text{Pt}(\text{NH}_3)_2\text{SCN}$.

H_2O ; can be cryst. from alcohol;
 l by HCl or H_2SO_4 + Aq. (Buck-

in hot H_2O . (Cleve.)

platous sulphocyanide,

$\text{Pt}(\text{NH}_3)_2\text{Ag}_4(\text{SCN})_6$.

platous sulphurous acid,

SO_2H_2 .

y in its salts.

platous amine sulphite.

platous amine sulphite,

SO_2NH_4 .

O. Insol. in alcohol.

platous amine sulphite,

$\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Ba} + 3\text{H}_2\text{O}$.

(Cleve.)

—, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{CO} +$

sol. in H_2O . Sol. in HCl + Aq.

— —, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Cu} +$

l. in H_2O ; sol. in HCl + Aq.

—, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Pb} + \text{H}_2\text{O}$.

— —, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Mn}$

ol. in H_2O .

—, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Ni} + 7\text{H}_2\text{O}$.

H_2O .

Sodium platousamine sulphite,

$\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Na} + 5\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O . 100 ccm. sat. solution at 20°
 contains 5.52 g. cryst. salt. (Haberland and
 Hanekop, A. 245. 235.)

Silver — —, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Ag} + \text{H}_2\text{O}$.

Ppt.

Uranyl — —, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{UO}_2 + \text{H}_2\text{O}$.

Ppt.

Zinc — —, $\text{Pt}(\text{NH}_3)_2(\text{SO}_3)_2\text{Zn} + 6\text{H}_2\text{O}$.

Ppt. Very sl. sol. in H_2O . (Cleve.)

Platousosemiamine potassium chloride,

$\text{Pt} \begin{smallmatrix} \text{NH}_3\text{Cl} \\ \text{Cl} \end{smallmatrix}$, $\text{KCl} + \text{H}_2\text{O}$.

Very sol. in H_2O ; insol. in alcohol. (Cossa,
 B. 23. 2507.)

Platousoxamine chloride, $\text{Pt} \begin{smallmatrix} \text{NH}_3\text{OCl} \\ \text{NH}_3\text{OCl} \end{smallmatrix}$.

Sol. in H_2O . Much less sol. in H_2O than
 platodioxamine chloride. (Alexander, A. 246. 239.)

Platousoxamine amine chloride,

$\text{Pt} \begin{smallmatrix} \text{NH}_3\text{O.NH}_3\text{Cl} \\ \text{NH}_3\text{O.NH}_3\text{OCl} \end{smallmatrix}$.

Easily sol. in H_2O . Insol. in alcohol and
 conc. HCl + Aq. (Alexander, A. 246. 239.)

— chloroplatinite, $\text{Pt} \begin{smallmatrix} \text{NH}_3\text{O.NH}_3\text{Cl} \\ \text{NH}_3\text{O.NH}_3\text{OCl} \end{smallmatrix} \text{PtCl}_2$

Ppt.

Platousulphurous acid.

Ammonium platousulphite, $(\text{NH}_4)_2\text{Pt}(\text{SO}_3)_2 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Birnbbaum, A. 139. 170.)
 $(\text{NH}_4)_2\text{Pt}(\text{SO}_3)_2 + \text{H}_2\text{O}$. Sol. in H_2O . (Lie-
 big, Pogg. 17. 108.)

Ammonium platousulphite chloride,

$(\text{NH}_4)_2\text{Pt}(\text{SO}_3)_2, 2\text{NH}_4\text{Cl}$.

Sol. in H_2O . (Birnbbaum.)
 $\text{PtClSO}_3\text{H}, 2\text{NH}_4\text{Cl}$. Deliquescent; sol. in
 H_2O . (Birnbbaum, A. 152. 143.)

See also Chloroplatousulphite, ammonium.

Potassium platousulphite, $\text{K}_2\text{Pt}(\text{SO}_3)_2 + 4\text{H}_2\text{O}$.

Sl. sol. in cold, easily in hot H_2O . Much
 more sol. than the Na salt. (Birnbbaum, A. 139. 168.)

+ $3\text{H}_2\text{O}$. (Lang, J. pr. 83. 415.)

$6\text{K}_2\text{O}, 2\text{PtO}, 10\text{SO}_2$. Sl. sol. in H_2O .

(Claus, J. B. 1847-48. 453.)

Does not exist. (Lang.)

$\text{K}_2\text{Pt}(\text{SO}_3)_2$. Sol. in H_2O .

Silver platosulphite, $\text{Ag}_2\text{Pt}(\text{SO}_3)_4$.

Ppt. Very sol. in cold $\text{NH}_4\text{OH} + \text{Aq.}$ (Lang. J, pr. 83. 415.)

Sodium platosulphite, $\text{Na}_4\text{Pt}(\text{SO}_3)_4$.

Very sl. sol. in cold, somewhat more easily in hot H_2O . Not decomp. by boiling KOH or $\text{NaOH} + \text{Aq.}$ Gradually sol. in $(\text{NH}_4)_2\text{S}$ or $\text{K}_2\text{S} + \text{Aq.}$ Insol. in $\text{NaCl} + \text{Aq}$ or alcohol. (Litton and Schnedermann, A. 42. 316.)

+ $1\frac{1}{2}\text{H}_2\text{O}$.

+ $7\text{H}_2\text{O}$.

$\text{Na}_2\text{Pt}(\text{SO}_3\text{H})_4$. Moderately sol. in H_2O . (Litton and Schnedermann.)

Platothiosulphuric acid.**Sodium platothiosulphate, $\text{Na}_4\text{Pt}(\text{S}_2\text{O}_3)_4 + 10\text{H}_2\text{O}$.**

Very sol. in H_2O (Schottländer, A. 140. 200.)

$\text{PtS}_2\text{O}_3, 4\text{Na}_2\text{S}_2\text{O}_3 + 10\text{H}_2\text{O}$.

$\text{PtS}_2\text{O}_3, 6\text{Na}_2\text{S}_2\text{O}_3 + 19\text{H}_2\text{O}$.

$2\text{Pt}_2\text{S}_2\text{O}_3, 7\text{Na}_2\text{S}_2\text{O}_3 + 18\text{H}_2\text{O}$. (Jochum, C. C. 1885. 642.)

Plumbic acid.**Barium plumbate, Ba_2PbO_4 .**

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$ with evolution of Cl . Sol. in acids in presence of a reducing substance. (Kassner, Arch. Pharm. 228. 109.)

Calcium plumbate.

Insol. in H_2O . $\text{HNO}_3 + \text{Aq}$ dissolves out CaO . (Crum, A. 55. 218.)

Ca_2PbO_4 . Properties as Ba_2PbO_4 . (Kassner, Arch. Pharm. 228. 109.)

+ $4\text{H}_2\text{O}$. Easily decomp. by HNO_3 . (Kassner, Arch. Pharm. 1894, 232. 378.)

Calcium hydrogen plumbate, $\text{H}_2\text{CaPb}_2\text{O}_6$.

Fairly stable; slowly sol. in HNO_3 in the cold. (Kassner.)

Calcium lead orthoplumbate, CaPbPbO_4 .

Insol. in H_2O . Sol. in HCl . HNO_3 , acetic and other acids cause a separation of PbO_2 . (Kassner, Arch. Pharm. 1903, 241. 147.)

Copper metaplumbate, CuPbO_3 .

Decomp. by acids. Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Acetic acid dissolves Cu . (Hoehnel, Arch. Pharm. 1896, 234. 399.)

Lead metaplumbate, PbPbO_3 .

Identical with lead sesquioxide. (Hoehnel, Arch. Pharm. 1896, 234. 399.)

Manganese metaplumbate.

Decomp. by acids. (Hoehnel, Arch. Pharm. 1896, 234. 399.)

Potassium plumbate, $\text{K}_2\text{PbO}_3 + 3\text{H}_2\text{O}$.

Very deliquescent. Decomp. by pure H into PbO_2 and KOH . Sol. in $\text{KOH} + \text{Aq}$ without decomp. (Freymy, J. Pharm. (3) 2. 32.)

Silver metaplumbate, Ag_2PbO_3 .

Ppt. (Grützner, Arch. Pharm. 1895, 1. 518.)

Sodium plumbate.

Sol. in H_2O with decomposition. Sl. sol. in alkalies + Aq. (Freymy, A. ch. (3) 12. 490.)

Sodium metaplumbate, $\text{Na}_2\text{PbO}_3 + 4\text{H}_2\text{O}$.

Decomp. by H_2O ; insol. in alcohol. (Hoehnel, Arch. Pharm. 1894, 232. 224.)

Strontium plumbate, Sr_2PbO_4 .

Properties as Ba_2PbO_4 . (Kassner, Arch. Pharm. 228. 109.)

Zinc metaplumbate, $\text{ZnPbO}_3 + 2\text{H}_2\text{O}$.

Decomp. by dil. acids; insol. in H_2O . (Hoehnel, Arch. Pharm. 1896, 234. 398.)

Plumbous acid.**Calcium plumbite.**

Sl. sol. in H_2O . (Karsten, Scher. J. 8.)

Potassium plumbite, $\text{PbO}, \pm \text{K}_2\text{O}$.

Known only in solution.

Silver plumbite, Ag_2PbO_3 .

Insol. in H_2O , $\text{NH}_4\text{OH} + \text{Aq}$ and H ; sol. in HNO_3 and acetic acid; and in H_2SO_4 , HI , and HF . (Bullheimer, B. 31. 1288.)

+ $2\text{H}_2\text{O}$. Insol. in H_2O . Decomp. (Kratwig, B. 15. 264.)

Sodium plumbite.

Known only in solution.

Potassium, K_2 .

Violently decomposes H_2O or H . Insol. in hydrocarbons. Sol. with action in acids.

Solubility in fused KOH at t° .

t°	G. K sol. in 100 g. fused
480	7.8—8.9
600	3 — 4
650	2 — 2.7
700	0.5—1.3

(Hevesy, Z. Elektrochem. 1909, 15.)

Sol. in liquid NH_3 . (Seely, C. N. 21. 829.) (Franklin, Am. Ch. J. 1896, 20. 829.)
1 gram atom of K dissolves in 4.7

at 0°; in 4.79 mol. at -50°; in -100°. (Ruff, B. 1906, 39. 839.)
liquid CO₂. (Büchner, Z. phys. 674.)

in ethylene diamine. Insol. in and in secondary and tertiary
raus, J. Am. Chem. Soc. 1907,

acetylide, K₂C₂.

C. R. 1898, 127. 917.)

acetylide acetylene, K₂C₂, C₂H₂.

C. R. 1898, 127. 915.)

amalgams.

K, Hg K, Hg₁₀K, Hg₁₂K and
ntz, C. R. 1900, 131. 183.)

stable up to 0°. Can be cryst.
out decomp. below 0°.

stable from 0° to 71° or 73°. Can
m Hg without decomp. at any
n these limits.

stable from 71° or 73-75°. Can
m Hg without decomp. at any
n these limits. (Kerp, Z. anorg.
)

arsenide, KH₂N.

by water or alcohol. Insol. in
s.

ammononickelate, Ni₂N₂K₂, 6NH₃.

by H₂O. Sl. sol. in liquid NH₃.
phys. Chem. 1915, 19. 559.)

ammonioargentate, AgNHK, NH₃,
I₂, KNH₂.

np. in the air. Decomp. by H₂O
NH₃ solutions of acids. (Frank-
hem. Soc. 1915, 37. 855.)

ammonobarate, BaNK, 2NH₃.

l by H₂O. Insol. in liquid NH₃.
d dissolved in a solution of
liquid NH₃. (Franklin, J. Am.
915, 37. 2297.)

ammonocadmiate, Cd(NHK)₂,

Franklin, Am. Ch. J. 1912, 47. 310.)

, 2NH₃. Decomp. by H₂O.
d NH₃. (Bohart, J. phys. Chem.
)

ammonocalciate, CaNK, 2NH₃.

l by H₂O. Readily sol. in a
H₄NO₃ in liquid NH₃. (Frank-
hem. Soc. 1915, 37. 2300.)

ammonocuprite, CuNK₂, 3NH₃.

liquid NH₃.

NH₃.

CuNK₂, NH₃. (Franklin, J. Am. Chem.
Soc. 1912, 34. 1503.)

CuNK₂, 2½NH₃. Ppt. (Franklin, Am.
Ch. J. 1912, 47. 311.)

Potassium ammonomagnesate, Mg(NHK)₂,
2NH₃.

Sl. sol. in liquid NH₃. Rapidly hydrolyzed
by H₂O. (Franklin, J. Am. Chem. Soc. 1913,
35. 1463.)

Potassium ammonoplumbite, PbNK, 2½NH₃.

Completely hydrolyzed by action of water
vapor. Violently decomp. by H₂O or dil.
acids. Sol. in liquid NH₃. (Franklin, J.
phys. Chem. 1911, 15. 519.)

Potassium ammonostannate, Sn(NK)₂, 4NH₃.

Decomp. by H₂O. Readily sol. in HCl+
Aq. Sl. sol. in liquid NH₃. Readily sol. in a
solution of NH₄I in liquid NH₃. (Fitzgerald,
J. Am. Chem. Soc. 1907, 29. 1696.)

Potassium ammonostrontiate, SrNK, 2NH₃.

Hydrolyzed vigorously by H₂O. Sol. in
solutions of NH₄NO₃ in liquid NH₃. Insol.
in liquid NH₃. (Franklin, J. Am. Chem. Soc.
1915, 37. 2299.)

Potassium ammonothallate, TlNK, 4NH₃.

Sensitive to action of air or moisture. Vio-
lently decomp. by H₂O or dil. acids. Mod-
erately sol. in liquid NH₃ at 20°, more sol.
at higher temp. and much less sol. at lower
temp. Decomp. by liquid NH₃ solutions of
acids. (Franklin, J. phys. Chem. 1912, 16.
689.)

Potassium ammonotitanate, (N) TiNHK.

Vigorously hydrolyzed by H₂O. Insol. in
liquid NH₃ solutions of either potassium
amide or NH₄Br. (Franklin, J. Am. Chem.
Soc. 1912, 34. 1500.)

Potassium ammonozincate, Zn(NHK)₂,
2NH₃.

Decomp. by water. Sl. sol. in liquid NH₃.
(Fitzgerald, J. Am. Chem. Soc. 1907, 29. 663.)

Decomp. by H₂O. Sol. in dilute acids.
Sl. sol. in liquid NH₃. Sol. in solutions of
ammonium salts in liquid NH₃. (Franklin,
Z. anorg. 1907, 55. 195.)

Potassium arsenide, K₂As.

(Hugot, C. R. 1899, 129. 604.)

K₂As₄. (Hugot.)

Potassium arsenide, ammonia, AsK₂, NH₃.

Nearly insol. in liquid NH₃. (Hugot.)

K₂As₄, NH₃. (Hugot.)

Potassium azoimide, KN₃.
Stable in aq. solution.
46.5 pts. are sol. in 100 pts. H₂O at 10.5°.
48.9 " " " " 100 " H₂O " 15.5.
49.6 " " " " 100 " H₂O " 17.
0.1375 " " " " 100 " abs. alcohol at 16°.
Insol. in pure ether. (Curtius, J. pr. 1898, (2) 58. 280.)

Potassium bromide, KBr.

Solubility of KBr in 100 pts. H₂O at t°.

t°	Pts. KBr	t°	Pts. KBr
0	53.48	60	85.35
20	64.52	80	93.46
40	74.63	100	102.0

(Kremers, Pogg. 97. 151.)

Solubility of KBr in 100 pts. H₂O at t°.

t°	Pts. KBr	t°	Pts. KBr
-13.4	46.17	43.15	77.0
-6.2	49.57	45.45	77.73
0	53.32	50.5	80.33
+3.4	55.60	54.8	82.78
5.2	56.63	60.15	85.37
12.65	61.03	66.75	88.22
13.0	61.17	71.45	90.69
13.3	61.45	74.85	92.25
18.3	64.11	86.5	97.28
26.05	68.31	97.9	102.9
30.0	70.35	110.0	110.3
37.9	74.46

Solubility is represented by a straight line of the formula 54.43+0.5128t. (Coppet, A. ch. (5) 30. 416.)
100 pts. KBr+Aq sat. at 15-16° contain 39.06 pts. KBr. (v. Hauer, J. pr. 98. 137.)

Solubility of KBr in 100 pts. H₂O at high temp.

t°	Pts. KBr
140	120.9
181	145.6

(Tilden and Shenstone, Phil. Trans. 1884. 23.)

Sat. solution boils at 112°. (Kremers.)
Sat. KBr+Aq contains at:
-12° -10° +3° 32° 39°
31.0 31.5 35.7 41.6 47.7% KBr,
55° 77° 140° 173° 220°
45.5 48.7 54.1 58.5 61.6% KBr.
(Étard, A. ch. 1894, (7) 2. 539.)

If solubility S=pts. KBr in 100 pts. solution, S=34.5+0.2420t from 0° to 40°, S=

41.5+0.1378t from 30° to 120°. (C. R. 98. 1432.)

Solubility of KBr in 100 g. H₂O at

t°	G. KBr.	t°	G
10.5	62.1	-11	
10	60.7	-10.5	
3.5	55.5	-10	
0	52.6	-8.5	
-5	50.1	-8	
-8	47.5	-6.5	
-11.5	45.3	...	

(Meusser, Z. anorg. 1906, 44. 80
68.74 g. KBr are sol. in 100 g. H₂O (Amadori and Pampanini, Rend. Ac 1911, V, 20. 473.)

Sp. gr. of KBr+Aq at 19°.

% KBr	Sp. gr.	% KBr	
5	1.037	30	
10	1.075	35	
15	1.116	40	
20	1.159	45	
25	1.207	..	

(Gérlach, Z. anal. 8. 285.)
Sp. gr. of KBr+Aq at 15° contain
5 10 20 30 36
1.0357 1.074 1.1583 1.2553 1.3
(Kohlrausch, W. Ann. 1879. 1

Sp. gr of KBr+Aq at t°.

G. KBr dissolved in 100 g. H ₂ O	G. KBr in 100 g. of the solution	t°
4.166	4	14.5
11.111	10	15.7
25.000	20	16.5
42.867	30	16.0

(de Lannoy, Z. phys. Ch. 1895, 18
KBr+Aq containing 6.99% KBr gr. 20°/20°=1.0521. (Le Blanc and Z. phys. Ch. 1896, 19. 278.)

Sp. gr. of KBr+Aq at 20.5°.

Normality of KBr+Aq.	G. KBr in 100 g. of solution	Sp.
4.29	37.97	1.
3.01	28.83	1.
2.00	20.49	1.
1.00	11.03	1.

(Oppenheimer, Z. phys. Ch. 1898, 1

Solubility of KBr+NH₄Br at 25°.

% NH ₄ Br	Sp. gr
0.0	1.3756
0.64	1.3745
2.46	1.3733
5.13	1.3721
15.29	1.3711
26.22	1.3715
34.76	1.3753
38.14	1.3753
41.78	1.3766
43.25	1.3777
48.08	1.3766
57.73	1.3763

(Kryst. Min. 1897, 28. 357.)

Solubility in KNO₃+Aq.

1 litre of the solution contains		
4.5°	at 25.2°	
Mol. KBr	Mol. KNO ₃	Mol. KBr
4.332	0.0	4.761
4.156	0.131	4.72
4.093	0.527	4.61
3.939	0.721	4.54
...	1.090	4.475
...	1.170	4.44
...	1.504	4.375

(Touren, C. R. 1900, 130. 911.)
under KNO₃.

KBr+KCl+Aq sat. at 15-16° con-
tains 10.0 pts. of the two salts; 10.0 pts.
-Aq sat. at 15-16° contain 57.96
two salts; 100 pts. KBr+KCl+
t. at 15-16° contain 57.88 pts. of
alts. (v. Hauer, J. pr. 98. 137.)

Solubility of KBr+KCl in H₂O at 25°.

% KCl	Sp. gr.
0.00	1.3756
2.34	1.3700
4.66	1.3648
8.26	1.3544
13.66	1.3320
16.69	1.3119
21.39	1.2689
25.09	1.2455
29.17	1.1977
31.13	1.1756

(Kryst. Min. 1897, 28. 357.)

Solubility of KBr+KCl in H₂O at t°.

t°	Sat. solution contains		
	% KCl	% KBr	% total salt
-14	10.7	18.8	29.5
-13.7	29.4
-13.5	29.5
-7	10.7	19.8	30.5
+5.2	34.4
+6	11.3	22.6	33.9
10	11.0	23.7	34.7
21	10.8	...	35.3
26	11.2	25.5	36.7
30	39.4
32	11.9	26.6	38.5
39	39.8
47	11.0	30.8	41.8
52	11.0	31.2	42.2
55	11.9	29.9	41.8
71	12.0	31.7	43.7
73	11.8	32.9	44.7
102	12.8	35.8	48.6
152	13.2	40.6	53.8
160	12.5	42.3	54.8
168	55.0
225	14.7	45.0	59.7

(Étard, A. ch. 1894, (7) 3. 281.)

Solubility of KBr+KCl in H₂O at 25°.

G. per 100 g. H ₂ O.	
KBr	KCl
68.47	0.0
62.26	5.43
58.50	8.46
52.45	12.48
45.42	17.17
38.70	21.23
26.62	25.88
12.94	31.02
0.0	36.12

(Amadori and Pampanini, Att. Acc. Linc.
1911, 20, II. 475.)

Solubility in KCl+Aq at 25.2°.

1 litre of the solution contains	
Mol. KCl	Mol. KBr
0.0	4.761
0.67	4.22
0.81	4.15
1.35	3.70
1.48	3.54
1.61	3.42
1.70	3.34
2.46	2.50
3.775	0.525

(Touren, C. R. 1900, 130. 1252.)

See also under KCl.

By repeatedly heating KBr + Aq sat. at 15–16° with KI and cooling to 15°, nearly all the KBr can be separated (v. Hauer.)

100 pts. H₂O sat. with KBr at 16° dissolve 13.15 pts. KI, but on addition of more KI, KBr is pptd. (van Melckebeke, C. C. 1872, 586.)

Solubility in KI + Aq at t°.

t°	Sat. solution contains		
	% KBr	% KI	% total salt
-22	8.3	42.6	50.9
-19	9.5	42.8	52.3
-6	9.3	44.7	54.0
-1.5	55.3
+3	10.3	45.9	56.2
13.6	10.1	46.2	55.9
25	10.8	48.0	58.8
44.2	11.1	50.1	61.2
51	12.1	50.0	62.1
66	10.8	53.1	63.9
70	11.6	51.9	63.5
80	12.3	52.5	64.8
93	13.0	53.7	66.7
116	13.2	54.6	67.8
125	13.7	54.8	68.5
150	15.1	55.1	70.2
175	16.0	57.2	73.2
175	72.7
196	16.7	56.5	73.2
220	17.6	57.0	74.6

(Étard, A. ch. 1894, (7) 3, 279.)

Solubility of KBr + KI in H₂O at 25°.

G per 100 g. H ₂ O	
KBr	KI
53.21	35.92
42.32	66.63
34.14	95.36
30.08	119.59
29.62	119
22.15	127.10
21.88	127.31
18.54	130.61
0.0	149.26

(Amadori and Pampanini, Att. Acc. Lino. 1911, 20, II, 475.)

Solubility of KBr in KOH + Aq.

G per 1000 g. H ₂ O		G per 1000 g. H ₂ O	
KOH	KBr	KOH	KBr
36.4	558.4	277.6	248.1
113.5	433.6	434.7	137.1
177.2	358.1	579.6	64.8
231.1	281.2	806.9	33.4

(Ditte, C. R. 1897, 124, 30.)

Sol. in Br₂ at 15°. (Walden, Z. anorg. 25, 220.)

Moderately sol. in liquid NH₃. (Am. Ch. J. 1898, 20, 829.)

Sol. in liquid NH₃. 45 pts. are a pts. liquid NH₃ at -50°. (Joann. 1905, 140, 1244.)

Attacked by liquid NO₂ with lib. Br₂. (Frankland, Chem. Soc. 1, 1361.)

Insol. in liquid CO₂. (Böchner, Ch. 1906, 54, 674.)

Sol. in SO₂. (Walden, Z. anorg. 217.)

Sol. in SO₂Cl(OH). (Walden, 1902, 29, 382.)

Difficultly sol. in AsBr₃. (W. anorg. 1902, 29, 374.)

Sol. in SbCl₃. (Walden, Z. anorg. 220.)

Sol. in liquid SO₂. (Walden, 1902, 30, 160.)

Hydrazine dissolves 56.4 pts. KBr 13°. (de Bruyn, R. t. c. 1899, 18, 2.)

Sl. sol. in alcohol. (Ballard.)

Sol. in 200 pts. cold, and 16 p 80% alcohol.

Sol. in 180 pts. 90% alcohol. (H. Dingl. 221, 89.)

Sol. in 750 pts. abs. alcohol at 15°.

100 pts. absolute methyl alcohol dissolve 1.51 pts. at 25°; 100 pts. absolute alcohol dissolve 0.13 pt. at 25°. (de phys. Ch. 10, 783.)

Solubility of KBr in methyl alcohol 25°.

P = % by wt. of alcohol in alc.
S = Sp. gr. of alcohol + Aq. sat.
L = millimole KBr in 100 ccm. solution.

P	S 25°/4°
0	1.3797
10.6	1.300
30.8	1.159
47.1	1.058
64.0	0.9801
78.1	0.8906
98.9	0.8411
100	0.8047

(Hers and Anders, Z. anorg. 1907,

100 g. KBr + CH₃OH contain 0 at the critical temp. (Centneruswa Ch. 1910, 72, 437.)

Solubility of KBr in ethyl alcohol + Aq.

% Sol.	Temp. = 30°		Temp. = 40°	
	G. KBr. per 100 g.		G. KBr per 100 g.	
	Solution	Solvent	Solution	Solvent
0	41.62	71.30	43.40	76.65
5	38.98	67.25	40.85	72.70
10	36.33	63.40	38.37	69.00
15	31.09	56.40	33.27	62.30
20	25.98	50.15	28.32	56.45
25	21.24	44.95	23.22	50.46
30	16.27	38.85	18.11	44.25
35	11.50	32.50	13.02	37.40
40	6.90	24.70	7.98	28.90
45	3.09	15.95	3.65	18.95
50	0.87	8.80	1.03	10.45

(Taylor, J. phys. Ch. 1896, 1. 724.)

room temp., 1 pt. KBr by weight is sol.

2 pts. methyl alcohol, D¹⁵ 0.7990.

0 " ethyl " D¹⁵ 0.8100.

8 " propyl " D¹⁵ 0.8160

Rohland, Z. anorg. 1898, 18. 325.)

Solubility of KBr in ethyl alcohol at 0°.

of alcohol l. g. per l. H ₂ O	G. KBr in 1 l. H ₂ O	Mol. solubility
—	536.75	4.51
1/4	529.25	4.45
1/2	502.85	4.22
1	491.75	4.13
2	455.25	3.82

Strong and Eyre, Proc. R. Soc. 1910,
[A] 84. 127.)

g. methyl alcohol dissolve 2.17 g. KBr

g. ethyl alcohol dissolve 0.142 g. KBr

g. propyl alcohol dissolve 0.035 g. KBr

g. isoamyl alcohol dissolve 0.003 g. KBr

Erner and Bissett, Chem. Soc. 1913, 103.

5 g. is sol. in 100 g. propyl alcohol.
Walden, Z. phys. Ch. 1894, 14. 276.)

in 5000 pts. ether (sp. gr. 0.729 at 15°.)
l. c.)

in 1700 pts. alcohol-ether (1 : 1) at 15°.
l. c.)

pts. acetone dissolve 0.023 pt. KBr at
Krug and M'Elroy, J. Anal. Ch. 6. 184.)

Solubility of KBr in acetone + Aq at 25°.

A = cc. acetone in 100 cc. acetone + Aq.

KBr = millimols KBr in 100 cc. of the
solution.

A	KBr	Sp. gr.
0	481.3	1.3793
20	366.7	1.2688
30	310.5	1.2118
40	259.0	1.1558
50	202.9	1.0918
60	144.9	1.0275
70	95.3	0.9591
80	46.5	0.89415
90	10.1	0.8340

(Herz and Knoch, Z. anorg. 1905, 45. 262.)

Solubility of KBr in glycerine + Aq at 25°.

G = g. glycerine in 100 g. glycerine + Aq.

KBr = millimols KBr in 100 cc. of the solu-
tion.

G	KBr	Sp. gr.
0	481.3	1.3793
13.28	444.3	1.3704
25.98	404.0	1.3655
45.36	340.5	1.3594
54.23	310.4	1.3580
83.84	219.25	1.3603
100	172.65	1.3691

(Herz and Knoch, Z. anorg. 1905, 45. 267.)

100 g. 95% formic acid dissolve 23.2 g.
KBr at 18.5°. (Aschan, Chem. Ztg. 1913, 37.
1117.)

Insol. in methyl acetate. (Naumann, B.
1909, 42. 3790); ethyl acetate. (Naumann,
B. 1904, 37. 3601.)

Insol. in benzonitrile. (Naumann, B.
1914, 47. 1370.)

100 ccm. of a sat. solution of KBr in fur-
fural at 25° contain 0.139 pts. by wt. KBr.
(Walden, Z. phys. Ch. 1906, 55. 713.)

Insol. in CS₂. (Arctowski, Z. anorg. 1894,
6. 257.)

Potassium rhodium bromide.

See Bromorhodite, potassium.

Potassium ruthenium tribromide.

See Bromoruthenite, potassium.

Potassium ruthenium tetrabromide.

See Bromoruthenate, potassium.

Potassium selenium bromide.

See Bromoselenate, potassium.

Potassium tellurium bromide.

See Bromotellurate, potassium.

Potassium thallic bromide, KBr , $\text{TlBr}_3 + 2\text{H}_2\text{O}$.

Sol. in H_2O .

3KBr , $2\text{TlBr}_3 + 3\text{H}_2\text{O}$. Sol. in H_2O . (Ram-melsberg.)

Potassium thorium bromide.

Sol. in H_2O . (Berzelius.)

Potassium tin (stannous) bromide, KBr , $\text{SnBr}_2 + \text{H}_2\text{O}$.

Sol. in H_2O . (Benas, C. C. 1884. 958.)

Can be recryst. from HBr or $\text{KBr} + \text{Aq}$. (Richardson, Am. Ch. J. 14. 95.)

2KBr , $\text{SnBr}_2 + 2\text{H}_2\text{O}$. Cannot be recryst. from $\text{HBr} + \text{Aq}$. (Richardson.)

Potassium tin (stannic) bromide, 2KBr , SnBr_4 .

See Bromostannate, potassium.

Potassium uranous bromide, KUBr_2 .

Very sol. in H_2O . (Aloy, Bull. Soc. 1899, (3) 21. 264.)

Potassium uranyl bromide, 2KBr , $\text{UO}_2\text{Br}_2 + 2\text{H}_2\text{O}$.

Very easily sol. in H_2O . (Sendtner.)

Potassium zinc bromide, KBr , $\text{ZnBr}_2 + 2\text{H}_2\text{O}$.

Not hygroscopic. (Ephraim, Z. anorg. 1908, 59. 60.)

2KBr , $\text{ZnBr}_2 + 2\text{H}_2\text{O}$. Not hygroscopic. (Ephraim.)

Potassium bromide ammonia, KBr , 4NH_3 .

(Joannis, C. R. 1905, 140. 1244.)

Potassium bromide ruthenium dihydronitrosochloride, $(\text{NO})\text{Ru}_2\text{H}_2\text{Cl}_2$, 2HCl , 3KBr .

Ppt. Sl. sol. in H_2O . (Brizard, A. ch. 1900, (7) 21. 359.)

Potassium bromoiodide, KBr_2I .

Decomp. rapidly on air. (Wells and Wheeler, Sill. Am. J. 143. 475.)

Potassium carbonyl, $\text{K}_2\text{C}_2\text{O}_2$.

Decomp. by H_2O with explosion. (Joannis, C. R. 116. 158.)

Potassium chloride, KCl .

Sol. in H_2O with absorption of heat.

30 pts. $\text{KCl} + 100$ pts. H_2O at 13.2° lower the temp. 12.6° . (Rüdorff, B. 2. 68.)

100 pts. H_2O dissolve 29.31 pts. KCl at 0° . (Gay-Lussac); 28.5 pts. KCl at 0° (Mulder; Gerardin.)

The saturated solution contains 58 boils at 107.6° (Mulder); contains 59 boils at 108.3° (Legrand); contains and boils at 109.6° (Gay-Lussac); 110° . (Kremers.)

Sol. in 3.016 pts. H_2O at 15° (Gay-Lussac); 3.03 pts. at 17.5° , or 100 pts. H_2O at solve 33 pts KCl . (Schiff.)

100 pts. H_2O at t° dissolve pts. K

t°	Pts. KCl	t°	Pts. KCl	t°
0	29.21	52.39	43.59	100.
19.35	34.53	79.58	50.93	

(Gay-Lussac, A. ch. (2) 11. 30)

100 pts. H_2O dissolve 34.6 pts. KCl at 13.8° ; 35 pts. at 15.6° . (Kopp.)

100 pts. H_2O at 17.5° dissolve 33.34 p sp. gr. of solution is 1.635. (Karsten.)

100 pts. H_2O at 12° dissolve 32 pts., and pts. (Otto-Graham.)

Sol. in 3 pts. H_2O at ord. temp., and 3 H_2O (Bergmann); in 3.33 pts. hot or cold croy; in 3 pts. at 15° , and 1.68 pts. at 110° P.)

Sol. in 3.5 pts. H_2O at 0° , and in less th H_2O (Schubarth); 100 pts. H_2O at 17.5° c 33.0 pts. KCl (Ure's Dict.).

100 pts. H_2O dissolve 35.405 pts. KCl solution has sp. gr. = 1.1809. (Michel an ch. (3) 41. 478.)

100 pts. H_2O dissolve at:

18°	30°	40°	57°
33.6	37.8	40.1	45.0 pts.

(Gerardin, A. ch. (4) 5. 13)

100 pts. H_2O dissolve 33.06–32.6 at 15.6° and sp. gr. of solution = 1. and Keightley, Chem. Soc. (2) 10.

Solubility in 100 pts. H_2O a

t°	Pts. KCl	t°	Pts. KCl	t°
0	28.5	17	33.9	3
1	28.7	18	34.2	3
2	29.0	19	34.4	3
3	29.3	20	34.7	3
4	29.5	21	35.0	3
5	30.0	22	35.3	3
6	30.5	23	35.5	4
7	31.0	24	35.8	4
8	31.5	25	36.1	4
9	31.7	26	36.4	4
10	32.0	27	36.6	4
11	32.3	28	36.9	4
12	32.5	29	37.2	4
13	32.8	30	37.4	4
14	33.1	31	37.7	4
15	33.4	32	38.0	4
16	33.6	33	38.2	5

Solubility in 100 pts., etc.—Continued.

t°	Pts. KCl	t°	Pts. KCl	t°	Pts. KCl
51	43.1	71	48.5	91	54.1
52	43.4	72	48.8	92	54.4
53	43.6	73	49.1	93	54.6
54	43.9	74	49.4	94	54.9
55	44.2	75	49.6	95	55.2
56	44.4	76	49.9	96	55.5
57	44.7	77	50.2	97	55.7
58	44.9	78	50.5	98	56.0
59	45.2	79	50.8	99	56.3
60	45.5	80	51.0	100	56.6
61	45.8	81	51.3	101	56.9
62	46.1	82	51.5	102	57.2
63	46.3	83	51.8	103	57.4
64	46.6	84	52.1	104	57.7
65	46.9	85	52.4	105	58.0
66	47.2	86	52.6	106	58.2
67	47.5	87	52.9	107	58.5
68	47.7	88	53.2	107.65	58.5
69	48.0	89	53.5
70	48.3	90	53.8

Mulder, calculated from his own and other observations, Scheik. Verhandel. 1864. 41.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. KCl	t°	Pts. KCl	t°	Pts. KCl
-11°	24.46	25.7	36.10	64.95	47.17
-6.4	25.78	29.25	37.31	71.65	48.76
0	27.9	38.0	39.71	74.25	49.27
+3.9	29.37	41.45	40.67	80.75	51.24
9.4	30.84	46.15	42.34	86.6	52.53
11.4	32.19	48.8	42.86	91.4	53.49
14.95	32.66	55.1	44.51
19.0	34.32	60.55	45.90

(Coppet, A. ch. (5) 30. 414.)

Solubility is represented by a straight line, of which the formula is 28.51+0.2837t. (Coppet.)

100 pts. H₂O dissolve 29.33 pts. KCl at 4°, 45.5 pts. at 60°. (Andreae, J. pr. (2) 29. 456.)

100 pts. H₂O dissolve at:

0°	100°	130°	180°
29.2	56.5	66	78 pts. KCl.

(Tilden and Shenstone, Lond, R. Soc. Proc. 35. 345.)

Solubility of KCl in 100 pts. H₂O at high temp.

t°	Pts. KCl	t°	Pts. KCl	t°	Pts. KCl
125	59.6	147	70.8	180	77.5
133	69.3	175	75.2

(Tilden and Shenstone, Phil. Trans. 1884. 23.)

If solubility S=pts. KCl in 100 pts. solution, S=20.5+0.1445t from -90° to 110°. (Étard, C. R. 98. 1432.)

Sat. KCl+Aq contains at:

142°	150°	175°	180°
38.6	38.8	41.2	41.8% KCl.

190°	200°	242°	732° (mpt.)
43.2	42.9	47.6	100% KCl.

(Étard, A. ch. 1894, (7) 2. 256.)

100 g. H₂O dissolve 0.488 gram-equivalent KCl at 25°. (van't Hoff and Meyerhoffer, Z. phys. Ch. 1904, 49. 315.)

Solubility of KCl in H₂O at t°.

G. KCl per 100 G. H₂O.

t°	KCl	Sp. gr.
0.70	28.29	1.1540
19.55	34.37	1.1738
32.80	38.32	1.1839
59.85	45.84	1.1980
74.80	49.58	1.2032
89.45	53.38	1.2069
108.0*	58.11	1.2118

* Bpt. of sat. solution.

(Berkeley, Phil. Trans. Roy. Soc. 1904, 203, A. 189.)

Solubility of KCl in 100 g. H₂O at t°.

t°	g. KCl	t°	g. KCl
+18.5	33.3	-4.5	25.9
11.5	31.2	-9	23.9
10	30.8	-8.5	21.5
7.5	29.8	-8	20.0
2.5	28.4	-7	17.5
0	27.5	-6	15.7
-1	27.2	-5.5	14.3

(Meusser, Z. anorg. 1905, 44. 80.)

Sat. KCl+Aq at 25° contains 26.46% KCl. (Foote, Am. Ch. J. 1906, 35. 238.)

28.01 g. KCl are contained in 100 g. solution sat. at 30°. (de Waal, Dissert. 1910.)

36.12 g. KCl are sol. in 100 g. H₂O at 25°. (Amadori and Pampanini, Rend. Acc. Linc. 1911, V. 20. 473.)

4.272 g. mol. are contained in 1 l. solution sat. at 25°. (Herz. Z. anorg. 1911, 73. 274.)

Solubility of KBr at 6°=23.06%; 28.4°=26.91%; 62.6°=31.57°. (Süss, Z. Krist. 1912, 51. 262.)

Solubility at 22°=25.68%. (Brönstedt, Z. phys. Ch. 1912, 80. 208.)

100 mol. H₂O dissolve at:

19.3°	29.7°	40.1°	54.5°
8.2	8.99	9.75	10.39 mol. KCl.

(Sudhaus, Miner. Jahrb. Beil.-Bd. 1914, 37. 18.)

KCl+Aq sat. at 16° has sp. gr.=1.077.
(Stolba, J. pr. 97. 503.)

Sp. gr. of KCl+Aq at 17.5°.

% KCl	Sp. gr.	% KCl	Sp. gr.	% KCl	Sp. gr.
1	1.0062	9	1.0586	17	1.1152
2	1.0125	10	1.0655	18	1.1225
3	1.0189	11	1.0725	19	1.1298
4	1.0254	12	1.0795	20	1.1372
5	1.0319	13	1.0866	21	1.1446
6	1.0385	14	1.0937	22	1.1521
7	1.0451	15	1.1008	23	1.1596
8	1.0518	16	1.1080	24	1.1673

(Schiff, A. 110. 76.)

Sp. gr. of KCl+Aq at 19.5°.

% KCl	Sp. gr.	% KCl	Sp. gr.
5.98	1.0382	21.31	1.1436
11.27	1.0733	25.133	1.1720
16.27	1.1075

(Kremers, Pogg. 95. 119.)

Sp. gr. of KCl+Aq at 15°.

% KCl	Sp. gr.	% KCl	Sp. gr.	% KCl	Sp. gr.
1	1.00650	10	1.06580	19	1.12894
2	1.01300	11	1.07271	20	1.13608
3	1.01950	12	1.07962	21	1.14348
4	1.02600	13	1.08654	22	1.15088
5	1.03250	14	1.09345	23	1.15828
6	1.03916	15	1.10036	24	1.16568
7	1.04582	16	1.10750	24.9*	1.17234
8	1.05248	17	1.11465
9	1.05914	18	1.12179

* Mother liquor.
(Gerlach, Z. anal. 8. 281.)

Sp. gr. of KCl+Aq at 20°, containing mols.
KCl to 100 mols. H₂O.

Mols. KCl	Sp. gr.	Mols. KCl	Sp. gr.
0.5	1.01310	4.0	1.09415
1.0	1.02568	5.0	1.11445
2.0	1.04959

(Nicol, Phil. Mag. (5) 16. 122.)

Sp. gr. of KCl+Aq at 18°.

% KCl	Sp. gr.	% KCl	Sp. gr.	% KCl	Sp. gr.
5	1.0308	15	1.0978	25	1.1408
10	1.0638	20	1.1335

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of KCl+Aq at 0°. S=pts. salt
pts. of solution; S₁=mols. salt i
mols. solution.

S	S ₁	Sp. gr.
20.7840	5.954	1.14
17.7214	4.940	1.12
14.4707	3.922	1.10
11.0757	2.918	1.07
7.5440	1.931	1.05
4.4968	1.123	1.03

(Charpy, A. ch. (6) 29. 23.)

Sp. gr. of KCl+Aq at 25°.

Concentration of KCl+Aq	Sp. gr.
1-normal	1.0466
1/2- "	1.0235
1/4- "	1.0117
1/8- "	1.0059

(Wagner, Z. phys. Ch. 1890, 5. 3)

KCl+Aq containing 5.05% KCl has
20°/20° = 1.0327.

KCl+Aq containing 20.55% KCl i
gr. 20°/20° = 1.1393.

(Le Blanc and Rohland, Z. phys. Ch
19. 272.)

Sp. gr. of KCl+Aq.

g. KCl in 1000 g. of solution	Sp. gr. 16°/16
0	1.000000
0.7140	1.000464
1.5042	1.000975
3.0724	1.001991
8.3165	1.006391

(Dijken, Z. phys. Ch. 1897, 24. 10

Sp. gr. of KCl+Aq at 20.1°, when
cent strength of solution; d=d
density, and w=volume conc. in
cc. $\left(\frac{pd}{100}\right) = w$

p	d	w
36.43	1.853	0.42
31.12	1.1554	0.32
24.79	1.1215	0.27
18.06	1.0866	0.19
13.17	1.0617	0.12
8.412	1.0386	0.08
6.610	1.0297	0.06
4.419	1.0193	0.42
3.456	1.0148	0.04
1.197	1.0040	0.01

(Barnea, J. phys. Ch. 1898, 2. 54

p. gr. of KCl+Aq at t°.

Normality of KCl+Aq.	g. KCl in 100 g. of solution	Sp. gr. t°/4°
3.74	23.93	1.1617
2.65	17.66	1.1166
1.87	12.82	1.0829
0.93	6.64	1.0424

mer, Z. phys. Ch. 1898, 27. 450.)

gr. of KCl+Aq at 18°/4°.

n 100 g. of solution	Sp. gr.
1963	1.0003
2459	0.9995
3342	0.99929
32343	0.99912

, Z. phys. Ch. 1900, 33. 559.)

1 containing 1 pt. KCl in 58.923 g. H₂O at 17° has sp. gr. = 1.0096. (Hittorf, Z. phys. Ch. 1902, 39. 628.)
Sp. gr. at 18°/4° of a normal solution of KCl = 1.0003. (Haigh, J. Am. Chem. Soc. 1912, 34. 1565.)

gr. of sat. KCl+Aq at t°.

g. KCl sol. in 100 g. H ₂ O	Sp. gr.
24.98	1.139
28.50	1.156
31.23	1.168
34.11	1.177
37.28	1.183
40.12	1.190
42.86	1.195
45.48	1.199
48.30	1.203

, J. Russ. phys. Chem. Soc. 1912, 44. 1565.)

r. of dil. KCl+Aq at 20.004°.
g. equiv. KCl per l. at 20.004° compared with H₂O at 20.004° = 1.

conc.	Sp. gr.
1000	1.000,000,0
1001	1.000,004,8
1002	1.000,009,7
1005	1.000,024,2
1010	1.000,048,5
1026	1.000,097,1
1050	1.000,242,6
1100	1.000,483,6

Lee, J. Am. Chem. Soc., 1913, 35. 1687.)

KCl+Aq containing 10% KCl boils at 101.1°; containing 20%, at 103.4°. (Gerlach.)
Sat. KCl+Aq containing 52.7 pts. KCl to 100 pts. H₂O forms a crust at 107.7°; highest temp. observed, 108.5°. (Gerlach, Z. anal. 26. 426.)

B.-pt. of KCl+Aq containing pts. KCl to 100 pts. H₂O. G=according to Gerlach (Z. anal. 26. 438); L=according to Legrand (A. ch. (2) 59. 426).

B.-pt.	G	L	B.-pt.	G	L
100.5°	4.9	4.7	105°	36.2	37.8
101.0	9.2	9.0	105.5	39.3	41.0
101.5	13.1	13.2	106	42.4	44.2
102	16.7	17.1	106.5	45.5	47.4
102.5	20.1	20.9	107	48.4	50.5
103	23.4	24.5	107.5	51.5	53.7
103.5	26.7	28.0	108	54.5	56.9
104	29.9	31.4	108.3	...	59.4
104.5	33.1	34.6	108.5	57.4	...

Precipitated from aqueous solution by HCl +Aq. Much less sol. in very dil. HCl+Aq than in H₂O. (Fresenius.)

Nearly insol. in conc. HCl+Aq.
100 cc. sat. HCl+Aq dissolve 1.9 g. KCl at 17°. (Ditte, A. ch. 1881, (5) 24. 226.)

Solubility of KCl in HCl+Aq at 0°.

G. per 100 cc. of solution	
HCl	KCl
0.0	25.73
1.42	22.69
2.41	20.84
2.59	20.51
4.05	17.71
8.39	11.93
12.40	7.46
14.95	5.60
23.88	1.49
54.20	1.52

(Engel, A. ch. 1888, (6) 33. 377.)

Solubility of KCl in HCl+Aq.

t°	Concentration of HCl g. mol. per 1000 g. H ₂ O	Wt. KCl per 1000 g. H ₂ O	Mol. solubility
0	0	283.55	3.81
"	1/4	267.25	3.59
"	1/2	250.00	3.36
"	1	214.25	2.88
25	0	359.25	4.82
"	1/4	341.55	4.59
"	1/2	324.30	4.35
"	1	289.60	3.89

(Armstrong and Eyre, Proc. R. Soc. 1910 (A) 84. 127.)

100 g. sat. HCl+Aq dissolve 1.9 g. KCl at 20°. (Stoltzenberg, B. 1912, 45. 2248.)

Solubility in HCl+Aq at 25°.

Millimols HCl in 10 ccm.	Millimols KCl in 10 ccm.
...	42.72
5.66	37.49
10.20	33.79
15.90	28.68
20.94	24.74
32.52	17.39

(Herz, Z. anorg. 1912, 73. 275.)

Solubility of KCl in HBr+Aq at 25°.

Millimols HBr in 10 ccm.	Millimols KCl in 10 ccm.
...	42.72
6.61	37.80
34.15	19.57

(Herz, Z. anorg. 1912, 73. 275.)

Sol. in sat. NH₄Cl+Aq with pptn. of NH₄Cl. When action has ceased, the solution at 18.75° contains 31.6% of the mixed salt; or 100 pts. H₂O dissolve 46.1 pts. of the mixed salt, viz, 16.27 pts. KCl and 29.83 pts. NH₄Cl. (Karsten.)

Solubility of KCl in NH₄Cl+Aq at 25°.

Dissolved in 1000 mols H ₂ O	
Mols KCl	Mols NH ₄ Cl
74.2	23.8
67.9	32.5
61.4	52.2
55.5	65.9
50.2	74.4
43.0	96.3
37.6	110.0
37.0	107.5
37.5	109.4
22.6	118.2

(Biltz, Z. anorg. 1911, 71. 174.)

See also NH₄Cl.

Sol. in sat. BaCl₂+Aq with pptn. of BaCl₂ until a state of equilibrium is reached, when 100 pts. H₂O at 16.8° dissolve 45.9 pts. mixed salts, viz. 18.2 pts. BaCl₂ and 27.7 pts. KCl.

See also BaCl₂.

Solubility of KCl in MgCl₂+Aq of given percentage composition.

t°	30%	21.2%	15%	11%
10	1.9%	5.3%	9.9%	14.3%
20	2.6	6.5	11.3	15.9
30	3.4	7.6	12.7	17.5
40	4.2	8.8	14.2	19.0
50	5.0	10.0	15.6	20.5
60	5.8	11.2	17.0	21.9
70	6.5	12.4	18.3	23.2
80	7.3	13.6	19.5	24.5
90	8.1	14.7	20.8	25.8
100	8.9	15.9	22.1	27.1

(Precht and Wittgen, B. 14. 1667.)

Solubility of KCl+NaCl in 20% MgCl₂+Aq

t°	% KCl	% NaCl	t°	% KCl	% NaCl
10	4.2	5.7	60	8.9	6.3
20	5.1	5.8	70	9.9	6.4
30	6.0	5.9	80	10.9	6.6
40	6.9	6.0	90	11.9	6.7
50	7.9	6.1	100	13.0	6.9

(P. and W.)

Sol. in sat. KNO₃+Aq with pptn. of KNO₃

1 litre of the solution contains			
at 14.5°		at 25.2°	
Mol. KCl	Mol. KNO ₃	Mol. KCl	Mol. KNO ₃
3.865	0.0	4.18	0.0
3.810	0.204	4.11	0.136
3.782	0.318	4.07	0.318
3.710	0.615	3.93	0.902
3.667	0.818	3.85	1.212
3.629	0.910	3.81	1.397
3.597	1.176	3.70	1.806
3.582	1.220

(Touren, C. R. 1900, 130. 909.)

Solubility of KCl in KNO₃+Aq.

t°	Concentration of KNO ₃ in g. mol. per 1000 g. H ₂ O	Wt. KCl in 1000 g. H ₂ O	Mol. solubility
0	0	283.55	3.81
"	1/4	284.25	3.81
"	1/2	283.60	3.81
"	1	287.60	3.86
25	0	364.15	4.89
"	1/4	355.00	4.90
"	1/2	361.65	4.86
"	1	358.80	4.81
"	1 1/2	355.20	4.77

(Armstrong and Eyre, Proc. R. Soc. 1910 [A] 84. 127.)

See also KNO₃.

n sat. $\text{NaNO}_3 + \text{Aq}$ without causing
See NaNO_3 .
n sat. $\text{Ba}(\text{NO}_3)_2 + \text{Aq}$ without causing

solubility in $\text{KBr} + \text{Aq}$ at 25.2° .

1 litre of the solution contains	
Mol. KBr	Mol. KCl
0.0	4.18
0.49	3.85
0.85	3.58
1.31	3.19
1.78	2.91
2.25	2.58
2.69	2.33

Touren, C. R. 1900, 130. 1252.)
so KBr.

pts. H_2O dissolve 133.2 pts. KI and
KCl at 21.5° , no matter how pre-
(Rüdorff, B. 6. 484.)
pts. KCl + Aq sat. at $15-16^\circ$ contain
5.37 pts. KCl. 100 pts. KCl + KI +
at $15-16^\circ$ contain 57.80 pts. of the two
(Cl is pptd. by KI. (v. Hauer, J. pr.
)

Solubility in $\text{KI} + \text{Aq}$ at t° .

Sat. solution contains		
% KCl	% KI	% total salt
4.8	50.8	56.6
5.1	51.1	56.2
...	...	57.9
4.2	54.6	58.8
4.7	55.0	59.7
5.7	56.0	61.7
...	...	62.5
4.4	59.5	63.9
5.0	59.6	64.6
...	...	66.2
...	...	66.8
...	63.3	...
7.6	64.8	72.4
8.7	65.4	74.1
8.6	66.0	74.6
10.0	66.5	76.5

Étard, A. ch. 1894, (7) 3. 281.)

solubility of $\text{KCl} + \text{KI}$ in H_2O at 25° .

G. per 100 g. H_2O		
KI	KCl	KI
149.26	19.64	...
144.03	23.75	43.89
137.79	29.56	23.83
132.60	31.38	14.83
133.90	33.68	7.00
106.91	36.12	0.00

ri and Pampanini, Att. Acc. Linc.
1911, 20, II. 475.)

Solubility of KCl in $\text{KOH} + \text{Aq}$ at 0° .

G. per 100 cc. solution	
KCl	KOH
26.83	0.0
23.44	1.33
21.39	2.64
17.39	5.56
13.89	8.46
10.91	11.23
8.64	13.83
6.78	16.43
4.74	19.72

(Engel, Bull. Soc. 1891, (3) 6. 16.)

Solubility in $\text{KOH} + \text{Aq}$ at 20° .

G. KOH in 1 litre	G. KCl in 1 litre	Sp. gr.	Degrees Baumé
10	293	1.185	22.5
20	285	1.185	22.5
30	276	1.190	23.0
40	265	1.192	23.0
50	255	1.195	23.5
60	245	1.200	24.0
70	235	1.200	24.0
80	226	1.205	24.5
90	219	1.205	24.5
100	211	1.210	25.0
110	203	1.210	25.0
120	199	1.215	25.5
130	192	1.215	25.5
140	185	1.220	26.0
150	178	1.225	26.5
160	171	1.225	26.5
170	165	1.230	27.0
180	158	1.235	27.5
190	153	1.240	28.0
200	148	1.245	28.5
210	142	1.250	29.0
220	137	1.255	29.5
230	133	1.260	30.0
240	128	1.265	30.5
250	124	1.270	30.8
260	120	1.275	31.3
270	115	1.280	31.7
280	112	1.285	32.0
290	108	1.290	32.5
300	104	1.295	33.0
310	100	1.300	33.5
320	93	1.305	34.0
340	89	1.315	34.6
350	81	1.320	35.0
360	78	1.325	35.5
370	74	1.330	36.0
390	71	1.340	36.7
400	68	1.345	37.1
410	64	1.350	37.5

Solubility in KOH+Aq at 20°.—Continued.

G. KOH in 1 litre	G. KCl in 1 litre	Sp. gr.	Degrees Baumé
420	61	1.355	38.0
430	58	1.360	38.5
440	55	1.365	38.9
450	53	1.370	39.2
460	50	1.375	39.5
470	47	1.380	40.0
480	44	1.385	40.2
490	42	1.390	40.6
500	40	1.397	41.0
510	38	1.405	41.5
520	35	1.410	42.0
530	33	1.415	42.3
540	31	1.420	42.6
550	29	1.425	43.0
560	27	1.430	43.5
570	25	1.435	43.7
580	24	1.440	44.0
590	23	1.445	44.3
600	22	1.450	44.6
610	21	1.455	45.0
620	20	1.460	45.5
630	18	1.465	45.9
640	17	1.470	46.2
650	16	1.475	46.5
660	15	1.480	46.8
670	15	1.485	47.0
680	15	1.490	47.5
690	15	1.495	47.9
700	14	1.500	48.2
710	14	1.505	48.5
720	13	1.510	48.8
730	13	1.515	49.1
740	13	1.520	49.5
750	13	1.525	49.7
760	12	1.530	50.0
770	12	1.535	50.3
780	12	1.540	50.6
790	11	1.545	51.0
800	11	1.550	51.3
810	10	1.560	51.5
820	10	1.565	51.8
830	9	1.570	52.2
840	9	1.575	52.6
850	9	1.580	53.0

(Winteler, Z. Elektrochem, 1900, 7. 360.)

KCl+NaCl.

100 pts. KCl+NaCl+Aq sat. at 13–16° contain 30.18 pts. of the two salts. (v. Hauer.)

100 pts. H₂O dissolve 13.92 pts. KCl and 30.65 pts. NaCl at 15.6°, and solution has sp. gr = 1.233. (Page and Keightley.)

100 pts. H₂O dissolve 10.11 pts. KCl, 32.15 pts. NaCl, and 4.69 pts. K₂SO₄, and solution has sp. gr. = 1.250. (P and K.)

100 pts. H₂O dissolve 29.9 pts. NaCl and 15.7 pts. KCl at 18.8°. (Rüdorff.)

Solubility of KCl+NaCl in H₂O at 100 pts. H₂O dissolve pts. KCl and pts. NaCl.

t°	Pts. KCl	Pts. NaCl	t°	Pts. KCl
10	12.5	29.7	60	24.6
20	14.7	29.2	70	27.3
30	17.2	28.7	80	30.0
40	19.5	28.2	90	32.9
50	22.0	27.7	100	34.7

(Precht and Wittgen, B. 14. 106)

100 pts. H₂O dissolve 13.99 pts. KCl and 30.65 pts. NaCl = 44.53 pts. mixed salts (Nicol, Phil. Mag. (5) 31. 385)

Solubility of KCl in NaCl+Aq at 100 pts. H₂O

G. per 100 g. H ₂ O	NaCl	KCl
0.0		34.52
6.5		29.37
13.0		4.71
19.5		0.42

(Nicol, Phil. Mag. 1891, 31. 36)

Solubility of KCl+NaCl at 100 pts. H₂O

t°	G. per 100 g. H ₂ O	KCl	NaCl
25	15.8	1	
"	29.0	3	
80	30.0	2	
"	26.4	3	

(Soch, J. phys. Ch. 1896, 2. 4)

Solubility of KCl+NaCl at 100 pts. H₂O

G. salts in 100 g. H ₂ O	NaCl	KCl	Solub.
9.89		28.34	KCl
18.35		22.75	
29.88		16.28	KCl+
31.57		10.91	NaCl
33.17		5.65	

(Uyeda, Mem. Col. Sc. Kyoto, 1910)

100 g. H₂O sat. with NaCl dissolves 13.92 g. KCl at 25°.

100 g. H₂O sat. with K₂SO₄ dissolves 30.65 g. NaCl at 25°. (phys. Ch. 1904, 49. 315)

Solubility in NaCl+Aq at 20°, 30°, 40°, 50°, 60°, 70°, 80°, 90°, 100°. Tables given in the original; each salt diminishes the solubility of the other. (Leather, Chem. Soc. 1914)

lity of KCl+NaCl in HCl+Aq at 25°.

HCl	% NaCl	% KCl
3.61	19.95	10.90
7.16	10.65	7.58
11.65	3.56	3.80
16.73	2.03	2.86
21.78	0.18	1.27

s, J. Am. Chem. Soc. 1915, 37. 846.)
also under NaCl.

l+SrCl₂.
pts. H₂O dissolve 11.2 pts. KCl and
ts. SrCl₂ at 14.5°. (v. Hauer.)

rCl₂+Aq. sat. at 14.5 is sat. with KCl
ie temp., 100 pts. H₂O dissolve:

...	33.2	11.2	...
...	...	48.6	50.7
...	...	59.8	...

Mulder, Scheik. Verhandel. 1864.)

+(NH₄)₂SO₄.
solution of KCl+(NH₄)₂SO₄ at b.-pt.
ooled to 14° has different composition
at. solution of (NH₄)Cl and K₂SO₄, and
position is changed by warming it with
KCl or (NH₄)₂SO₄. (Rüdorff.)

+K₂SO₄.
s. H₂O contain the following amounts
lt at 18.75°: (1) sat. with KCl alone;
) sat. first with KCl then with K₂SO₄;
) sat. with K₂SO₄ and KCl together;
) sat. first with K₂SO₄ then with KCl;
) sat. with K₂SO₄ alone.

	1	2	3	4	5
...	34.5	32.96	33.12	33.12	...
...	...	1.79	1.75	1.83	10.8

(Karsten.)

s. H₂O sat. with both K₂SO₄ and KCl
contain the following amounts.

At 14.8°			
...	33.5	28.2	...
...	...	2.0	10.3

At 15.8°			
...	33.6	27.9	...
...	...	2.3	10.4

At 16.1°			
...	33.6	27.1	...
...	...	3.3	10.4

(Kopp, A. 34. 264.)

Sat. K₂SO₄+Aq dissolves KCl only with
pptn. of K₂SO₄, but sat. KCl+Aq dissolves
some K₂SO₄ without any separation. (Kar-
sten.)

Solubility of KCl+K₂SO₄: 100 pts. H₂O
dissolve at t°.

t°	Pts. KCl	Pts. K ₂ SO ₄	t°	Pts. KCl	Pts. K ₂ SO ₄
10	30.9	1.32	60	43.8	1.94
20	33.4	1.43	70	46.5	2.06
30	36.1	1.57	80	49.2	2.21
40	38.7	1.68	90	52.0	2.38
50	41.3	1.82	100	54.5	2.53

(Precht and Wittgen.)

100 g. H₂O dissolve 34.76 g. KCl+2.93 g.
K₂SO₄ at 25°. (Van't Hoff and Meyerhoffer,
Z. phys. Ch. 1898, 27. 75.)

Sol. in 20% KC₂H₃O₂+Aq. (Stromeyer.)
Quickly attacked by liquid NO₂ in the
presence of traces of moisture, with evolution
of Cl₂. (Frankland, Chem. Soc. 1901, 79.
1361.)

Sl. sol. in liquid NH₃. (Franklin, Am. Ch.
J. 1898, 20. 829.)

Easily sol. in liquid HF. (Franklin, Z.
anorg. 1905, 46. 2.)

100 g. hydrazine dissolves 8.5 pts. KCl at
12.5-13°. (de Bruyn, R. t. c. 1899, 18. 297.)

100 g. anhyd. hydroxylamine dissolve 12.3
g. KCl at 17-18°. (de Bruyn, Z. phys. Ch.
1892, 10. 782.)

100 pts. alcohol of 0.900 sp. gr. dissolve 4.62 pts.;
0.872, 1.66 pts.; 0.834, 0.38 pt.; 0.817, 0.00 pt. KCl.
(Kirwan.)

Sol. in 48 pts. boiling alcohol. (Wenzel.)
Insol. in absolute alcohol containing LiCl. (Mits-
cherlich.)

At 15°, 100 pts. alcohol of p percentage by
volume (S=sp. gr.) dissolve pts. KCl as
follows:

p	10	20	30	40
S	0.984	0.972	0.958	0.940
KCl	19.8	14.7	10.7	7.7

p	50	60	80
S	0.918	0.896	0.848
KCl	5.0	2.8	0.45

(Schiff, A. 118. 365.)

100 pts. of a mixture of 40% alcohol with
60% H₂O dissolve 9.2 pts. KCl at 15°.
(Schiff.)

Insol. in absolute alcohol or in 96% alcohol
at 15° or below. At 20°, 100 pts. of the latter
dissolve 0.04 pt.; at 25°, 0.06 pt.; at 30°, 0.20
pt. KCl. Dilute alcohol dissolves less KCl
than the contained H₂O would dissolve by
itself.

Solubility in dil. alcohol. D = sp. gr. of alcohol; S = solubility in 100 pts. alcohol at t°.

D = 0.9904		D = 0.9848		D = 0.9793		D = 0.9726	
t°	S	t°	S	t°	S	t°	S
0	23.2	4	20.9	4	16.4	3	12.2
4	24.8	20	25.5	21	20.3	5	12.7
22	29.4	27	26.6	28	22.0	16	15.4
25	30.2	30	27.5	43	25.6	20	16.1
34	32.8	37	29.0	25	17.3
52	37.5	60	35.2	34	19.0

D = 0.9573		D = 0.9390		D = 0.8967		D = 0.8244	
t°	S	t°	S	t°	S	t°	S
10	8.8	2	4.2	12	2.87	4	0.00
11	9.0	7	5.1	31	4.35	15	0.00
17	10.3	16	6.4	47	4.88	20	0.04
30	12.5	30	8.5	65	5.65	25	0.06
40	13.9	38	9.6	32	0.20
60	16.7	57	11.3

(Gerardin, A. ch. (4) 5. 140.)

Solubility of KCl in dil. alcohol at 14.5°

Sp. gr.	100 ccm. contain		
	Alcohol	Water	KCl
1.1720	...	88.10	29.10
1.1542	2.79	85.78	26.85
1.1365	4.98	84.00	24.67
1.1075	10.56	79.63	20.56
1.1085	15.57	75.24	17.24
1.0545	20.66	70.52	14.27
1.0455	24.25	67.05	13.25
0.9695	40.42	50.18	6.35
0.9315	48.73	40.60	3.82
0.8448	68.63	15.55	0.30

(Bodländer, Z. phys. Ch. 7. 316.)

Solubility of KCl in ethyl alcohol.

(G. KCl per 100 g. alcohol + Aq.)

Wt. % alcohol	at 30°	at 40°	Wt. % alcohol	at 30°	at 40°
0	38.9	41.8	43.1	11.1	13.1
5.28	33.9	35.9	55.9	6.8	8.2
9.43	30.2	33.3	65.9	3.6	4.1
16.9	24.9	27.6	78.1	1.3	1.6
25.1	19.2	21.8	86.2	0.4	0.5
34.1	15.6	17.2

(Bathrick, J. phys. Ch. 1896, 1. 160.)

Solubility of KCl in ethyl alcohol

Concentration of alcohol Mol. g. alcohol per 1000 g. H ₂ O	Solubility in 1000 g. H ₂ O	Mol solu
...	285.15	3
0.25	277.95	3
0.50	271.10	3
1.00	265.50	3
3.00	208.80	2

(Armstrong and Eyre, Proc. Roy. Soc. (A) 84. 127.)

Solubility of KCl in ethyl alcohol 25°.

Wt. % alcohol	G. KCl per 100 cc. sat. solution	Wt. % alcohol	G. sat
0	31.18	60	...
10	23.93	70	...
20	17.89	80	...
30	13.27	90	...
40	9.40	100	...
50	6.26		

(McIntosh, J. phys. Ch. 1903, 7.

100 pts. absolute methyl alcohol 0.5 pt. at 18.5°; 100 pts. absolute alcohol dissolve 0.034 pt. at 18.5°. (Z. phys. Ch. 10. 783.)

100 pts. 40% wood alcohol dissolve KCl. (Schiff.)

Solubility of KCl in methyl alcohol 25°.

P = % by wt. of alcohol in alcohol
S = Sp. gr. alcohol + Aq sat. with KCl
L = millimols KCl in 100 ccm. of solution.

P	S 25°/4°	
0	1.1782	41
10.6	1.125	32
30.8	1.033	18
47.1	0.9679	10
64.0	0.9064	4
78.1	0.8607	2
98.9	0.8242	
100	0.7937	

(Hera and Anders, Z. anorg. 1907, 8

Solubility of KCl in methyl alcohol.

Concentration of alcohol. Mol. g. alcohol per 1000 g. H ₂ O	Solubility 1 in 1000 g. H ₂ O	Molecular solubility
...	283.55	3.81
0.25	280.00	3.76
0.50	276.35	3.71
1.00	267.85	3.60
3.00	238.10	3.18
...	364.15	4.89
0.25	361.90	4.86
0.50	357.10	4.79
1.00	348.70	4.67
3.00	324.15	4.35

Strong and Eyre, Proc. Roy. Soc. 1910 (A) 84. 127.)

Room temp. 1 pt. by weight is sol. in:
pts. methyl alcohol, D¹⁵ 0.7990.
" ethyl " D¹⁵ 0.8035.
1. in propyl alcohol. (Rohland, Z. 1899, 18. 325.)
g. methyl alcohol dissolve 0.53 g. KCl
g. ethyl alcohol dissolve 0.022 g. KCl
g. propyl alcohol dissolve 0.004 g. KCl
g. isoamyl alcohol dissolve 0.0008 g. KCl at 25°.
Mer and Bissett, Chem. Soc. 1913, 103.

1. in propyl alcohol. (Schlamp, Z. Ch. 1894, 14. 276.)

Solubility of KCl in propyl alcohol.

Concentration of alcohol. Mol. g. per 1000 g. H ₂ O	Solubility in 1000 g. H ₂ O	Molecular solubility
...	283.55	3.81
0.25	274.10	3.68
0.50	265.45	3.56
1.00	248.0	3.33
...	365.10	4.90
0.25	355.40	4.77
0.50	347.70	4.67
1.00	331.50	4.45

Strong and Eyre, Proc. Roy. Soc. 1910, (A) 84. 127.)

1. in fusel-oil. (Gooch, Am. Ch. J. 9. 184.)
sl. sol. in mixture of equal pts. ab-
solute alcohol and ether. (Berzelius.)

500 mg. KCl treated with 10 g. of above mixture yield only 0.3 mg. to the liquid. (Lawrence Smith, Am. J. Sci. 16. 56.)
Insol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184; Eidmann, C. C. 1899, II. 1014.)

Solubility of KCl in acetone+Aq at t.°

t°	% acetone	100 g. of the solution contain		
		G. H ₂ O	G. acetone	G. KCl
30°	0	72.73	0.00	27.27
	5	71.15	3.74	25.11
	9.09	69.62	6.96	23.42
	20	64.88	16.22	18.90
	30	59.49	25.45	15.06
	40	53.17	35.52	11.31
	50	45.98	45.98	8.04
	60	37.97	56.91	5.12
	70	29.22	68.18	2.60
	80	19.82	79.43	0.76
	90	9.98	89.88	0.13
	100	0.00	100.00	0.00
40°	0	71.31	0.00	28.69
	5	69.62	3.67	26.72
	9.09	67.88	6.79	25.33
	15	65.15	11.51	32.34
	20	62.97	15.75	21.28
	80	19.81	79.34	0.58
	85	14.94	84.66	0.40
	90	10.00	89.84	0.16
	95	4.97	94.96	0.07
	100	0.00	100.00	0.00

Since there is but one liquid phase here, these figures represent the solubility of KCl in acetone+Aq at 30° and 40°.

(Snell, J. phys. Chem. 1898, 2. 484.)

The addition of KCl to mixtures of acetone and H₂O will cause a division into two layers. The following table gives the temp. at which sat. solutions of KCl in acetone+Aq of varying concentrations separate into two layers and also the compositions of the sat. solutions of KCl in acetone+Aq.

% acetone	Temp. of division	100 g. of solution contain		
		G. H ₂ O	G. acetone	G. KCl
26	46.5°
30	40.0	59.36	25.44	15.20
40	34.2	53.21	35.47	11.32
50	32.6	45.97	45.97	8.06
60	33.3	37.86	56.80	5.34
70	35.5	29.09	68.25	2.66
75	39.0
80	45.6	19.80	79.20	1.00

(Snell.)

The following table gives the compositions of the solutions of KCl in acetone+Aq at the points at which the solution just divides into two layers. Temp.=40°.

100 g. of the solution contain		
G. H ₂ O	G. acetone	G. KCl
56.68	28.63	14.68
53.05	35.67	11.29
50.34	39.82	9.83
47.60	43.83	8.58
44.35	48.36	7.29
42.68	50.75	6.57
38.53	56.26	5.21
36.59	58.84	4.57
32.37	64.18	3.45
30.62	66.43	2.95
28.12	69.45	2.44

(Snell.)

The addition of KCl to aqueous acetone causes the separation of the liquid into two layers. The following table gives the composition of these layers at 40°.

Upper layer contains per 100 g. of solution		
G. H ₂ O	G. acetone	G. KCl
55.20	31.82	12.99
54.27	36.69	12.03
53.27	35.44	11.29
51.69	37.76	10.55
51.23	48.50	10.27
50.34	39.88	9.77
49.08	41.67	9.26
48.02	43.18	8.79
47.62	43.73	8.64
46.49	45.34	8.17
45.65	46.52	7.83
45.64	46.57	7.79
58.99	25.24	15.77

Lower layer contains per 100 g. of solution		
G. H ₂ O	G. acetone	G. KCl
28.14	69.42	2.44
29.45	67.83	2.72
30.96	65.97	3.07
31.83	64.83	3.33
32.64	63.79	3.56
34.07	62.01	3.92
35.27	60.49	4.24
37.44	57.67	4.89
38.00	56.96	5.04
38.68	56.17	5.25
39.98	54.36	5.66
40.41	53.78	5.81
23.66	74.91	1.43

(Snell.)

Solubility in acetone+Aq at 20°. A=ccm. acetone in 100 ccm. acetone solution. KCl=millimols KCl in 100 ccm. solution.

A	KCl
0	410.5
10	351.7
20	286.6
30	223.7
40	166.5
50	115.4
60	71.2
70	38.5
80	12.9
90	2.0
100	...

(Herz and Knoch, Z. anorg. 1904, 41)

Solubility of KCl in glycerine+Aq G=g. glycerine in 100 g. glycerine KCl= millimols KCl in 100 cc. of the solution.

G	KCl	Sp.
0	424.5	1.1
13.28	383.4	1.1
25.98	339.3	1.1
45.36	271.4	1.2
54.23	238.5	1.2
83.84	149.0	1.2
100	110.6	1.2

(Herz and Knoch, Z. anorg. 1905, 4)

Insol. in CS₂. (Baeyer; Arcton anorg. 1894, 6. 257.) Insol. in benzonitrile. (Naumann, 47. 1370.) Insol. in methyl acetate. (Naum 1909, 42. 3790); ethyl acetate. (N B. 1910, 43. 314.)

Solubility of KCl in organic compounds at 25°.

Compound	G. comp. per l. H ₂ O	G. H ₂ O
Water	...	
Acetaldehyde	11.01	
Paraldehyde	11.07	
Glycerol	13.01	
Glycol	15.51	
"	62.05	
Mannitol	45.53	
"	136.50	

(Armstrong and Eyre, Proc. Roy. Soc. A, 84. 234.)

solubility in pyridine+Aq at 10°.

Solvent	100 g. of the solution contain g. KCl
Pyridine	
0	23.79
10	19.76
20	16.37
30	13.19
40	10.05
50	6.34
60	3.335
70	1.245
80	0.24
90	0.039
100	...

voeder, J. pr. 1908, (2) 77. 268.)

in anhydrous pyridine and in 97% Aq.

sol. in 95% pyridine+Aq.

in 93% pyridine+Aq. (Kahlenm. Chem. Soc. 1908, 30. 1107.)

n. of a sat. solution of KCl in fur-25° contain 0.085 pts. by wt. KCl. Z. phys. Ch. 1906, 55. 713.)

H₂O dissolve 246.5 g. sugar+44.8 g. 1.25°; 100 g. sat. solution contain sugar+11.33 g. KCl. (Köhler, Z. erind, 1897, 47. 447.)

solubility in glucose+Aq at 25°.

ion of mol. H ₂ O	Solubility in 1000 g. H ₂ O	Molecular solubility
	362.70	4.86
	366.10	4.91
	369.85	4.96
	376.25	5.04
	402.25	5.39

ing and Eyre, Proc. Roy. Soc. 1910, 84. 127.)

manganese chloride, K₂MnCl₄. H₂O; less sol. in NH₄Cl+Aq; un-Neuman, M. 1894, 15. 492.)

rhodium chloride. ororhodite, potassium.

ruthenium sesquichloride. ororuthenite, potassium.

ruthenium tetrachloride. ororuthenate, potassium.

tellurium chloride. xrotellurate, potassium.

thallic chloride, 3KCl, TiCl₃+I₂O. Not decomp. by boiling H₂O. berg.)

Potassium thorium chloride, KCl, 2ThCl₄+18H₂O. Deliquescent; sol. in H₂O and alcohol. (Berzelius.)

Potassium tin (stannous) chloride (Potassium chlorostannite), KCl, SnCl₂+H₂O. Decomp. by H₂O; sol. in hot HCl or KCl+Aq. (Remsen and Richardson, Am. Ch. J. 14. 90.) 2KCl, SnCl₂+H₂O. Partially decomp. by dissolving in H₂O. (Rammelsberg, Pogg. 94. 507.) +2H₂O. Very sol. in hot, and but slightly in cold HCl+Aq or KCl+Aq. (Remsen and Richardson.) 4KCl, SnCl₂+3H₂O. (Poggiale, C. R. 20. 1182.) Does not exist. (Remsen and Richardson.)

Potassium tin (stannic) chloride, 2KCl, SnCl₄. See Chlorostannate, potassium.

Potassium tungsten chloride, K₂(OH)WCl₄. Decomp. by moisture. Insol. in organic solvents. (Olsson, B. 1913, 46. 581.) K₂W₂Cl₄. Sol. in H₂O. Nearly insol. in most organic solvents. (Olsson.)

Potassium uranium chloride, UCl₄, 2KCl. Very hygroscopic; sol. in H₂O with decomp.; sol. in acetic acid. Decomp. by alcohol. Nearly insol. in ether. (Aloy, Bull. Soc. 1899, (3) 21. 264.)

Potassium uranyl chloride, K₂(UO₂)Cl₄. Very sol. in H₂O. Moderately sol. in dil. alcohol. (Aloy, Dissert. 1901.) +2H₂O. Very sol. in H₂O and alcohol. (Arfvedson.) Sol. in H₂O, with decomp. and separation of KCl, unless H₂O is acidulated with HCl. (Peligot, A. ch. (3) 5. 37.)

Solubility in H₂O at t°.

t°	100 pts. of the solution contain			Solid phase
	Pts. UO ₂	Pts. Cl	Pts. K	
0.8	38.57	13.59	3.86	UO ₂ Cl ₂ , 2KCl, 2H ₂ O +KCl
14.9	33.71	13.51	...	"
17.5	37.36	14.50	5.27	"
25.0	35.01	15.26	...	"
41.5	35.27	15.92	7.39	"
50.0	34.18	16.56	...	"
60.0	34.19	17.25	9.14	UO ₂ Cl ₂ , 2KCl, 2H ₂ O.
71.5	33.55	17.44	9.28	"
78.5	35.26	18.24	9.95	"

(Rimbach, B. 1904, 37. 483.)

Potassium uranyl chloride is decomp. by H_2O at temp. below 60° . Above 60° , it is sol. in H_2O without decomp.

Potassium vanadium chloride, $VK_2Cl_3 + H_2O$.

Difficultly sol. in H_2O and alcohol. (Stähler, B. 1904, 37. 4412.)

Potassium yttrium chloride.

Sol. in H_2O with evolution of heat.

Potassium zinc chloride, $2KCl, ZnCl_2$.

Very deliquescent. Sol. in 1 pt. cold, and in all proportions of hot H_2O . (Pierre, A. ch. (3) 16. 248.)

+ H_2O . Not very deliquescent. Can be recryst. (Ephraim, Z. anorg. 1908, 59. 58.)

$KCl, ZnCl_2 + 2H_2O$. Not deliquescent. Cannot be recryst. without decomp. (Ephraim.)

Potassium chloriodide, KCl_2I .

Very unstable. (Wells and Wheeler, Sill. Am. J. 143. 475.)

KCl_2I . Sol. in H_2O with decomp. Ether dissolves out ICl_3 . (Filhol, J. Pharm. 25. 433.)

Potassium fluoride, KF or K_2F_2 .

Very deliquescent. Very sol. in H_2O . Sl. sol. in $HF + Aq$. Easily sol. in conc. $KC_2H_3O_2 + Aq$. Insol. in alcohol. (Berzelius.) Sol. in dilute alcohol. (Stromeyer, A. 100. 83.)

Sp. gr. of aqueous solution of KF at 18° containing—

5	10	20	30	40% KF
1.041	1.064	1.117	1.272	1.378

(Kohlrausch, W. Ann. 1879. 1.)

Solubility in $HF + Aq$ at 21° .

(G. per 100 g. H_2O .)

HF	KF	HF	KF
0.0	96.3	13.95	31.4
1.21	72.0	15.98	33.4
1.61	61.0	17.69	35.6
3.73	40.4	20.68	38.4
4.03	32.5	28.60	46.9
6.05	30.4	41.98	61.8
9.25	29.9	53.71	74.8
11.36	29.6	74.20	105.0
12.50	30.5	119.20	169.5

(Ditte, C. R. 1896, 123. 1282.)

Easily sol. in liquid HF . (Franklin, Z. anorg. 1905, 46. 2.)

Very al sol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

KF will "salt out" acetone from solution. The table shows the composition of the solutions at the point at which inhomogeneous solutions of acetone and H_2O just become homogeneous at 20° .

100 g. of the solution contain:

G. KF	G. H_2O	G. acetone	G. KF	G. H_2O
5.75	58.91	35.34	0.61	31.92
5.00	56.28	38.72	0.50	29.92
3.84	52.25	43.91	28.42	69.76
3.06	49.05	47.89	25.74	71.24
2.61	46.84	50.55	22.35	72.96
2.22	44.79	52.99	20.28	73.80
14.95	73.66	11.39	18.71	74.10
11.46	70.77	17.77	16.31	73.95
9.17	67.30	23.53	12.40	72.01
7.72	64.01	28.27	33.86	65.73
7.07	62.03	30.90	29.97	68.54
6.43	60.50	33.07	22.06	73.41
1.38	40.55	58.06	17.82	74.01
0.979	36.42	62.60	14.34	73.29
0.693	32.69	66.61	44.24	55.52
0.57	31.50	67.93	33.34	66.66
0.89	35.74	63.36	29.86	68.54
0.75	33.84	65.41	24.38	72.16

At the first quadruple point of hydrate, acetone, water and vapor equilibrium the upper layer contains acetone, while the lower layer contains 100 g., 46.3 g. KF . A saturated solution will thus dehydrate acetone to the extent of 98%.

(Frankforter and Cohen, J. Am. Ch. 1914, 36. 1115.)

Similar data are given for KF in propyl alcohol by Frankforter and Cohen (J. phys. Ch. 1913, 17. 402.)

+ $2H_2O$. Very deliquescent. (C. ch. (6) 3. 20.)

Sat. aq. solution at 18° contains KF (de Forcrand, C. R. 1911, 152. 1075.)

Sp. gr. of solution sat. at $18^\circ = 1.1$ contains 48% KF . (Mylius and 1897, 30. 1718.)

+ $4H_2O$. Not deliquescent. (de C. R. 1911, 152. 1075.)

Sat. aq. solution at 18° contains KF . (de Forcrand, C. R. 1911, 152. 1075.)

Potassium hydrogen fluoride, KHF_2 .

Easily sol. in H_2O . Sl. sol. in containing HF . Easily sol. in conc. K_2Aq . Sol. in dil. alcohol, but insol. in alcohol.

$KF, 2HF$. Deliquescent. Dissolves in H_2O with absorption of heat. (Moissan, 106. 547.)

$KF, 3HF$. As above. (Moissan,

n manganic fluoride.
omanganate, potassium.

n scandium fluoride, K_2ScF_6 .
 H_2O . Decomp. by acids. (R. J. anorg. 1914, 86. 275.)

n silicon fluoride.
osilicate, potassium.

n tantalum fluoride.
otantalate, potassium.

n tellurium fluoride, KF, TeF_4 .
 sp. by H_2O . (Högbom, Bull. Soc. J.)

n thallic fluoride, $2TlF_3, KF$.
 sp. by moisture. Insol. in HF . e, A. 1909, 366. 226.)

n thorium fluoride, $2KF, ThF_4 + 2$.
 insol. in H_2O . Sol. in $HF + Aq.$ hF_4 . Precipitate. (Chydenius.)

m tin (stannous) fluoride, $2KF, F_2 + H_2O$.
 H_2O . (Wagner, B. 19. 896.)

m tin (stannic) fluoride.
uostannate, potassium.

m titanium tetrafluoride.
uotitanate, potassium.

m titanium sesquifluoride, $4KF, F_6$.
 itate. Very sl. sol. in H_2O . Sol. in s. (Piccini, C. R. 97. 1064.)
so Fluosesquititanate, potassium.

m titanyl fluoride.
uoxypertitanate, potassium.

m tungstyl fluoride.
uoxytungstate, potassium.

m uranium fluoride, KF, UF_4 .
 in H_2O and dil. acids. Difficultly conc. $HCl + Aq.$ Sol. in conc. H_2SO_4 . J. B. 1866. 212.)

m uranyl fluoride.
uoxyuranate, potassium.

m vanadium sesquifluoride.
uovanadate, potassium.

m vanadium tetrafluoride (?).
 sol. in H_2O . Insol. in alcohol. is.)

Potassium zinc fluoride, KF, ZnF_2 .
 Sol. in H_2O . (R. Wagner.)
 $2KF, ZnF_2$. Sol. in H_2O . (Berzelius.)

Potassium zirconium fluoride.
 See Fluozirconate, potassium.

Potassium fluoride hydrogen peroxide, KF, H_2O_2 .
 Not hygroscopic. Very sol. in H_2O . Is not decomp. at 70° and only partially so at 110° . (Tanatar, Z. anorg. 1901, 28. 255.)

Potassium fluoride vanadic acid.
 See Fluoxyvanadate, potassium.

Potassium hydride, KH .
 Decomp. by H_2O . Insol. in oil of turpentine, benzene, ether and CS_2 . (Moissan, C. R. 1902, 134. 18.)

Potassium hydrosulphide, KSH .
 Very deliquescent, and sol. in H_2O with gradual decomp. Crystallizes with $\frac{1}{2}H_2O$. Sol. in alcohol.

Potassium hydroxide, KOH .
 Very deliquescent, and sol. in H_2O with evolution of much heat. 100 pts. KOH , exposed over H_2O at $16-20^\circ$ take up 460 pts. H_2O in 56 days. (Mulder.)
 1 pt. KOH dissolves in 0.5 pt. cold H_2O (Lowitz); in 0.47 pt. cold H_2O (Bineau, C. R. 41. 509); in 1 pt. H_2O . (Abl.)

Solubility of KOH in H_2O at t° .

t°	G. KOH per 100 g.		Solid phase
	H_2O	solution	
—22	3.7	3.6	Ice
—20.7	22.5	18.4	"
—65.2	44.5	30.8	"
—36.2	36.2	26.6	$KOH.4H_2O$
—32.7	77.94	43.8	"
—33	80	44.4	$KOH.4H_2O + KOH.$ $2H_2O$
—23.2	85	45.9	$KOH.2H_2O$
0	97	49.2	"
10	103	50.7	"
15	107	51.7	"
20	112	52.8	"
30	126	55.76	"
32.5	135	57.44	$KOH.2H_2O + KOH.$ H_2O
50	140	58.33	$KOH.H_2O$
100	178	64.03	"
125	213	68.06	"
143	311.7	75.73	"

(Pickering, Chem. Soc. 1893, 63. 908.)

100 pts. KOH are sol. in 93.4 pts. H_2O at 15° or 100 pts. H_2O dissolve 107 pts. KOH at 15° . Sp. gr. = 1.5355 at 15° .

All higher values found in solubility tables are incorrect. (Ferchland, Z. anorg. 1902, 30. 133.)

100 g. sat. aq. solution at 15° contain 50.48 g. KOH. (de Forcrand, C. R. 1909, 149. 719.)

Sat. KOH + Aq boils at 157.7° (Griffiths); 340° (Gerlach).

B.-pt. of KOH + Aq containing pta. KOH to 100 pta. H₂O.

B.-pt.	Pta. KOH	B.-pt.	Pta. KOH
105°	20.5	215°	210.5
110	34.5	220	219.8
115	46.25	225	230.0
120	57.5	230	240.9
125	67.5	235	251.9
130	76.8	240	263.1
135	85.0	245	274.4
140	92.5	250	285.7
145	99.8	255	298.5
150	106.5	260	312.5
155	114.05	265	328.0
160	121.7	270	343.5
165	129.35	275	359.0
170	137.0	280	375.0
175	144.8	285	391.0
180	152.6	290	408.2
185	160.4	295	425.5
190	168.2	300	444.4
195	176.5	310	484.0
200	185.0	320	526.3
205	193.5	330	571.5
210	202.0	340	623.6

(Gerlach, Z. anal. 26. 464.)

Sp. gr. and b.-pt. of KOH + Aq according to Dalton.

% K ₂ O	Sp. gr.	B.-pt.	% K ₂ O	Sp. gr.	B.-pt.
4.7	1.00	100.58°	36.8	1.44	123.80°
9.5	1.11	101.11	39.6	1.47	129.44
13.0	1.15	101.00	42.0	1.52	135.50
16.2	1.19	103.33	46.7	1.60	143.33
19.5	1.23	104.44	51.2	1.68	160.00
23.4	1.28	106.00	56.8	1.78	188.22
26.3	1.33	109.44	63.6	1.88	215.56
29.4	1.36	112.22	72.4	2.00	315.56
32.4	1.39	115.56	84.0	2.2	red heat
34.4	1.42	118.80	100	2.4	

Sp. gr. of KOH + Aq at 15°.

% K ₂ O	Sp. gr.	% K ₂ O	Sp. gr.	% K ₂ O	Sp. gr.
0.568	1.0030	10.750	1.1050	20.935	1.2208
1.607	1.0153	11.862	1.1182	21.500	1.2342
2.820	1.0300	13.013	1.1308	22.032	1.2493
3.961	1.0469	14.145	1.1437	23.764	1.2648
5.002	1.0678	15.277	1.1568	24.895	1.2805
6.224	1.0880	16.408	1.1702	26.027	1.2966
7.455	1.0703	17.540	1.1839	27.158	1.3131
8.487	1.0819	18.671	1.1979	28.290	1.3300
9.619	1.0934	19.803	1.2122		

(Zimmermann, N. J. Pharm. 10. 2. 5.)

Sp. gr. of KOH + Aq.

% K ₂ O	Sp. gr.	% K ₂ O	Sp. gr.	% K ₂ O	Sp. gr.
2.44	1.02	23.14	1.22	37.97	1.43
4.77	1.04	24.77	1.24	40.17	1.44
7.02	1.06	26.34	1.26	42.31	1.46
9.20	1.08	27.86	1.28	44.40	1.48
11.28	1.10	29.34	1.30	46.44	1.50
13.30	1.12	30.74	1.32	48.46	1.52
15.38	1.14	32.14	1.34	50.09	1.54
17.40	1.16	33.46	1.36	51.58	1.56
19.34	1.18	34.74	1.38	53.06	1.58
21.25	1.20	35.99	1.40		

(Richter.)

Sp. gr. of KOH + Aq at 15°. a = sp. gr. if % is K₂O; b = sp. gr. if % is KOH.

%	a	b	%	a	b
1	1.010	1.009	31	1.370	1.309
2	1.020	1.017	32	1.385	1.311
3	1.030	1.025	33	1.403	1.324
4	1.039	1.033	34	1.418	1.336
5	1.048	1.041	35	1.431	1.349
6	1.058	1.049	36	1.445	1.361
7	1.068	1.058	37	1.460	1.374
8	1.078	1.065	38	1.475	1.387
9	1.089	1.074	39	1.490	1.400
10	1.099	1.083	40	1.504	1.411
11	1.110	1.092	41	1.522	1.425
12	1.121	1.110	42	1.539	1.438
13	1.132	1.111	43	1.564	1.450
14	1.143	1.119	44	1.570	1.462
15	1.154	1.128	45	1.584	1.472
16	1.166	1.137	46	1.600	1.484
17	1.178	1.146	47	1.615	1.496
18	1.190	1.155	48	1.630	1.511
19	1.202	1.166	49	1.645	1.527
20	1.215	1.177	50	1.660	1.539
21	1.230	1.188	51	1.676	1.552
22	1.242	1.198	52	1.690	1.565
23	1.256	1.209	53	1.705	1.577
24	1.270	1.220	54	1.720	1.590
25	1.285	1.230	55	1.733	1.604
26	1.300	1.241	56	1.746	1.618
27	1.312	1.252	57	1.762	1.630
28	1.326	1.264	58	1.780	1.641
29	1.340	1.278	59	1.795	1.655
30	1.355	1.289	60	1.810	1.667

(Calculated by Gerlach, Z. anal. 3. 27, after Zimmermann, N. J. Pharm. 10. 2. 5, and Schiff, A. 107. 300.)

Sp. gr. of KOH + Aq at 15°.

% KOH	Sp. gr.	% KOH	Sp. gr.
4.2	1.0382	21.0	1.2006
8.4	1.0776	22.4	1.2430
12.6	1.1177		1.2880
16.8	1.1588		

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of KOH + Aq at 15°.

KOH	Sp. gr.	% KOH	Sp. gr.
10	1.077	50	1.539
20	1.175	60	1.667
30	1.238	70	1.790
40	1.411

Wielach, Z. anal. 27. 275, calculated from Schiff, A. 107. 300.)

Sp. gr. of K₂O + Aq at 15°.

% K ₂ O	Sp. gr.	% K ₂ O	Sp. gr.
5	1.054	30	1.358
10	1.111	35	1.428
15	1.171	40	1.500
20	1.231	45	1.576
25	1.294

Hager, Adjumenta varia, Leipsic, 1876.)

Sp. gr. of KOH + Aq at 20° containing 2 la. KOH to 100 mols. H₂O = 1.05325. Col, Phil. Mag. (5) 16. 122.)

Sp. gr. of KOH + Aq at 15°.

H	Sp. gr.	% KOH	Sp. gr.	% KOH	Sp. gr.
1	1.53822	34	1.33313	16	1.14925
2	1.52822	33	1.32236	15	1.13955
3	1.51430	32	1.31166	14	1.12991
4	1.50245	31	1.30102	13	1.12031
5	1.49067	30	1.29046	12	1.11076
6	1.47896	29	1.27997	11	1.10127
7	1.46733	28	1.26954	10	1.09183
8	1.45577	27	1.25918	9	1.08240
9	1.44429	26	1.24888	8	1.07302
10	1.43289	25	1.23866	7	1.06371
11	1.42150	24	1.22849	6	1.05443
12	1.41025	23	1.21838	5	1.04517
13	1.39906	22	1.20834	4	1.03593
14	1.38793	21	1.19837	3	1.02671
15	1.37686	20	1.18839	2	1.01752
16	1.36586	19	1.17855	1	1.00834
17	1.35485	18	1.16875	0	0.99918
18	1.34396	17	1.15898		

Wickering, Phil. Mag. 1894, (5) 37. 375.)

Sp. gr. of N solution at 18°/4° = 1.0481.oomis, W. Ann 1896, 80. 55C.)

Sp. gr. of KOH + Aq.

% KOH	6.87	12.10
Sp. gr. 20°/20°	1.0601	1.1025

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 272.)

KOH + Aq containing equal pts of KOH and H₂O freezes at -54°. (Guyton-Morveau, Gm.-K. 2, 1. 18.)

KOH is completely miscible with NaOH and with RbOH in both the liquid and the solid states. (Hevesy, Z. phys. Ch. 1910, 73. 667.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 828.)

Abundantly sol in strong alcohol or wood-spirit.

See below under KOH + 2H₂O.

Readily sol in glycerine.

Sol in not less than 25 pts. of ether. (Boullay.) Sol. in much more than 25 pts. of ether. (Connell.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.; ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Insol. in acetone. Readily sol. in fusel oil.

Insol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Sol. in aqueous solution of mannite. (Favre, A. ch. (3) 11. 76.)

The composition of the hydrates formed by KOH at different dilutions is calculated from determinations of the lowering of the fr.-pt. produced by KOH and of the conductivity and sp. gr. of KOH + Aq. (Jones, Am. Ch. J. 1905, 34. 337.)

+H₂O.

+2H₂O. Very deliquescent, and sol. in H₂O with absorption of much heat

100 g. sat. solution in H₂O at 30° contain 55.75 g. anhyd. KOH. (de Waal, Dissert. 1910.)

Solubility of KOH + 2H₂O in alcohol + Aq at 30°.

% KOH	% alcohol	% H ₂ O
55.75	0	44.25
54.81	0.43	44.76
..*
31.0	57.50	11.50
28.99	65.07	5.94
27.67	69.92	2.41
27.20	73.01	negative
26.25	81.98	"

*Separates into two layers.

(de Waal, Dissert, 1910.)

+4H₂O.

Potassium hydrogen titanium diimide,
Ti(NH)NK.
Decomp. by H₂O and alcohol. Insol. in all ord. indifferent organic solvents. (Ruff, B. 1912, 45. 1371.)

Potassium iodide, KI.
Deliquescent only in very moist air. Very sol. in H₂O with absorption of heat.
The temp. of H₂O can be lowered 24° by dissolving KI. (Baup.)
140 pts. KI dissolved in 100 pts. H₂O at 10.8° lower the temp. 22.5°. (Rüdorff, Pogg. 136. 276.)
100 pts. H₂O dissolve 126.6 pts. KI at 0° (Kremers); 127.8 pts. KI at 0° (Mulder); 127.9 pts. KI at 0°. (Gerardin.)
By boiling, 100 pts. H₂O dissolve 221 pts. KI at 120° (Baup); 222.2 pts. KI at 120° (Gay-Lussac); 222.6 pts. KI at 118.4° (Mulder); 223.58 pts. KI at 117° (Legrand); 223.6 pts. KI at 117°. (Gerardin.)
Between these temps. the solubility increases proportional to temp.

Sol. in 0.735 pt. H₂O at 12.5°; in 0.709 pt. H₂O at 16°; in 0.7 pt. H₂O at 18°; in 0.45 pt. H₂O at 120°. (Graham-Otto.)
100 pts. KI + Aq sat. at 15-16° contain 58.07 pts. KI. (v. Hauer, J. pr. 98, 137.)
100 pts. H₂O at 12.5° dissolve 136 pts.; at 16°, 141 pts. KI. (Baup.)
100 pts. H₂O at 18° dissolve 143 pts. KI; at 120°, 271 pts. (Gay-Lussac.)
Sol. in 0.79 pt. H₂O at 0°; in 0.70 pt. H₂O at 20°; in 0.63 pt. H₂O at 48°; in 0.57 pt. H₂O at 60°; in 0.53 pt. H₂O at 80°; in 0.51 pt. H₂O at 100°. (Kremers, Pogg. 97. 15.)

Sol. in 0.71 pt. H₂O at 15°. (Eder, Dingl. 221. 89.)

Solubility of KI in 100 pts. H ₂ O at t°.					
t°	Pts. KI	t°	Pts. KI	t°	Pts. KI
0	127.9	19	143.4	38	159
1	128.7	20	144.2	39	160
2	129.6	21	145.1	40	160
3	130.4	22	145.9	41	161
4	131.2	23	146.7	42	162
5	132.1	24	147.5	43	163
6	132.9	25	148.3	44	164
7	133.7	26	149.1	45	164
8	134.5	27	149.9	46	165
9	135.3	28	150.7	47	166
10	136.1	29	151.5	48	167
11	137.0	30	152.3	49	168
12	137.8	31	153	50	168
13	138.6	32	154	51	169
14	139.4	33	155	52	170
15	140.2	34	156	53	171
16	141.0	35	156	54	172
17	141.8	36	157	55	172
18	142.6	37	158	56	173

Solubility of KI in 100 pts., etc.—Cont.					
t°	Pts. KI	t°	Pts. KI	t°	P
57	174	78	191	99	
58	175	79	192	100	
59	175	80	192	101	
60	176	81	193	102	
61	177	82	194	103	
62	178	83	195	104	
63	179	84	196	105	
64	180	85	197	106	
65	180	86	197	107	
66	181	87	198	108	
67	182	88	199	109	
68	183	89	200	110	
69	184	90	201	111	
70	184	91	202	112	
71	185	92	202	113	
72	186	93	203	114	
73	187	94	204	115	
74	188	95	205	116	
75	188	96	206	117	
76	189	97	207	...	
77	190	98	208	...	

(Mulder, calculated from his own and observations, Scheik. Verhandel. 1864.

Solubility of KI in 100 pts. H ₂ O at				
t°	Pts. KI	t°	Pts. KI	t°
-22.65	107.2	21.05	143.3	71.1
-22.35	106.6	25.6	146.6	74.75
-16.8	111.1	29.1	149.6	81.6
-11.35	116.3	37.3	156.7	86.35
-5.9	120.4	42.3	160.3	93.5
0	126.1	45.75	163.6	100.7
+3.25	130.1	51.8	167.6	110.2
9.55	134.0	55.05	169.1	113.7
12.75	137.1	60.55	173.4	...
12.9	137.9	65.0	178.3	...

(Coppet, A. ch. (5) 80. 417.)

Solubility is represented by a straight of the formula 126.23+0.8088t. (Copp

Solubility of KI in 100 pts. H ₂ O at high			
t°	Pts. KI	t°	Pts.
124	233.9	144	264
133	249.3	175	311

(Tilden and Shenstone, Phil. Trans. 1881

If solubility S = pts. KI in 100 pts. sol
S = 55.8+0.122t from 0° to 165°. (C
C. R. 98. 1432.)

Sat. KI + Aq contains % KI at t°.

t°	% KI	t°	% KI
21	50.7	78	64.8
21	51.0	96	66.9
21	51.2	150	70.6
19	52.2	151	70.9
15	53.2	175	71.6
-9	54.5	176	72.7
0	56.9	190	73.8
21	59.3	193	74.5
44	60.8	213	75.7
72	64.3

(Étard, A. ch. 1894, (7) 2, 542.)

Solubility of KI in 100 g. H₂O at t°.

t°	g. KI	t°	g. KI
1	122.2	-11.5	64.7
-5	119.8	-9.5	51.5
-4	117.4	-7	42.6
10	115.1	-6	34.4
-4	75.8	-5	25.7

(Meusser, Z. anorg. 1905, 44, 80.)

70 pts. by weight are contained in 100 g. of sat. aq. at 25°, or 59.54 pts. in 100 g. of aq. at 0°; sp. gr. = 1.7254.

15 pts. by weight are contained in 100 cc. of aq. sat. at 0°, or 56.34 pts. in 100 g. of aq. at 0°; sp. gr. = 1.6699. (Walden, Z. phys. Chem., 65, 715.)

Solubility of KI in H₂O at low temperatures.

% KI	Solid phase	t°	% KI	Solid phase
53.8	Ice	-22	52.1	KI
41.2	"	-20	52.6	"
54.6	"	-15	53.5	"
48	"	-10	54.5	"
55.2	"	-5	55.4	"
55.9	" + KI	0	56.4	"

(Mann and Kerschbaum, Z. anorg. 1907, 64, 218.)

26 g. KI dissolve in 100 g. H₂O at 25° (dori and Pampanini, Rend. Acc. Linc. V, 20, 473.)

19 g. in 100 g. KI + Aq sat. at 25° (Mann and Whittemore, J. Am. Chem. Soc. 53, 1934.)

1 g. in 100 g. KI + Aq sat. at 0°, 60.35 g. KI + Aq sat. at 30°. (Van Dam and Monk, Chem. Weekbl. 1911, 8, 848.)

Sp. gr. of KI + Aq at 21°.

% KI	Sp. gr.	% KI	Sp. gr.	% KI	Sp. gr.
1	1.0075	21	1.1807	41	1.4224
2	1.0151	22	1.1911	42	1.4371
3	1.0227	23	1.2016	43	1.4520
4	1.0305	24	1.2122	44	1.4671
5	1.0384	25	1.2229	45	1.4825
6	1.0464	26	1.2336	46	1.4982
7	1.0545	27	1.2445	47	1.5142
8	1.0627	28	1.2556	48	1.5305
9	1.0710	29	1.2669	49	1.5471
10	1.0793	30	1.2784	50	1.5640
11	1.0877	31	1.2899	51	1.5810
12	1.0962	32	1.3017	52	1.5984
13	1.1048	33	1.3138	53	1.6162
14	1.1136	34	1.3262	54	1.6343
15	1.1226	35	1.3389	55	1.6528
16	1.1318	36	1.3519	56	1.6717
17	1.1412	37	1.3653	57	1.6911
18	1.1508	38	1.3791	58	1.7109
19	1.1605	39	1.3933	59	1.7311
20	1.1705	40	1.4079	60	1.7517

(Schiff, A. 110, 75.)

Sp. gr. of KI + Aq. S = according to Schiff (A. 108, 340) at 21°; K = according to Kremers (Pogg. 96, 62), interpolated by Gerlach (Z. anal. 8, 285.)

	5	10	15	20	25	30% KI,
S	1.038	1.079	1.123	1.171	...	1.279
K	1.038	1.078	1.120	1.166	1.218	1.271

	35	40	45	50	55	60% KI,
S	1.483
K	1.331	1.396	1.469	1.546	1.636	1.734

Sp. gr. of KI + Aq at 18°.

% KI	Sp. gr.	% KI	Sp. gr.	% KI	Sp. gr.
10	1.0363	30	1.273	55	1.630
10	1.0762	40	1.3966		
20	1.1679	50	1.545		

(Kohlrausch, W. Ann. 1879, 1.)

Sp. gr. of KI + Aq at 18°.

% KI	Sp. gr.
1.044	1.0062
5.0	1.0363

(Giotrian, W. Ann. 1883, 18, 191.)

Sp. gr. at 16°/4° of KI + Aq containing 32.4875% KI = 1.30238. (Schönrock, Z. phys. Ch. 1893, 11, 781.)

KI + Aq containing 9.35% KI has sp. gr. 20°/20° = 1.0726

KI + Aq containing 11.35% KI has sp. gr. 20°/20° = 1.0892. (Le Blanc and Rohland, Z. phys. Ch. 1896, 19, 278.)

B.-pt. of KI+Aq containing pts. KI to 100 pts. H₂O.

B.-pt.	Pts. KI	B.-pt.	Pts. KI	B.-pt.	Pts. KI
101°	15	108°	111.5	115	185°
102	30	109	123	116	195
103	45	110	134	117	205
104	60	111	145	118	215
105	74	112	155	118.5	220
106	87	113	165
107	99.5	114	175

(Gerlach, Z. anal. 26. 439.)

Sat. KI+Aq boils at 119°. (Kremers.)

Sat. KI+Aq forms a crust at 117.5°, and contains 210 pts. KI to 100 pts. H₂O; highest temp. observed, 118.5°. (Gerlach, Z. anal. 26. 426.)

Solubility of KI in I₂+Aq at 25°.

KI mol./l.	I g.-atoms/l.
6.15	0.00
6.23	3.64
6.40	11.11
6.36	13.16
6.33	13.2
6.24	17.03

(Abegg, Z. anorg. 1906, 50. 428.)

Solubility of KI+I₂ in H₂O at 25°.

% KI	% I	Solid phase	% KI	% I	Solid phase
29.45	64.34	KI+KI ₂	25.88	68.79	KI ₂ +I ₂
28.91	63.88	"	25.57	69.01	"
26.84	66.54	KI ₂ +KI ₂	27.86	66.56	KI ₂
27.18	67.14	"	27.27	66.91	"
27.14	66.60	"	26.95	67.17	KI ₂
			25.71	67.91	"

(Foote and Chalker, Am. Ch. J. 1908, 39. 564.)

See also under Iodine.

KI+Aq sat. at 14.5° containing 139.8 pts. KI to 100 pts. H₂O dissolves 1.0 pt. K₂SO₄ with separation of 2.2 pts. KI, so that solution contains 137.6 pts. KI and 1.0 pt. K₂SO₄ to 100 pts. H₂O. (Mulder, Rotterdam, 1864.)

100 pts. H₂O dissolve 86.3 pts. KI and 2.1 pts. Na₂SO₄ at 14.5°. (Mulder, J. B. 1866. 67.)

Sol. in AsCl₃, SnCl₄ and POCl₃. (Walden, Z. anorg. 1900, 25. 214.)

Attacked by dry liquid NO₂ with liberation of I₂. (Frankland, Chem. Soc. 1901, 79. 1361.)

Sol. in liquid SO₂. (Walden, B. 1 2864.)

Solubility in SO₂ decreases with temp. (Walden, Z. phys. Ch. 1903, Insol. in liquid CO₂. (Büchner, Ch. 1906, 54. 674.)

Very easily sol. in liquid NH₃. (Am. Ch. J. 1898, 20. 829.)

Hydrazine dissolves 135.7 pts. KI 13°. (de Bruyn, R. t. c. 1899, 18.)

100 pts. alcohol of 0.85 sp. gr. dissolve at 12.5°. 100 pts. absolute alcohol dissolve KI at 13.5°. Much more sol. in hot alcohol.

100 pts. alcohol of D sp. gr. at 18°—

D 0.9904 0.9851 0.9726 0.9665 0.9513
130.5 119.4 100.1 89.9 76

D 0.9390 0.9088 0.8464 0.8321
66.4 48.2 11.4 6.2

That is, aqueous alcohol dissolves the same amount of KI that is present in the alcohol would dissolve therefore probable that KI is insoluble in absolute alcohol. (Gerardin.)

Solubility in 100 pts. alcohol of gr. at:

8° 13° 25° 46° 55° 67°
67.4 69.2 75.1 84.7 87.5 90

(Gerardin, A. ch. (4) 5. 1.)

Sol. in 68.3 pts. absolute alcohol (Dingl. 221. 89); in 370 pts. ethyl alcohol (0.729), (Eder, l. c.); in 120 pts. alcohol (1:1), (Eder, l. c.)

Sol. in 10-12 pts. 90% alcohol, absolute alcohol. (Hager, Comm.)

100 pts. absolute methyl alcohol dissolve 16.5 pts. at 20.5°; 100 pts. absolute alcohol dissolve 1.75 pts. at 20.5°. (Z. phys. Ch. 10. 783.)

Solubility of KI in methyl alcohol

P = % by wt. of alcohol in alcohol
S = Sp. gr. of alcohol+Aq sat.
L = millimols KI in 100 ccm. solution.

P	S 25°/4°
0	1.7213
10.6	1.634
30.8	1.460
47.1	1.325
64.0	1.185
78.1	1.066
98.9	0.9700
100	0.9018

(Hertz and Anders, Z. anorg. 1904, 104)

Solubility of KI in CH_3OH .

. KI in 100 g. of the solution.
temp. of complete solution.
temp. at which salt begins to separate

	t_1	t_2
14	...	266°
15	0°	..
	20	262
17	25	256
	85	242
	115	229
	144	196
	188	...

ressner, Z. phys. Ch. 1910, 72. 432.)

Solubility of KI in methyl alcohol at t° .

g. KI in 100 g. alcohol	t°	g. KI in 100 g. alcohol
14.50	180	30.7
16.20	200	29.1
18.9	220	27.5
22.5	240	24.8
25.0	245	22.6
27.2	247	21.0
29.2	250	13.8
30.6	252.5°	7.6

sol temp. of solution

ressner, Chem. Soc. 1910, 97. 626.)

at temp. 1 pt. KI by weight is sol. in
1. methyl alcohol D^{16} 0.7990.
ethyl " D^{16} 0.8322.
propyl " D^{16} 0.8160.

and, Z. anorg. 1898, 18. 325.)

in mixtures of methyl and ethyl alcohol at 25°.

methyl alcohol in the solvent
KI in 10 ccm. of the solution.
gr. of the sat. solution at 25°.

	G	S 25°/4°
10	0.155	0.8015
17	0.191	0.8041
20	0.225	0.8071
22	0.494	0.8295
29	1.013	0.8794
7	1.072	0.8795
5	1.184	0.8908
0	1.316	0.9018

1 Kuhn, Z. anorg. 1908, 60. 155.)

Solubility in mixtures of methyl and propyl alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. KI in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	1.316	0.9018
11.11	1.096	0.8823
23.8	0.854	0.8629
65.2	0.262	0.8187
91.8	0.060	0.8045
93.75	0.058	0.8041
100	0.043	0.8041

(Hers and Kuhn.)

Solubility in mixtures of propyl and ethyl alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. KI in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution

P	G	S 25°/4°
0	0.155	0.8015
8.1	0.146	0.7983
17.85	0.137	0.7991
58.6	0.075	0.7988
88.6	0.052	0.8022
91.2	0.049	0.8027
95.2	0.044	0.8029
100	0.043	0.8041

(Hers and Kuhn.)

100 g. methyl alcohol dissolve 18.04 g. KI at 25°.

100 g. ethyl alcohol dissolve 2.16 g. KI at 25°.

100 g. propyl alcohol dissolve 0.43 g. KI at 25°.

100 g. isopropyl alcohol dissolve 0.09 g. KI at 25°.

(Turner and Bissett, Chem. Soc. 1913, 103. 1909.)

0.455 g. is sol. in 100 g. propyl alcohol. (Schlamp, Z. phys. Ch. 1894, 14. 276.)

Alcoholic solution can be mixed with $\frac{1}{2}$ vol. ether without pptn.

100 g. 95% formic acid dissolve 38.2 g. KI at 18.5°. (Aschan, Chem. Ztg. 1913, 37. 1113.)

Solubility in organic solvents at t° .

C = pts. by wt. of KI in 100 ccm. of the sat. solution.

L = no. of litres which at the saturation temp. hold in solution 1 mol. KI.

S = sp. gr. of the solution at t° , referred to H_2O at t° .

p = pts. by wt. of KI in 100 g. of the solution.

Solvent	t°	C	L	S	p
Water	25°	102.70	0.162	1.7254	59.54
	0°	94.05	0.177	1.6699	56.32
Methyl alcohol	25°	13.48	1.231	0.9003	14.97
"	25°	14.26			
	0°	11.61	1.430	0.8964	12.95
Ethyl alcohol	25°	1.520	10.92	0.7908	1.922
"	0°	1.197	13.87	0.8085	1.479
Glycol	25°	45.85	0.362	1.3888	33.01
"	25°	47.23	0.351		
	0°	43.28	0.383	1.3954	31.03
Acetonitrile	25°	1.551	10.70		
"	25°	1.590	10.44	0.7936	2.003
"	0°	1.852	9.00	0.8198	2.259
Propionitrile	25°	0.316	52.53	0.7821	0.404
"	25°	0.355	46.76		
"	0°	0.344	48.26	0.8005	0.429
"	0°	0.412	40.29		
Benzonitrile	25°	0.051	325.5	1.0076	0.050
Nitromethane	25°	0.349	47.56	1.1367	0.307
"	25°	0.289	57.44		
"	0°	0.366	45.36	1.1627	0.315
"	0°	0.314	52.87		
Nitrobenzene	25°	0.0019	87.40		
Acetone	25°	1.038	16.0	0.7968	1.302
"	0°	1.732	9.58	0.8227	2.105
Furfural	25°	5.93	2.80	1.2014	4.94
"	0°	15.10	1.10		
Benzaldehyde	25°	0.343	48.4	1.0446	0.328
Salicylaldehyde	25°	0.549	30.24	1.1373	0.483
"	0°	1.257	13.21	1.1501	1.093
Anisaldehyde	25°	0.720	23.06	1.1180	0.644
"	0°	1.520	10.92	1.1223	1.355
Ethyl acetate	25°	0.0013	12.80		
Methyl cyanacetate	25°	2.459	6.75	1.1358	2.165
"	0°	3.256	5.10	1.1521	2.827
Ethyl cyanacetate	25°	0.888	18.7	1.0579	0.839
"	25°	1.090	15.23	1.0678	1.021

(Walden, Z. phys. Ch. 1906, 55. 715.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3789.)

Sol. in ethyl acetate. (Casasaca, C. R. 30. 821.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in ethylamine. (Shinn, J. phys. Chem. 1907, 11. 538.)

100 pts. acetone dissolve 2.930 pts. KI at 25°. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

Sol. in acetone, insol. in methylal. (Eidmann, C. C. 1899. II. 1014.)

3.08 pts. sol. in 100 pts. acetone at -2.5° .

2.38 " " " 100 " " " $+22^{\circ}$.

1.21 " " " 100 " " " 56° .

0.26 " " " 100 " pyridine " 10° .

0.11 " " " 100 " " " 119° .

(Luszczyński, B. 1894, 27. 2287.)

Freely sol. in glyc. rine. Insol. in acetic acid. (Berthelot.)

Sol. in 3 pts. glycerine; insol. in olive oil. (Cap and Garot.)

100 g. glycerol dissolve 40 g. KI at 18°. (Ossendowski, Pharm. J. 1907, 79. 575.)

Potassium triiodide, KI_3 .

Very deliquescent; very sol. in H_2O and alcohol. (Johnson, Chem. Soc. 1877, 1. 249.)

Solution of I in KI contains this salt (as KI). Decomp. by heat or shaking with CH_2Cl_2 , ether, chloroform. Sol. in alcohol, from which CS_2 does not remove I. (Jørgensen, I. pr. (2) 2. 247.)

Potassium periodide.

Solubility determinations show that the compds. KI_5 and KI_7 are the only periodides of potassium which form solids at 25°.

See under $KI+I$. (Foote and Chappin, Am. Ch. J. 1908, 39. 566.)

KI_7 . See above.

Potassium mercuric iodide ammonia, $K_2HgI_4, 2NH_3$.

(Peters, Z. anorg. 1912, 77. 188.)

Potassium silver iodide, KI, AgI .

Sol. in $KI+Aq$. Sol. in hot alcohol. (Boullay, A. ch. 34. 377.)

$2KI, AgI$. Sol. in $KI+Aq$. Decomp. by H_2O (Boullay.)

Hygroscopic. (Hellwig, Z. anorg. 1902, 21. 180.)

$3KI, AgI$. Decomp. by H_2O . (Ditte, C. R. 93, 415.)

$KI, 2AgI$. Sol. in methylethylketone. (Marsh, Chem. Soc. 1913, 103. 783.)

Potassium silver polyiodide, $AgK_2I_{12}, 3KI+5H_2O$.

Very deliquescent. (Johnson, Chem. Soc. 33. 183.)

Potassium tellurium iodide.

See Iodotellurate, potassium.

Potassium thallic iodide, KI, TlI_3 .

Decomp. by H_2O . Can be crystallised from alcohol. (Willm.)

$3KI, 2TlI_3+3H_2O$. Partially decomp. by H_2O . (Rammelsberg.)

Potassium (tin) stannous iodide, $KI, SnI_4+1\frac{1}{2}H_2O$.

When treated with a small quantity of H_2O , KI dissolves out; but when more H_2O is added, the substance is completely dissolved. More sol. in warm than cold alcohol. (Boullay.)

am zinc iodide, KI, ZnI₂.
deliquescent. (Rammelsberg, Pogg.
[₄+2H₂O. Hydrosopic. (Ephraim,
1910, 67. 382.)

am iodide sulphur dioxide, KI, SO₂.
ard, C. R. 1900, 130. 1188.)
SO₂. (Walden, Z. phys. Ch. 1903,
150. (Walden.)

am nitride, K₂N.
ap. violently by H₂O. (H. Davy.)

am ruthenium dihydronitrosobromide,
H₂, NOBr₂, 2HBr, 3KBr.
(Brizard, A. ch. 1900, (7) 21. 362.)

am ruthenium nitroschloride,
H₂NOCl₂, 3KCl, 2HCl.
in H₂O. (Brizard, C. R. 1899, 129.

am suboxide.
aposes H₂O.
not exist. (Lupton, Chem. Soc. 1876,

am oxide, K₂O.
sol. in H₂O with much heat.
potassium hydroxide.

am dioxide, K₂O₂.
rescent. Sol. in H₂O.
compound K₂O₂, 2H₂O₂. (Schöne,
241.)

am peroxide, K₂O₄.
rescent. Very sol. with decomp. in

am silicon oxyfluoride, SiF₂(OK)₂ and
(F)OK.
and Bechi, A. Suppl. 4. 33.)

am tantalum oxyfluoride, K₄Ta₄O₆F₁₄.
in boiling water. Easily sol. in HF +
arignac, A. ch. (4) 9. 268.)

am phosphide, KP₃.
ap. by H₂O. (Joannis, C. C. 1894,

Easily decomp. by H₂O. (Hugot,
95, 121. 208.)

am hydrogen phosphide, PH₂K.
ap. by H₂O. (Joannis, C. R. 1894,
)

am phosphoselenide, KSeP = K₂Se,
3.

cold H₂O with rapid decomp. Sol.
l with slight decomp. (Hahn, J. pr.

Potassium phosphotriselenide, 2K₂Se,
P₂Se₃.

Deliquescent. Decomp. violently with H₂O.
Sol. in alcohol or ether, or in a mixture of the
two, with slight decomp., but decomp. gradu-
ally on the air. (Hahn, J. pr. 93. 430.)

Potassium phosphopentaseleenide, K₄P₂Se₇ =
2K₂Se, P₂Se₃.

Deliquescent; immediately decomp. by
H₂O, alcohol, or ether. (Hahn.)

Potassium phosphosulphide, 4K₂S₂, P₂S₃.

Deliquescent. Sol. in H₂O with decomp.

Potassium selenide, K₂Se.

Sol. in H₂O with subsequent decomp. on the
air.

Insol. in liquid NH₃; sol. in air free H₂O
to a colorless liquid. (Hugot, C. R. 1899,
129. 299.)

+2H₂O. Sol. in H₂O with decomp.
(Clever, Z. anorg. 1895, 10. 143.)

+9, 14, or 19H₂O. (Fabre, C. R. 102.
613.)

Potassium tetraselenide, K₂Se₄.

Easily sol. in H₂O. Decomp. on standing.
Sol. in liquid NH₃. (Hugot, C. R. 1899,
129. 299.)

Potassium monosulphide, K₂S.

Deliquescent. Sol. in H₂O and alcohol.
H₂O solution decomp. on air.

Sol. in 10 pts. glycerine. (Cap and Garot,
J. Pharm. (3) 26. 81.)

Moderately sol. in liquid NH₃. (Franklin,
Am. Ch. J. 1898, 20. 829.)

Insol. in acetone and in methylal. (Eid-
mann, C. C. 1899, II. 1014.)

Insol. in methyl acetate. (Naumann, B.
1909, 42. 3790.)

+5H₂O. (Schöne, Pogg. 131. 380.)

All potassium sulphides are sol. in glycerine;
insol. in ether and ethyl acetate.

Potassium disulphide, K₂S₂.

Sol. in H₂O and alcohol, with gradual de-
comp.

Potassium trisulphide, K₂S₃.

Sol. in H₂O and alcohol, with gradual de-
comp. on the air.

Potassium tetrasulphide, K₂S₄.

Sol. in H₂O and alcohol.

+2H₂O. Sol. in H₂O. Sl. sol. in alcohol.

+8H₂O. Sol. in H₂O. Alcohol takes out
water. (Schöne.)

Potassium pentasulphide, K₂S₅.

Sol. in H₂O and alcohol.

Potassium palladium sulphide.

See Sulphopalladate, potassium.

Potassium platinum sulphide.

See Sulphoplatinate, potassium.

Potassium silver sulphide,

Decomp. by H_2O . (Ditte, C. R. 1895, 120. 91.)

Potassium rhodium sulphide, $3\text{K}_2\text{S}, \text{Rh}_2\text{S}_3$.

Decomp. by H_2O . (Leidié.)

Potassium tellurium sulphide.

See Sulphotellurate, potassium.

Potassium thallium sulphide, $\text{K}_2\text{S}, \text{Tl}_2\text{S}_3$.

Not decomposed by H_2O , or hot NH_4OH , or $\text{KOH} + \text{Aq}$. Decomp. by HCl or moderately conc. $\text{H}_2\text{SO}_4 + \text{Aq}$. Hot $\text{HNO}_3 + \text{Aq}$ decomp. with separation of S. (Schneider, J. pr. 110. 168.)

Potassium tin (stannic) sulphide.

See Sulphostannate, potassium.

Potassium zinc sulphide, $\text{K}_2\text{S}, 3\text{ZnS}$.

Not attacked by H_2O , but easily decomp. by the most dil. acids. (Schneider, J. pr. (2) 8. 29.)

Potassium telluride, K_2Te .

Sol. in H_2O . (Demarçay, Bull. Soc. (2) 40. 99.)

Sol. in H_2O and liquid NH_3 . (Hugot, C. R. 1899, 129. 388.)

Praseocobaltic chloride,

Easily sol. in H_2O .

Dil. $\text{HCl} + \text{Aq}$ dissolves traces; conc. $\text{HCl} + \text{Aq}$ dissolves more. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ with decomp. Sol. in conc. H_2SO_4 without decomp. Sl. sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Rose.)

— mercuric chloride, $\text{Co}(\text{NH}_3)_4\text{Cl}_2, \text{HgCl}_2$.

Sl. sol. in cold H_2O ; insol. in $\text{HgCl}_2 + \text{Aq}$. (Vortmann, B. 15. 1892.)

— chloride dichromate,

Scarcely sol. in cold, easily sol. in warm H_2O . (Vortmann, B. 15. 1897.)

Praseocobaltic chloride nitrate,

Much less sol. in H_2O than the chloride. Precipitated from aqueous solution by dil. $\text{HNO}_3 + \text{Aq}$. (Vortmann, B. 15. 1896.)

Praseodymicotungstic acid.**Ammonium praseodymicotungstate,**

Very sl. sol. in H_2O . Decomp. by and alkalies. (E. F. Smith, J. Am. Soc. 1904, 26. 1478.)

Barium praseodymicotungstate, $4\text{BaO}, 16\text{WO}_3 + 7\text{H}_2\text{O}$.

Ppt. Insol. in H_2O .

$6\text{BaO}, \text{Pr}_2\text{O}_3, 16\text{WO}_3 + 9\text{H}_2\text{O}$. Ppt. Smith.)

Silver praseodymicotungstate, $4\text{Ag}_2\text{O}, 16\text{WO}_3 + 8\text{H}_2\text{O}$.

Insol. in H_2O . (E. F. Smith.)

Praseodymium, Pr.**Praseodymium bromide, $\text{PrBr}_3 + 6\text{H}_2\text{O}$.**

Very sol. in H_2O ; sol. in HBr . (von S. Z. anorg. 1898, 18. 353.)

Praseodymium carbide, PrC_2 .

Decomp. by H_2O ; insol. in conc. H decomp. by dil. HNO_3 . (Moissan, 1900, 131. 597.)

Praseodymium chloride, PrCl_3 .

Very sol. in H_2O . Insol. in PCl_5 or S . Sol. in alcohol. Insol. in ether and organic solvents. (Matignon, C. R. 1902 427.)

2.14 g. PrCl_3 dissolve in 100 g. pyrid 15° . (Matignon, Int. Cong. App. Chem 2. 53.)

$+ \text{H}_2\text{O}$.

$+ 3\text{H}_2\text{O}$.

$+ 7\text{H}_2\text{O}$. 100 g. H_2O dissolve 33

$\text{PrCl}_3 + 7\text{H}_2\text{O}$ or 103.9 g. of the anhyd salt at 13° . The aqueous solution at 14° has a sp. gr. $16^\circ/16^\circ = 1.687$. At 10 solubility in H_2O is unlimited. (Mat A. ch. 1906, (8) 8. 388.)

Sol. in conc. HCl . (von Schule, Z. 1898, 18. 352.)

100 pts. of a solution of the salt in HCl contain at 13° , 41.05 pts. of anhydrous and 7.25 pts. HCl . Sp. gr. of this soln $16^\circ = 1.574$. (Matignon, A. ch. 1906, 388.)

Praseodymium hydride, PrH_3 (?).

(Muthmann, A. 1904, 331. 59.)

Praseodymium hydroxide.

Sol. in citric acid. (Baskerville, J. Chem. Soc. 1904, 26. 49.)

Praseodymium nitride, PrN .

Decomp. in moist air with evolution of NH_3 . (Muthmann, A. 1904, 331. 59.)

um oxide, Pr_2O_3 .

l. in H_2O . (v. Welsbach, M. 8.

by heating in the air. (Scheele, 1798, 17. 322.)

um monoperoxide, $\text{Pr}(\text{OH})_2\cdot\text{H}_2\text{O}$.
, Chem. Soc. 1902, 82. (2) 140.)

um superoxide, $\text{Pr}(\text{OH})_2(\text{OOH})$.
elickoff, C. C. 1902, I. 172.)

um trisuperoxide, $\text{Pr}(\text{OOH})_3$.
elickoff.)

um peroxide, Pr_4O_7 .
acids with evolution of O. (v.

um oxysulphide, Pr_2SO_2 .
anorg. 1911, 71. 436.)

um disulphide, PrS_2 .
by heat. (Biltz, Z. anorg. 1911,

obaltic salts.

· purpureocobaltic salts, see—

rpureocobaltic salts.

rpureocobaltic salts.

urpureocobaltic salts.

purpureocobaltic salts.

baltic cobalticyanide,
 $[\text{I}_3]_2\text{Co}(\text{CN})_6 + 1\frac{1}{2}\text{H}_2\text{O}$.
 H_2O .

anide, $\text{Co}(\text{NH}_3)_4\text{Fe}(\text{CN})_6$.
cold H_2O . Probably belongs to

ric hydroxychloride,
 $[\text{I}_{11}(\text{HgCl})_2(\text{HgOH})\text{Cl}_2$.
ortmann and Morgulis, B. 22.

$(\text{HgOH})_4\text{Cl}_2$. Ppt. (V. and M.)

riodide, basic,
 $[\text{I}_{11}(\text{HgI}_2)_2(\text{HgOH})_2\text{I}_2$.
sol. in acids. Sol. in $\text{KI} + \text{Aq}$.
and Borsbach, B. 23. 2804.)

date, $\text{Co}_2\text{O}_3(\text{NH}_3)_{10}$, $7\text{MoO}_3 +$
(?).

H_2O or dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Car-
109. 109.)

ite.

atopurpureocobaltic salts.

ate, $\text{Co}(\text{NH}_3)_4\text{O}(\text{WO}_4)$.
sol. in cold or hot H_2O . (Gibbs.)
 $[\text{H}_3]_{10}$, $10\text{WO}_3 + 9\text{H}_2\text{O}$ (?). Insol.

in H_2O , or dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, or $\text{NH}_4\text{OH} +$
 Aq . (Carnot, C. R. 109. 147.)

Purpureocobaltic vanadate, $\text{Co}_2\text{O}_3(\text{NH}_3)_{10}$,
 $5\text{V}_2\text{O}_5 + 9\text{H}_2\text{O}$ (?).

Ppt. Insol. in H_2O . (Carnot, C. R. 109.
147.)

Purpureocobaltic octamine salts.

See Octamine cobaltic purpureo salts.

Pyrosulphuric acid, $\text{H}_2\text{S}_2\text{O}_7$.

See Disulphuric acid.

Radium, Ra.

Radium A.

More sol than Radium B and C in all
solvents; sol. even in organic solvents, espe-
cially CS_2 . (Ramstedt, Le Radium, 1913, 10.
159.)

Radium B.

More quickly sol. than Radium C in H_2O
and acids; less quickly sol. in alkaline solu-
tions; very sl. sol. in organic solvents. (Ram-
stedt, Le Radium, 1913, 10. 159.)

Radium C.

Sol. in common acids, less so in alkaline
solutions and in H_2O , only very sl. sol. in
organic solvents. (Ramstedt, Le Radium,
1913, 10. 159; Chem. Soc. 1913, 104. (2) 659.)

Radium bromide.

Less sol. in H_2O than corresponding Ba
comp. (Curie, Dissert. 1903.)

Radium chloride.

Less sol. in H_2O than corresponding Ba
comp. (Curie, Dissert. 1903.)

Radium emanation.

Coefficient of absorption for $\text{H}_2\text{O} = 0.245$ at
 3° ; 0.23 at 20° ; 0.17 at 40° ; 0.135 at 60° ; 0.12
at 70° ; 0.12 at 80° . (Hofmann, Phys. Zeit.
1905, 6. 339.)

Solubility in H_2O at t° .

Coefficient of solubility = conc. of the
emanation in the liquid: conc. of the emana-
tion in the gas.

t°	Coefficient of solubility
0	0.506
4.3	0.424
5.7	0.398
10.0	0.340
14.0	0.303
17.6	0.280
20.0	0.245
26.8	0.206
31.6	0.193
39.1	0.160

Coefficient of solubility in sea-water of
sp. gr. at $14^\circ = 1.022$ is 0.255.

(Boyle, Phil. Mag. 1911, (6) 22. 850.)

Solubility in H_2O .

Temp.	0.5°	17.5°	35°	41°	51°
Sol.	0.526	0.283	0.183	0.161	0.138

Temp.	60°	74°	79°	82°	91°
Sol.	0.127	0.112	0.111	0.111	0.108

(Kofler, M. 1913, 34. 389.)

Coefficient of solubility of radium emanation at 14° in various solvents.

Ethyl alcohol	7.34
Amyl alcohol	9.31
Toluene	13.7
Sea water	0.255
Mercury	0

(Boyle, Phil. Mag. 1911, (6) 22. 851.)

Coefficient of solubility emanation of radium in various solvents at t° .

Solvents	$t = 18^\circ$	$t = 0^\circ$	$t = -18^\circ$
Ethyl acetate	7.35	9.41	13.6
Acetone	6.30	7.99	10.8
Absolute alcohol	6.17	8.28	11.4
Aniline	3.80	4.43	...
Benzene	12.82	16.54 at 3°	...
Chloroform	15.08	20.5	28.5
Cyclohexane	18.04 at 80°
Water	0.285	0.52	...
Ether	15.08	20.09	29.1
Glycerine	0.21
Hexane	16.56	23.4	35.2
Paraffine oil	9.2	12.6	...
Carbon-bisulphide	23.14	33.4	50.3
Toluene	13.24	18.4	27
Xylene	12.75

(Ramstedt, Le Radium, 1911, 8. 255.)

Solubility in various oils, etc., at t° .

Rape oil		Poppy seed oil		Oil of turpentine	
t°	Solubility	t°	Solubility	t°	Solubility
-3	51.2	-5	50.5	-21	42.5
10	35.3	16	30.2	0	23.1
20	26.1	40	19.1	18	16.6
100	6.2	65	12.4	50	7.5
200	3.3	90	8.4	6.5	4.08

Solubility in 10% dammar resin in oil of turpentine = 16.7 at 18°.

Solubility in 5% colophony in amyl alcohol = 11.2 at 20°.

Solubility in amyl alcohol = 10.6 at 18°.

Solubility in 20% colophony in amyl alcohol = 11.1 at 20°.

(Curie, Thesis. 1910.)

Coefficient of absorption for petrole

22.70 at -21° .12.87 at $+3^\circ$.9.55 at 20° .8.13 at 40° .7.01 at 60° .

(Hofmann, Phys. Zeit. 1905, 6. 339.)

Rhodicyanhydric acid, $H_2Rh(CN)_4$.

Not known in the free state.

Potassium rhodicyanide, $K_2Rh(CN)_4$.Sol. in H_2O . Easily decomp. by acidVery sol. in H_2O . (Leidié, C. R. 19089.)

Rhodium, Rh.

Insol. in all acids, including aqua regia

Rhodium "sponge" is sol. in HNO_3 and somewhat in $HCl + Aq$ when exp. air.

Rhodium ammonia compounds.

See—

Bromopurpleorhodium comp., $BrRh(NH_3)_4X_2$.Chloropurpleorhodium comp., $ClRh(NH_3)_4X_2$.Iodopurpleorhodium comp., $IRh(NH_3)_4X_2$.Luteorhodium comp., $Rh(NH_3)_4X_2$.Nitratopurpleorhodium comp., $(NO_3)Rh(NH_3)_4X_2$.Roseorhodium comp., $Rh(NH_3)_4(OH)$.Xanthorhodium comp., $(NO_3)Rh(NH_3)_4$.Rhodium tribromide, $RhBr_3 + 2H_2O$.Very sol. in H_2O . (Goloubkine, Soc. 1911, 100. (2) 45.)

Rhodium rubidium bromide.

See Bromorhodite, rubidium.

Rhodium sodium bromide.

See Bromorhodite, sodium.

Rhodium dichloride, $RhCl_2$ (?).Insol. in H_2O , HCl , or $HNO_3 + Aq$ attacked by boiling KOH or K_2CO_3 (Fellenberg.)Decomp. by boiling $KOH + Aq$. (Ber)

Does not exist. (Leidié, C. R. 108.)

Rhodium trichloride, $RhCl_3$.Insol. in acids, even aqua regia. boiled for a long time with $KOH + Aq$ comes sl. sol. in $HCl + Aq$.Insol. in H_2O and acids; sol. in alkalis (Leidié, C. R. 1899, 129. 1251.)+ $4H_2O$. Very sl. deliquescent. Sol. in H_2O , $HCl + Aq$, or alcohol. Insol. in H_2SO_4 only when boiling. (J. pr. 80. 282.)

ite amount of crystal H_2O . (Leidié 17. 271.)

chloride with MCl .
rhodite, M .

dihydroxide, $RhO_2 \cdot 2H_2O$, or
rhodate, $Rh_2O_3 \cdot RhO_2 + 6H_2O$.
 $ICl + Aq$.

sesquihydroxide, $Rh_2O_3 \cdot H_2O$.

sol. in conc. $HCl + Aq$. (Claus.)
Easily sol. in HCl , H_2SO_4 , H_2SO_3 ,
 $HSCN + Aq$; also when moist, in
 H_2O . Sol. in conc. $KOH + Aq$; very
sl. in H_3BO_3 , H_3PO_4 , $H_2C_4H_4O_6$, and HCN .
Insol. in acid alkali oxalates $+ Aq$.
R. 107. 234.)

triiodide, RhI_3 .

Goloubkine, Chem. Soc. 1911, 100.

monoxide, RhO .

attacked by acids. (Deville and De
la Rüe. (3) 61. 83.)

sesquioxide, Rh_2O_3 .

Insol. in H_2O , boiling $KOH + Aq$, or any
aqua regia. (Claus.)

dioxide, RhO_2 .

Insol. in all acids or alkalies.

trioxide, RhO_3 .

“acid.” Known only in solution of
rhodate,” which is very easily
(Claus.)

oxybromide, $Rh(OH)_2Br + 2H_2O$.

Insol. in H_2O . (Goloubkine, Chem. Soc.
(2) 45.)

monosulphide, RhS .

Insol. in aqua regia.

sesquisulphide, Rh_2S_3 .

Insol. in alkali sulphides $+ Aq$. (Debray, C.
2.)

Insol. in alkali sulphides $+ Aq$. Not at-
tacked by HNO_3 , aqua regia, or $Br_2 + Aq$.
Ill. Soc. (2) 50. 664.)

sodium sulphide, $3Na_2S, Rh_2S_3$.

Insol. in H_2O . (Leidié.)

sesquisulphydroxide, $Rh_2S_3 \cdot H_2O$.

Insol. in aqua regia or $Br_2 + Aq$. Insol.
in alkali sulphides $+ Aq$ or acids. (Leidié,
(2) 50. 664.)

Rhodochromium bromide,

$HOCr_2(NH_3)_{10}Br_5 + H_2O$.

Rather difficultly sol. in H_2O . Decomp. by
boiling or standing. Sol. in $NH_4OH + Aq$
or $NaOH + Aq$. Insol. in dil. $HBr + Aq$,
 $KBr + Aq$, or alcohol. (Jørgensen, J. pr. (2)
25. 321.)

— bromide, basic, $HOCr_2(NH_3)_{10}(OH)Br_4 + H_2O$.

Sl. sol. in H_2O . Sol. in NH_4OH or $NaOH + Aq$. Insol. in alcohol. (Jørgensen.)

— bromoplatinate, $HOCr_2(NH_3)_{10}Br_3PtBr_6$,
 $HOCr_2(NH_3)_{10}Br_3(PtBr_6)_2 + 4H_2O$.

Ppt. (Jørgensen.)

— chloraurate, $HOCr_2(NH_3)_{10}Cl_3(AuCl_4)_2 + 2H_2O$.

Difficultly sol. but not insol. in H_2O . (Jørgensen.)

— chloride, $HOCr_2(NH_3)_{10}Cl_5 + H_2O$.

Sol. in about 40 pts. of cold H_2O . Insol. in
cold dil. $HCl + Aq$, $NH_4Cl + Aq$, or alcohol.
Sol. in $NH_4OH + Aq$. (Jørgensen, J. pr. (2)
25. 321.)

— chloriodide, basic,
 $HOCr_2(NH_3)_{10}(OH)Cl_3I_2$.

Sl. sol. in cold H_2O ; insol. in alcohol.
(Jørgensen.)

— chloroplatinate, $HOCr_2(NH_3)_{10}Cl_3PtCl_6$,
 $HOCr_2(NH_3)_{10}Cl(PtCl_6)_2 + 4H_2O$.

Precipitate. (Jørgensen.)

— dithionate, $[HOCr_2(NH_3)_{10}]_2(S_2O_6)_5 + 2H_2O$.

Nearly insol. in H_2O .

— dithionate, basic,
 $HOCr_2(NH_3)_{10}OH(S_2O_6)_2 + H_2O$.

Insol. in H_2O , cold $NH_4OH + Aq$, or
 $NaOH + Aq$.

— iodide, $HOCr_2(NH_3)_{10}I_5 + H_2O$.

Very difficultly sol. in H_2O . Insol. in very
dil. $HI + Aq$ or alcohol. Sl. sol. in NH_4OH or
 $KOH + Aq$. (Jørgensen.)

— nitrate, $HOCr_2(NH_3)_{10}(NO_3)_5$.

Rather difficultly sol. in H_2O , from which
it is precipitated by a few drops of $HNO_3 + Aq$.
Sol. in hot dil. $NH_4OH + Aq$.

— nitrate chloroplatinate,
 $HOCr_2(NH_3)_{10}(NO_3)(PtCl_6)_2 + 4H_2O$.

Precipitate. (Jørgensen.)

Rhodochromium sulphate,

$[HOCr_2(NH_3)_{10}]_2(SO_4)_5 + 2H_2O$.

Very sl. sol. in cold H_2O . Easily sol. in
cold dil. $H_2SO_4 + Aq$.

Almost insol. in a mixture of 3 vols. H_2O , 1 vol. alcohol, and $\frac{1}{2}$ vol. dil. $H_2SO_4 + Aq.$ (Jørgensen.)

Rhodonitrous acid.

Ammonium rhodonitrite, $(NH_4)_2Rh_2(NO_2)_{12}$.

Nearly insol. in cold, sl. sol. in hot H_2O . Insol. in conc. NH_4Cl or $NH_4C_2H_3O_2 + Aq.$ Insol. in alcohol. (Leidié, C. R. 111. 108.)

Barium rhodonitrite, $Ba_2Rh_2(NO_2)_{12}$.

Sl. sol. in cold, more easily in hot H_2O . (Lamy.)

+ $12H_2O$. Sol. in 50 pts. H_2O at 16° , and 6.5 pts. at 100° . (Leidié, C. R. 111. 108.)

Potassium rhodonitrite, $K_2Rh_2(NO_2)_{12}$.

Nearly insol. in cold, very sl. sol. in boiling H_2O . Completely insol. in $KNO_3 + Aq.$ and in $KCl + Aq$ (30% KCl), or $KC_2H_3O_2 + Aq$ (50% $KC_2H_3O_2$). Insol. in alcohol. (Leidié, C. R. 111. 106.)

Sodium rhodonitrite, $Na_2Rh_2(NO_2)_{12}$.

Sol. in $2\frac{1}{2}$ pts. H_2O at 17° , and 1 pt. at 100° . Insol. in alcohol. Decomp. by $HCl + Aq.$ (Leidié, C. R. 111. 107.)

Rhodosochromium bromide.

Sol. in H_2O ; insol. in dil. $HBr + Aq$ (1 : 1). (Jørgensen, J. pr. (2) 45. 260.)

— chloraurate, $Cr_2(NH_4)_6(OH)_8Cl_3$, $2AuCl_3 + 2H_2O$.

Not insol. in cold H_2O . (Jørgensen.)

— chloride, $Cr_2(NH_4)_6(OH)_8Cl_3 + 2H_2O$.

Sol. in 10.6 pts. H_2O at 18° ; decomp. by boiling. Pptd. by $\frac{1}{2}$ to 1 vol. dil. $HCl + Aq.$ Sol. in cold dil. $NH_4OH + Aq.$ (Jørgensen, J. pr. (2) 45. 260.)

— chloroplatinate, $2Cr_2(NH_4)_6(OH)_8Cl_3$, $3PtCl_4 + 6H_2O$.

Insol. in H_2O . (Jørgensen.)

$Cr_2(NH_4)_6(OH)_8Cl_3$, $2PtCl_4 + 2H_2O$. Insol. in 95% alcohol. (Jørgensen.)

— chromate, $[Cr_2(NH_4)_6(OH)_8]_2(CrO_4)_3 + 7H_2O$. (Jørgensen.)

Very sl. sol. in H_2O . (Jørgensen.)

— iodide, $Cr_2(NH_4)_6(OH)_8I_3 + 2H_2O$.

Sol. in H_2O . Insol. in dil. $HI + Aq.$ (Jørgensen.)

— nitrate, $Cr_2(NH_4)_6(OH)_8(NO_3)_3 + H_2O$.

Much less sol. in cold H_2O than the chloride. Insol. in dil. $HNO_3 + Aq.$ (Jørgensen.)

— oxalate, $[Cr_2(NH_4)_6(OH)_8]_2(C_2O_4)(HC_2O_4)_4 + 2H_2O$.

Sol. in cold H_2O , but not very easily. (Jørgensen.)

Rhodosochromium sulphate,

$[Cr_2(NH_4)_6(OH)_8]_2(SO_4)_3 + 5H_2O$.

Very sl. sol. in cold H_2O . Easily sol. $NH_4Cl + Aq.$ (Jørgensen.)

$[Cr_2(NH_4)_6(OH)_8]SO_4 \cdot HSO_4 + 1\frac{1}{2}H_2O$ comp. by H_2O into H_2SO_4 and above pound. (Jørgensen.)

— persulphide, $[Cr_2(NH_4)_6(OH)_8]_2S_{11} \cdot 4H_2O$.

Ppt. Insol. in H_2O . (Jørgensen.)

Rhodosulphuric acid.

Potassium rhodosulphate, $K_2Rh_2(SO_4)_4$.

Two modifications:

(a) Slowly sol. in cold, easily in hot H

(b) Insol. in H_2O .

Does not exist. (Leidié, C. R. 107. 2)

Sodium rhodosulphate.

Insol. in H_2O , HCl , HNO_3 , or aqua (Claus.)

Does not exist. (Leidié.)

$Na_2Rh_2(SO_4)_4$. Insol. in H_2O . (Seubert and Kobbé, B. 23. 2560.)

Rhodosulphurous acid.

Potassium rhodosulphite, $K_2Rh_2(SO_3)_4 \cdot 6H_2O$.

Nearly insol. in H_2O . Slowly sol. in Not decomp. by boiling $KOH + Aq.$ (Claus.)

Sodium rhodosulphite,

$Na_2Rh_2(SO_3)_4 + 4\frac{1}{2}H_2O = 3Na_2SO_3 + 2RhSO_3$.

Insol. in cold, very sl. sol. in hot Easily sol. in $HNO_3 + Aq.$ (Seubert Kobbé, B. 23. 2558.)

Roseochromium bromide,

$Cr(NH_4)_3Br_3 + H_2O$.

Easily sol. in H_2O . Insol. in HBr (Christensen, J. pr. (2) 23. 26.)

— bromochromate, $Cr(NH_4)_3Br(CrO_3)$

Somewhat sol. in H_2O , but decomp. standing. (Jørgensen, J. pr. (2) 25. 306)

— bromoplatinate, $Cr(NH_4)_3Br(PtCl_6) \cdot 2H_2O$.

Precipitate. Difficultly sol. in H_2O . (Christensen, l. c.)

— chloride, $Cr(NH_4)_3Cl_3 + H_2O$.

Easily sol. in H_2O with subsequent decomp. Insol. in alcohol. (Christensen, J. pr. (2) 26.)

— mercuric chloride, $Cr(NH_4)_3Cl_3 \cdot HgCl_2 + 2H_2O$.

Sl. sol. in H_2O . Sol. in dil. $HCl + Aq.$ decomposition. (Christensen, l. c.)

omium dithionate, basic,
 $\text{Co}(\text{NH}_3)_6(\text{OH})_2\text{S}_2\text{O}_8 + \text{H}_2\text{O}$.

sol. in very dil. $\text{HCl} + \text{Aq}$. (Jörgensen, (2) 25. 398.)

de, $\text{Cr}(\text{NH}_3)_6\text{I}_3$.

sol. in H_2O ; decomp. by boiling. (Jörgensen, l. c.)

rate, $\text{Cr}(\text{NH}_3)_6(\text{NO}_3)_3 + \text{H}_2\text{O}$.

easily sol. in H_2O . (Christensen,

$\text{I}_3)(\text{NO}_3)_3(\text{OH}_2)_3$, HNO_3 . Decomp. in alcohol. (Jörgensen, J. pr. (2) 44.

hate, $[\text{Cr}(\text{NH}_3)_6]_2(\text{SO}_4)_3 + 5\text{H}_2\text{O}$.

sol. in H_2O . Precipitated by alcohol. (Jörgensen, l. c.)

hate bromoplatinate,
 $\text{Co}(\text{NH}_3)_6(\text{SO}_4)_2\text{PtBr}_6$.

slightly sol. in H_2O . (Christensen, l. c.)

hate chloroplatinate,
 $\text{Co}(\text{NH}_3)_6(\text{SO}_4)_2\text{PtCl}_6$.

slightly sol. in H_2O . (Christensen, l. c.)

baltic bromide,

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Br}_3$.

H_2O ; insol. in $\text{HBr} + \text{Aq}$. (Jörgensen, (2) 31. 49.)

**oplatinate, $\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Br}_3$,
 $4 + \text{H}_2\text{O}$.**

slightly sol. in H_2O or dil. alcohol. Insol. in alcohol. (Jörgensen.)

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Br}_3$, $3\text{PtBr}_4 + 4\text{H}_2\text{O}$. Ppt. (Jörgensen.)

nosulphate,

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Br}(\text{SO}_4)$.

H_2O . (Krok.)

nosulphate bromaurate,

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)(\text{SO}_4)\text{Br}$, AuBr_3 .

onate.

sol. in H_2O .

aurate, $\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$, AuCl_3 .
 easily sol. in cold H_2O .

ride, $\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$.

4.8 pts. H_2O at 10.1° , but decomp. g.

1. H_2O dissolve 16.12 pts. at 0° , and 16.19° at 16.19° . (Kurnakoff, J. russ. Soc.

in 1000 pts. fuming $\text{HCl} + \text{Aq}$, more 20% $\text{HCl} + \text{Aq}$. (Rose.)

Roseocobaltic mercuric chloride,

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$, $3\text{HgCl}_2 + \text{H}_2\text{O}$.

More easily sol. in solvents than the anhydrous purpureo salt. (Carstanjen.)

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$, HgCl_2 . Sol. in $\text{HCl} + \text{Aq}$ with decomp. into above salt. (Jörgensen.)

— **chloroplatinate,**

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$, $\text{PtCl}_4 + \frac{1}{2}\text{H}_2\text{O}$.

Decomp. by H_2O . (Jörgensen.)

$2\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$, $\text{PtCl}_4 + 2\text{H}_2\text{O}$. Decomp. by H_2O .

$2\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Cl}_3$, $3\text{PtCl}_4 + 6\text{H}_2\text{O}$. Not difficultly sol. in warm H_2O . (Gibbs.)

$\text{Co}(\text{NH}_3)_6\text{Cl}_3$, $\text{PtCl}_4 + \text{H}_2\text{O}$. (Gibbs.)

— **chlorosulphate, $\text{Co}(\text{NH}_3)_6\text{Cl}(\text{SO}_4)$.**

Easily sol. in H_2O .

— **chlorosulphate mercuric chloride,**

$\text{Co}(\text{NH}_3)_6\text{Cl}(\text{SO}_4)$, $\text{HgCl}_2 + 3\text{H}_2\text{O}$.

Sol. in hot H_2O , and can be recrystallized without decomp. (Krok.)

— **dichromate,**

$[\text{Co}(\text{NH}_3)_6]_2(\text{Cr}_2\text{O}_7)_3 + 5\text{H}_2\text{O}$.

Can be recrystallized out of weak acetic acid.

— **cobalticyanide, $\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{Co}(\text{CN})_6$.**

Nearly absolutely insol. in cold H_2O . (Jörgensen.)

$+ \text{H}_2\text{O}$. (Gibbs and Gentile.)

— **dithionate, $\text{Co}(\text{NH}_3)_6(\text{S}_2\text{O}_8)(\text{OH})$.**

Decomp. by H_2O . (Rammelsberg, Pogg. 58. 296.)

$\text{Co}(\text{NH}_3)_6(\text{OH}_2)(\text{S}_2\text{O}_8) + 2\text{H}_2\text{O}$. Ppt. (Jörgensen.)

— **hydroxide, $\text{Co}(\text{NH}_3)_6(\text{OH})_3$.**

Known only in aqueous solution.

— **mercuric hydroxychloride,**

$\text{CoN}_5\text{H}_{12}(\text{HgOH})_2\text{Cl}_3$.

Ppt. Sol. in dil. acids. (Vortmann and Morgulis, B. 22. 2646.)

$\text{CoN}_5\text{H}_{12}(\text{HgOH})_2\text{Cl}_2(\text{OH})$. Ppt. Sol. in dil. acids. (Vortmann and Morgulis.)

— **iodide, $\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{I}_3$.**

Less sol. in H_2O than bromide. Insol. in $\text{HI} + \text{Aq}$. (Jörgensen.)

— **iodosulphate, $\text{Co}(\text{NH}_3)_6(\text{OH}_2)\text{I}(\text{SO}_4)$.**

Easily sol. in H_2O . (Krok.)

— **mercuriodide, $[\text{CoN}_5\text{H}_{12}]_2(\text{HgI})_2\text{I}_3$.**

Ppt. (Vortmann and Borsbach, B. 23. 2805.)

$\text{CoN}_5\text{H}_{12}(\text{HgI})_2\text{I}_3$. Ppt. (Vortmann and Borsbach.)

$\text{CoN}_5\text{H}_{12}(\text{HgI})_2\text{I}_2(\text{OH})$. Ppt.

Roseocobaltic nitrate,

Three modifications:

α . Sol. in 20 pts. H_2O at 15° . (Jørgensen.)

β . Known only in solution. Insol. in cold $\text{HNO}_3 + \text{Aq.}$ (Gibbs.)

γ . Easily sol. in hot H_2O . (Gibbs.) (Purpureo salt?)

$\text{Co}(\text{NH}_3)_5(\text{OH}_2)(\text{NO}_3)_2$, HNO_3 . Decomp. by H_2O or alcohol. (Jørgensen, J. pr. (2) 44. 63.)

— **nitrate chloroplatinate,**



Ppt. (Jørgensen.)

— **nitratosulphate,**



Sl. sol. in cold, easily in hot H_2O .

— **oxalate,** $[\text{Co}(\text{NH}_3)_5(\text{OH}_2)]_2(\text{C}_2\text{O}_4)_3 + 2\text{H}_2\text{O}.$

Nearly insol. in H_2O .



— **oxalochloroplatinate,**



Sol. in hot H_2O .

— **oxalosulphate,** $[\text{Co}(\text{NH}_3)_5]_2(\text{SO}_4)_2\text{C}_2\text{O}_4, \text{H}_2\text{C}_2\text{O}_4 + 2\text{H}_2\text{O}.$

Sol. in hot H_2O .

$[\text{Co}(\text{NH}_3)_5]_2(\text{SO}_4)_2(\text{C}_2\text{O}_4)(\text{OH})_2 + 6\text{H}_2\text{O}$. Sl. sol. in H_2O .

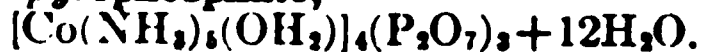
— **orthophosphate,**



Nearly insol. in H_2O .

$[\text{Co}(\text{NH}_3)_5(\text{OH}_2)]_2(\text{PO}_4\text{H})_3 + 4\text{H}_2\text{O}$. Very sl. sol. in cold H_2O ; easily in H_2O containing HCl . (Jørgensen.)

— **pyrophosphate,**



Insol. in H_2O . (Jørgensen.)

$\text{Co}(\text{NH}_3)_5(\text{OH}_2)(\text{P}_2\text{O}_7\text{Na}) + 12\text{H}_2\text{O}$. Nearly insol. in cold, easily sol. in hot H_2O containing NH_4OH . (Jørgensen, J. pr. (2) 23. 252.)

— **sulphate,** $[\text{Co}(\text{NH}_3)_5(\text{OH}_2)]_2(\text{SO}_4)_3 + 3\text{H}_2\text{O}.$

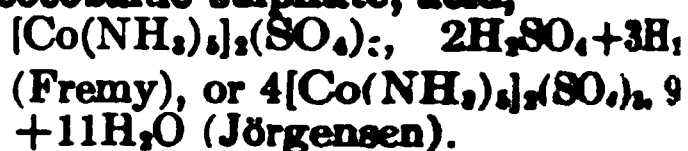
Three modifications:

α . Sl. sol. in cold H_2O . Sol. in 58 pts. at 27° (Gibbs); 83.5 pts. at 20.2° , and 94.6 pts. at 17.2° (Jørgensen); more easily sol. in hot H_2O , and still more easily in $\text{NH}_4\text{OH} + \text{Aq.}$

β . Sol. in 1–2 pts. H_2O . (Gibbs.)

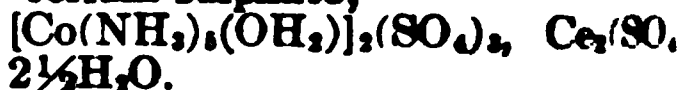
γ . Less sol. than lutesulphate. (Jørgensen.)

$+ 2\text{H}_2\text{O}$. Easily sol. in H_2O . (Vortmann.)

Roseocobaltic sulphate, acid,

More easily sol. in H_2O than neutral sulphate, into which it is converted by neutralization. Sol. in about 13 pts. H_2O . (Jørgensen.)

— **cerium sulphate,**



Sl. sol. in cold, practically insol. in hot H_2O . Sol. in acids. (Gibbs, Am. Ch. 560.)

$[\text{Co}(\text{NH}_3)_5(\text{OH}_2)]_2(\text{SO}_4)_3, \text{Ce}(\text{SO}_4)_3 + 2\frac{1}{2}\text{H}_2\text{O}$. As above. (Gibbs.)

— **sulphate chloraurate.**

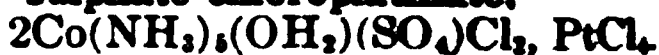
Three modifications:

α . $\text{Co}(\text{NH}_3)_5(\text{OH}_2)(\text{SO}_4)\text{Cl}$, AuCl_3 . (Jørgensen.)

β . $\text{Co}(\text{NH}_3)_5(\text{SO}_4)$, $\text{AuCl}_3 + 2\text{H}_2\text{O}$. Sl. in cold H_2O . (Gibbs.)

γ . As above. Can be recrystallized from hot H_2O .

— **sulphate chloroplatinate.**

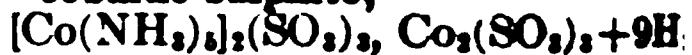


Three modifications, all difficultly sol. in cold H_2O . (Jørgensen.)

— **sulphite,** $[\text{Co}(\text{NH}_3)_5(\text{OH}_2)]_2(\text{SO}_3)_2 + 2\text{H}_2\text{O}.$

Sl. sol. in cold, decomp. by hot H_2O . (Gibbs.)

— **cobaltic sulphite,**



Insol. in cold, decomp. by hot H_2O . (Jørgensen.)

Roseocobaltic octamine compounds

See Roseotetramine cobaltic compounds.

Roseoiridium compounds.

See Irido-aquopentamine compounds.

Roseorhodium bromide,

Sol. in cold H_2O . (Jørgensen, J. pr. (2) 394.)

Roseorhodium cobalticyanide,

Scarcely sol. in H_2O .

— **iodosulphate,** $\text{Rh}(\text{NH}_3)_5(\text{OH}_2)\text{I}(\text{SO}_4).$

Very sl. sol. in H_2O ; easily sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Jørgensen.)

— **nitrate,** $\text{Rh}(\text{NH}_3)_5(\text{OH}_2)(\text{NO}_3)_2.$

Moderately sol. in cold H_2O . (Jørgensen.)



or alcohol. (Jørgensen, J. pr. (2) 44.

iodium nitrate chloroplatinate,
 $(\text{NH}_4)_4(\text{OH})_2(\text{NO}_3)_2\text{PtCl}_6 + 2\text{H}_2\text{O}$.
 (Jørgensen.)

hypophosphate,
 $(\text{NH}_4)_4(\text{OH})_2(\text{HPO}_3)_2 + 4\text{H}_2\text{O}$.
 sl. sol. in H_2O .

dium pyrophosphate,
 $(\text{NH}_4)_4(\text{OH})_2\text{NaP}_2\text{O}_7 + 23\text{H}_2\text{O}$.
 Very sl. sol. in cold H_2O . Easily sol.
 dil. acids.

lphate,
 $(\text{NH}_4)_4(\text{OH})_2(\text{SO}_4)_2 + 3\text{H}_2\text{O}$.
 sl. sol. in cold, much more in hot H_2O .

lphate chloroplatinate,
 $(\text{NH}_4)_4(\text{OH})_2(\text{SO}_4)_2\text{PtCl}_6$.
 Nearly insol. in H_2O or alcohol.

etramine cobaltic bromide,
 $(\text{NH}_4)_4(\text{OH})_2\text{Br}_2$.
 n H_2O ; insol. in $\text{HBr} + \text{Aq}$. Nearly in-
 lcohol. (Jørgensen, Z. anorg. 2. 295.)

— **chloride,** $\text{Co}(\text{NH}_4)_4(\text{OH})_2\text{Cl}_2$.
 y sol. in H_2O ; insol. in conc. $\text{HCl} + \text{Aq}$;
 at. $\text{HgCl}_2 + \text{Aq}$. (Jørgensen.)

— **cobalticyanide,**
 $(\text{NH}_4)_4(\text{OH})_2\text{Co}(\text{CN})_6$.
 ensen.)

— **oxalate sulphate,**
 $(\text{NH}_4)_4(\text{OH})_2(\text{SO}_4)_2\text{C}_2\text{O}_4$.
 (Jørgensen.)

— **pyrophosphate,**
 $(\text{NH}_4)_4(\text{OH})_2(\text{P}_2\text{O}_7)_2 + 6\text{H}_2\text{O}$.
 ly insol. in H_2O , but easily sol. in very
 ls + Aq . (Jørgensen.)

— **sulphate,**
 $(\text{NH}_4)_4(\text{OH})_2(\text{SO}_4)_2 + 3\text{H}_2\text{O}$.
 n about 35 pts. H_2O , and more easily
 tion of dil. HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$. (Jör-
)

— **sulphate bromaurate,**
 $(\text{NH}_4)_4(\text{OH})_2(\text{SO}_4)_2\text{AuBr}_4$.
 ol. in cold H_2O ; insol. in alcohol.
 sen.)

— **sulphate chloroplatinate,**
 $(\text{NH}_4)_4(\text{OH})_2(\text{SO}_4)_2\text{PtCl}_6$.
 e bromaurate. (Jørgensen.)

Rubidium, Rb.

Decomp. H_2O with violence. Insol. in
 hydrocarbons. Sol. in liquid NH_3 . (Seely,
 C. N. 23. 169); (Franklin, Am. Ch. J. 1898,
 20. 829).

Rubidium acetylide acetylene RbC_2 , C_2H_2 .

Very hygroscopic.
 Insol. in CCl_4 and in ether. (Moissan, C.
 R. 1903, 136. 1220.)

Rubidium amalgam, RbHg_{12} .

Stable in contact with Hg below 0° .
 Above 0° the composition of the amalgam
 varies. Can be cryst. from Hg without de-
 comp. below 0° . (Kerp, Z. anorg. 1900, 25.
 68.)

Rubidium amide, RbNH_2 .

Very deliquescent. Violently decomp. by
 H_2O ; less violently acted on by alcohol.
 (Titherley, Chem. Soc. 1897, 71. 470.)

Rubidium ammonia, RbNH_3 .

Decomp. by H_2O .
 Very sol. in liquid NH_3 . (Moissan, C. R.
 1903, 136. 1178.)

Rubidium azoimide, RbN_3 .

Sl. hygroscopic.
 Stable in aq. solution.
 107.1 pts. are sol. in 100 pts. H_2O at 16° .
 114.1 " " " " 100 " H_2O " 17° .
 0.182 " " " " 100 " abs. alcohol at
 16° .

Insol. in pure ether.
 (Curtius, J. pr. 1898, (2) 58. 281.)

Rubidium bromide, RbBr .

100 pts. H_2O dissolve 98 pts. at 5° ; 104.8
 pts. at 16° . (Reissig, A. 127. 33.)

Solubility in H_2O .

100 pts. of the solution contain at:

0.5°	5.0°	16.0°
47.26	49.50	51.17 pts. RbBr ,

39.7°	57.5°	113.5°
56.87	60.39	67.24 pts. RbBr .

(Rimbach, B. 1905, 38. 1557.)

Sp. gr. of RbBr containing g. equiv. RbBr
 per l.

G. equiv. RbBr	Sp. gr. $6^\circ/6^\circ$	Sp. gr. $18^\circ/18^\circ$	Sp. gr. $30^\circ/30^\circ$
0.508	1.06448	1.06389	1.06326
1.020	1.12931	1.12799	1.12626
2.031	1.25622	1.25366	1.25187
4.072	1.50574	1.50107	1.49870

(Clausen, W. Ann. 1914, (4) 44. 1070.)

RbBr + Aq. containing 6.60% RbBr has sp. gr. $20^{\circ}/20^{\circ} = 1.0525$.

RbBr + Aq. containing 14.36% RbBr has sp. gr. $20^{\circ}/20^{\circ} = 1.1226$.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 279.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

Difficultly sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Rubidium tribromide, RbBr₃.

Very sol. in H₂O; decomp. by alcohol and ether. (Wells and Wheeler, Sill. Am. J. 143. 475.)

Rubidium ruthenium bromide.

See Bromoruthenate and bromoruthenite, rubidium.

Rubidium selenium bromide.

See Bromoselenate, rubidium.

Rubidium tellurium bromide.

See Bromotellurate, rubidium.

Rubidium thallic bromide, RbBr, TlBr₃ + H₂O.

Recryst. from H₂O unchanged. (Pratt, Am. J. Sci. 1895, (3) 49. 403.)

3RbBr, TlBr₃ + H₂O. Very sol. in H₂O. (Pratt.)

Rubidium stannic bromide.

See Bromostannate, rubidium.

Rubidium bromochloride, RbBr₂Cl.

Easily decomp., even by H₂O. (Wells and Wheeler.)

RbBrCl₂. Sol. in H₂O; decomp. by alcohol and ether. (Wells and Wheeler.)

Rubidium bromochloriodide, RbBrClI.

Sol. in H₂O and alcohol. Decomp. by ether. (Wells and Wheeler.)

Rubidium bromiodide, RbBr₂I.

Very sol. in H₂O. Sat. solution contains about 44% RbBr₂I, and sp. gr. = 3.84. (Wells and Wheeler.)

Rubidium carbide, Rb₂C₂.

Decomp. violently by H₂O. (Moissan, C. R. 1903, 136. 1221.)

Rubidium chloride, RbCl.

100 pts. H₂O dissolve 76.38 pts. at 1°; 82.89 pts. at 7°. (Bunsen.)

Solubility in H₂O at t°.
100 pts. of the solution contain pts.

t°	Pts. RbCl
0.4	43.61
15.5	46.56
57.3	53.71
114.9	59.48

(Rimbach, B. 1902, 35. 1304.)

Solubility of RbCl in H₂O at

t°	G. RbCl per 100 g.		t°	G. RbCl
	H ₂ O	Solution		H ₂ O
0	77.0	43.5	60	115.5
10	84.4	45.8	70	121.4
20	91.1	47.7	80	127.2
30	97.6	49.4	90	133.1
40	103.5	50.9	100	138.9
50	109.3	52.2	112.9	146.6

(Berkeley, Phil. Trans. Roy. Soc. 19 A. 189.)

Sat. RbCl + Aq at 25° contains RbCl. (Foote, Am. Ch. J. 1906, 35.

Sp. gr. of RbCl + Aq containing in H₂O:

13.14 25.88 33.13 pts. Rb
1.1066 1.2156 1.2675 sp. gr.

(Tammann, W. Ann. 24. 1885.)

A normal solution of RbCl has at 25° = 1.0610. (Wagner, Z. phys. Cl 5. 39.)

RbCl + Aq containing 6.64% Rb sp. gr. $20^{\circ}/20^{\circ} = 1.0502$.

RbCl + Aq containing 10.59% Rb sp. gr. $20^{\circ}/20^{\circ} = 1.0815$. (Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 278.)

Sp. gr. $20^{\circ}/4^{\circ}$ of a normal solution = 1.085405. (Haigh, J. Am. Chem. Soc. 1912, 34. 1151.)

Sp. gr. of RbCl + Aq sat. at t°

t°	Sp. gr.	t°	
0.55	1.4409	60.25	1
18.7	1.4865	75.15	1
31.5	1.5118	89.35	1
44.7	1.5348	114°	1

* Boiling point.

(Berkeley.)

Sp. gr. of RbCl containing g. equiv. per l.

G. equiv. RbCl	Sp. gr. 6°/6°	Sp. gr. 18°/18°	
0.5123	1.06410	1.04538	1
1.001	1.08916	1.08810	1
2.073	1.18200	1.17959	1
3.984	1.34334	1.33067	1

(Clausen, W. Ann. 1914, (4) 44. 1)

sol. in liquid NH_3 . (Franklin, Am. 98, 20. 829.)

ity in alcohols at 25° .

methanol dissolve 1.41 g.

ethyl alcohol dissolve 0.078 g.

propyl alcohol dissolve 0.015 g.

isoamyl alcohol dissolve 0.0025 g.

r and Bissett, Chem. Soc. 1913, 103.

in anhydrous pyridine and in 97%

+Aq. Very sl. sol. in 95% pyridine

sol. in 93% pyridine+Aq. (Kah-

l. Am. Chem. Soc. 1908, 30. 1107.)

in methyl acetate. (Naumann, B.

3790); acetone; (Naumann, B. 1904,

; (Eidmann, C. C. 1899, II. 1014).

ruthenium trichloride.

lororuthenite, rubidium.

ruthenium tetrachloride.

ororuthenate, rubidium.

oxyruthenium chloride,

RuO_4Cl_4 .

decomp. by H_2O ; sol. in cold HCl .

Am. Chem. Soc. 1901, 23. 779.)

tellurium chloride.

orotellurate, rubidium.

thallous chloride, 2RbCl , TlCl_3 +

recryst. from H_2O without change.

m. J. Sci. 1895, (3) 49. 399.)

TiCl_3 . Crystallizes from HCl solu-

ummann, A. 244. 348.)

Very sol. in cold H_2O . (Pratt,

i. 1895, (3) 49. 398.)

Efflorescent in dry air. Sol. in

H_2O at 18° , and 1.6 pts. at 100° .

y, Zeitschr. d. allgem. österr.

rv. 1880. No. 9.)

stannous chloride.

orostannate, rubidium.

titanium chloride, 2RbCl , TiCl_3 ,

H_2O . (Stähler, B. 1904, 37. 4408.)

tungsten chloride, $\text{Rb}_2\text{W}_2\text{Cl}_6$.

in cold, more sol. in hot H_2O .

very dil. NaOH +Aq.

insol. in most organic solvents.

. 1913, 46. 574.)

uranous chloride, Rb_2UCl_6 .

It. (Aloy, Bull. Soc. 1899, (3) 21.

Rubidium uranyl chloride, 2RbCl , $(\text{UO}_2)\text{Cl}_2$,
+ $2\text{H}_2\text{O}$.

Solubility in H_2O .

100 pts. of the solution contain at:

24.8° 80.3°

57.8 65.73 pts. UO_2Cl_2 , 2RbCl .

(Rimbach, B. 1904, 37. 467.)

Rubidium vanadium chloride, Rb_2VCl_6 ,
+ H_2O .

Sl. sol. in H_2O and alcohol.

Decomp. by H_2O on standing so that it
dissolves. (Stähler, B. 1904, 37. 4411.)

Rubidium zinc chloride, 2RbCl , ZnCl_2 .

Easily sol. in H_2O and HCl +Aq. (Godef-
froy, B. 8. 9.)

Rubidium chloride selenium dioxide, RbCl ,
 2SeO_2 + $2\text{H}_2\text{O}$.

Sol. in H_2O . (Muthmann, B. 1893, 26.
1013.)

Rubidium chloroiodide, RbCl_2I .

Properties are similar to those of RbBrClI .
(Wells.)

RbCl_2I . Sol. in alcohol, not attacked by
ether. (Wells and Wheeler, Sill. Am. J. 144.
42.)

Sol. in POCl_3 . (Walden, Z. anorg. 1900,
25. 212.)

Nearly insol. in AsBr_3 . (Walden, Z. anorg.
1902, 29. 374.)

Very stable; sl. sol. in H_2O at 0° ; only very
sl. sol. in HCl . (Erdmann, Arch. Pharm.
1894, 232. 32.) (C. C. 1894, I. 670.)

Rubidium fluoride, RbF .

Very sol. in H_2O . (Eggeling, Z. anorg.
1905, 46. 174.)

100 g. H_2O dissolve 130.6 g. RbF at 18° .
(de Forcrand, C. R. 1911, 152. 1210.)

Sol. in dil. HF . (Pennington, J. Am. Chem.
Soc. 1896, 18. 57.)

Insol. in liquid NH_3 . (Core, Am. Ch. J.
1898, 20. 829.)

Rubidium hydrogen fluoride, RbF , HF .

Very deliquescent.

Insol. in alcohol and ether. (Chabrié,
C. R. 1905, 140. 91.)

Very hygroscopic. Sol. in H_2O . (Eggeling,
Z. anorg. 1905, 46. 175.)

RbF , 2HF . Very sl. sol. in H_2O . (Egge-
ling, Z. anorg. 1905, 46. 176.)

Rubidium silicon fluoride.

See Fluosilicate, rubidium.

Rubidium tantalum fluoride.

See Fluotantalate, rubidium.

Rubidium uranyl fluoride, 4RbF , UO_2F_2 +
 $6\text{H}_2\text{O}$.

(Ditte, C. R. 91. 115.)

Ruthenium dihydronitrosohydroxide,
 $\text{NO} \cdot \text{Ru}_2\text{H}_2(\text{OH})_2 + 2\text{H}_2\text{O}.$

Sl. sol. in cold H_2O with decomp. (Brizard, A. ch. 1900, (7) 21. 349.)

Ruthenium dihydronitrosooxychloride,
 $\text{NO} \cdot \text{Ru}_2\text{H}_2\text{Cl}_2\text{OH} + 2\text{H}_2\text{O}.$

Ppt. (Brizard, A. ch. 1900, (7) 21. 349.)

Ruthenium monoxide, RuO .

Insol. in acids. (Claus, A. 59. 236.)

Ruthenium sesquioxide, Ru_2O_3 .

Insol. in acids. Mixture of Ru and RuO_2 . (Debray and Joly, C. R. 106. 1494.)

See Ruthenium nitrosos sesquioxide.

Ruthenium dioxide, RuO_2 .

Insol. in acids. (Debray and Joly.)

Ruthenium trioxide, RuO_3 .

"Ruthenic acid." Known only in its salts.

Ruthenium tetroxide, RuO_4 .

Rather difficultly and slowly sol. in H_2O . (Claus.)

Decomp. in aqueous solution into $\text{Ru}_2\text{O}_7 + 2\text{H}_2\text{O}$. (Debray and Joly.)

Ruthenium pentoxide, Ru_2O_5 .

(Debray and Joly, C. R. 106. 1494.)

+ $2\text{H}_2\text{O}$. Ppt. (Debray and Joly.)

Ruthenium heptoxide, Ru_2O_7 .

"Perruthenic acid." Known only in its salts.

Ruthenium oxide, Ru_4O_9 .

(Debray and Joly.)

Ruthenium oxychloride, $\text{Ru}(\text{OH})\text{Cl}_2$.

Very sol. in H_2O , but decomp. by an excess. (Joly, C. R. 114. 293.)

Ruthenium silicide, RuSi .

Insol. in boiling acids; slowly attacked by a mixture of fused KHSO_4 and KNO_3 . (Moissan, C. R. 1903, 137. 231.)

Ruthenium trisulphide, RuS_3 .

Ppt. (Antony, Gazz. ch. it. 1900, 30. 539.)

Ruthenomonamine hydroxide,

$\text{Ru}(\text{OH})_2(\text{NH}_3)_2 + 4\text{H}_2\text{O}.$

See Ruthenosamine hydroxide.

Ruthenodiamine carbonate,

$\text{Ru}(\text{N}_2\text{H}_4)_2\text{CO}_3 + 5\text{H}_2\text{O}.$

Easily sol. in H_2O . Insol. in alcohol. (Claus.)

Ruthenodiamine chloride,

$\text{Ru}(\text{N}_2\text{H}_4\text{Cl})_2 + 3\text{H}_2\text{O}.$

Not very sol. in cold, easily sol. in hot
 Insol. in alcohol.

See Ruthenonitrosodiamine comp.

— mercuric chloride, $\text{Ru}(\text{N}_2\text{H}_4\text{Cl})_2, \text{Hg}$

Nearly insol. in cold, sol. in hot
 (Gibbs, Sill. Am. J. (2) 34. 350.)

— chloroplatinate, $\text{Ru}(\text{N}_2\text{H}_4\text{Cl})_2, \text{Pt}$

Sl. sol. in H_2O . (Claus.)

— hydroxide, $\text{Ru}(\text{N}_2\text{H}_4\text{OH})_2$.

Known only in aqueous solution.

— nitrate, $\text{Ru}(\text{N}_2\text{H}_4\text{NO}_3)_2 + 2\text{H}_2\text{O}.$

Somewhat difficultly sol. in cold, eas
 hot H_2O . Insol. in alcohol.

— sulphate, $\text{Ru}(\text{N}_2\text{H}_4)_2\text{SO}_4 + 4\text{H}_2\text{O}.$

Moderately sol. in H_2O . Insol. in alc
 (Claus.)

Ruthenocyanhydric acid, $\text{H}_4\text{Ru}(\text{CN})_6$.

Easily sol. in H_2O and alcohol. Less
 in ether. (Claus, J. B. 1855. 444.)

**Potassium ruthenocyanide, $\text{K}_4\text{Ru}(\text{CN})_6$
 $3\text{H}_2\text{O}.$**

Sl. efflorescent. Very sol. in H_2O ; sl
 in dil. alcohol. (Claus.)

Ruthenonitrosodiamine bromide,

$\text{Ru}(\text{NO})(\text{NH}_3)_4\text{Br}_2.$

Sl. sol. in H_2O . (Joly, C. R. 111. 969.)

$\text{Ru}(\text{NO})\text{OH}(\text{NH}_3)_4\text{Br}_2$. Less sol. than
 corresponding chloride. (Joly, C. R. 108. 31)

— chloride, $\text{Ru}(\text{NO})(\text{NH}_3)_4\text{Cl}_2.$

Sl. sol. in H_2O . (Joly, C. R. 111. 969.)

$\text{Ru}(\text{NO})\text{OH}(\text{NH}_3)_4\text{Cl}_2$. Sol. in H
 (Joly, C. R. 108. 1300.)

$\text{Ru}(\text{NO})(\text{NH}_3)_4\text{Cl}_2 + 2\text{H}_2\text{O} =$
 $\text{Ru}(\text{NO})(\text{OH})(\text{NH}_3)_4\text{Cl}_2, \text{HCl} + \text{H}_2\text{O}$
 Very sol. in H_2O . (Joly, C. R. 111. 969.)

— chloroplatinate,

$\text{Ru}(\text{NO})\text{OH}(\text{NH}_3)_4\text{PtCl}_6.$

Scarcely sol. in boiling H_2O . (Joly, C.
 108. 1300.)

$\text{Ru}(\text{NO})(\text{NH}_3)_4\text{Cl}_2, \text{PtCl}_6$. Ppt. (C.
 R. 111. 969.)

— iodide, $\text{Ru}(\text{NO})(\text{NH}_3)_4\text{I}_2.$

Sl. sol. in H_2O . (Joly, C. R. 111. 969.)

$\text{Ru}(\text{NO})\text{OH}(\text{NH}_3)_4\text{I}_2$. Less sol. than
 corresponding bromide. (Joly, C. R.
 1300.)

— nitrate, $\text{Ru}(\text{NO})(\text{NH}_3)_4(\text{NO}_3)_2.$

More sol. in H_2O than
 $\text{Ru}(\text{NO})(\text{OH})(\text{NH}_3)_4(\text{NO}_3)_2$. (Joly, C.
 111. 969.)

$\text{OH}(\text{NH}_4)_4(\text{NO}_3)_2$. Sl. sol. in cold in conc. $\text{HNO}_3 + \text{Aq.}$ (Joly, C. R.

rosodiamine sulphate,
 $\text{O}(\text{NH}_4)_4(\text{SO}_4)_2 + 10\text{H}_2\text{O}$.

H_2O . (Joly, C. R. 111. 969.)
 $(\text{NH}_4)_4(\text{SO}_4)_2, \text{H}_2\text{SO}_4 + \text{H}_2\text{O}$. De-
 cold H_2O . (Joly.)
 $(\text{OH})_4(\text{NH}_4)_4\text{SO}_4 + \text{H}_2\text{O}$. Most sol.
 this class of salts. (Joly, C. R.

itrous acid.

1 ruthenonitrite, $\text{RuH}_2(\text{NO}_2)_4$,
 $\text{NO}_2 + 2\text{H}_2\text{O}$.

sol. in H_2O ; practically insol. in
 (Brizard, A. ch. 1900, (7) 21.

ruthenonitrite,
 $(\text{NO}_2)_{12} = 6\text{KNO}_2, \text{Ru}_2(\text{NO}_2)_3$.
 sol. in H_2O , alcohol, or ether.
 l. Am. J. (2), 34. 344.)
 n H_2O . Easily sol. in $\text{KNO}_2 + \text{Aq.}$

$(\text{O}_2)_{10} = \text{Ru}_2\text{O}_2(\text{N}_2\text{O}_2)_2, 4\text{KNO}_2$.
 1 H_2O . (Joly and Vèzes, C. R. 109.

$(\text{O}_2)_{14} = \text{Ru}_2\text{O}_2(\text{N}_2\text{O}_2)_2, 8\text{KNO}_2$. Sl.
 O. Sol. in cold dil. acids. (Joly
)
 $(\text{O}_2)_4, 3\text{KNO}_2 + 4\text{H}_2\text{O}$. Very sol.
 Aqueous solution decomp. sl. on
 ig. Almost insol. in conc. $\text{KCl} +$
 ard, C. R. 1899, 129. 216.)

enonitrite, $\text{NO.Ru}_2\text{H}_2(\text{NO}_2)_4$,
 $\text{O}_2 + 2\text{H}_2\text{O}$.
 Brizard, A. ch. 1900, (7) 21. 368.)

thenonitrite, $\text{Ru}_2(\text{NO}_2)_4, 4\text{NaNO}_2 +$
 l. in H_2O without decomp. (Joly,
 t, 118. 469.)

samine hydroxide,
 $\text{H}_2\text{OH})_2 + 4\text{H}_2\text{O}$ (?).
 liquescent, and sol. in H_2O . (Claus.)

n, Sm.
 ment has not been isolated.

bromide, $\text{SmBr}_3 + 6\text{H}_2\text{O}$.
 liquescent. (Cleve.)

carbide, SmC_2 .
 by water and acids. (Moissan,
 0, 131. 925.)

Samarium dichloride, SmCl_2 .

Decomp. by H_2O with liberation of H_2 and
 formation of samarium oxide and samarium
 oxychloride. Insol. in CS_2 , CHCl_3 , benzene,
 abs. alcohol, pyridine and toluene. (Matig-
 non, C. R. 1906, 142. 85.)

Samarium trichloride, SmCl_3 .

The anhydrous salt is very hygroscopic
 and easily sol. in H_2O . (Matignon, C. R.
 1902, 134. 1309.)

Very sol. in H_2O . Very sol. in abs. alcohol.
 6.38 g. are sol. in 100 grams pyridine at ord.
 temp.; insol. in quinoline. (Matignon, A. ch.
 1906, (8) 8. 406.)
 $+3\text{H}_2\text{O}$. Deliquescent.

Samarium chloride ammonia, $\text{SmCl}_3 + \text{NH}_3$;
 $+2\text{NH}_3$; $+3\text{NH}_3$; $+4\text{NH}_3$; $+5\text{NH}_3$;
 $+8\text{NH}_3$; $+9.5\text{NH}_3$; $+11.5\text{NH}_3$.
 (Matignon, C. R. 1905, 140. 143.)

Samarium fluoride, $\text{SmF}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Precipitate. Insol. in H_2O and dil. acids.
 (Cleve.)

Samarium iodide, SmI_3 .

(Matignon, A. ch. 1906, (8) 8. 413.)

Samarium hydroxide, $\text{Sm}_2(\text{OH})_6$.

Insol. in alkalis; easily sol. in acids, and
 decomposes ammonium salts. (Cleve, C. N.
 51. 145.)

Samarium oxide, Sm_2O_3 .

Easily sol. in acids. (Cleve, C. N. 51. 145.)

Samarium peroxide, Sm_4O_7 .

Precipitate. (Cleve.)

Samarium oxychloride, SmOCl .

(Matignon, A. ch. 1906, (8) 8. 412.)

Samarium sulphide, Sm_2S_3 .

(Matignon, A. ch. 1906, (8) 8. 415.)

Scandium, Sc.

Element has not been isolated.

**Scandium bromide, $\text{Sc}_2\text{Br}_6 + 3\text{H}_2\text{O}$, and
 $+12\text{H}_2\text{O}$.**

(Crookes, Roy. Soc. Proc. 1908, 80. A, 518.)

**Scandium chloride, $\text{Sc}_2\text{Cl}_6 + 3\text{H}_2\text{O}$, and
 $+12\text{H}_2\text{O}$.**

(Crookes, Roy. Soc. Proc. 1908, 80. A, 518.)

Scandium hydroxide.

Easily sol. in conc. HNO_3 or $\text{HCl} + \text{Aq.}$
 (Crookes, Roy. Soc. Proc. 1908, 80. A, 518.)

Scandium oxide, Sc_2O_3 .

Easily sol. by boiling with conc. HNO_3 or $\text{HCl} + \text{Aq}$.

Scandium sulphide, Sc_2S_3 .

Decomp. by H_2O and by acids with evolution of H_2S . (Wirth, Z. anorg. 1914, 87. 5.)

Selenantimonic acid.**Sodium selenantimonate, $\text{Na}_3\text{SbSe}_4 + 9\text{H}_2\text{O}$.**

Sol. in 2 pts. cold H_2O . Insol. in alcohol. (Hofacker, A. 107. 6.)

Sl. sol. in H_2O ; unstable. (Pouget, A. ch. 1899, (7) 18. 562.)

Selenantimonous acid.**Potassium orthoselenantimonite, K_2SbSe_3 .**

Ppt. Decomp. by H_2O . (Pouget, A. ch. 1899, (7) 18. 560.)

Potassium paraselenantimonite, $\text{K}_2\text{Sb}_2\text{Se}_7 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O ; unstable. (Pouget, A. ch. 1899, (7) 18. 560.)

Sodium orthoselenantimonite, $\text{Na}_2\text{SbSe}_3 + 9\text{H}_2\text{O}$.

Very sol. in H_2O . Aqueous solution on standing deposits red crystals of sodium selenantimonate, $\text{Na}_3\text{SbSe}_4 + 9\text{H}_2\text{O}$. (Pouget, A. ch. 1899, (7) 18. 562.)

Sodium paraselenantimonite, $\text{Na}_2\text{Sb}_2\text{Se}_7$.

(Pouget, A. ch. 1899, (7) 18. 561.)

Selenic acid, H_2SeO_4 .

Very sol. in H_2O with evolution of heat.

If aqueous solution is evaporated at temp. of 165° , acid has 2.524 sp. gr., at temp. of 207° , acid has 2.60 sp. gr., at temp. of 285° , acid has 2.625 sp. gr. Decomp. to H_2SeO_3 at higher temp. (Mitscherlich, Pogg. 9. 623.)

By evaporation at 265° , acid of 2.600 sp. gr. containing 95% H_2SeO_4 is obtained. If brought at same temp. in vacuo over H_2SO_4 , acid of 2.627 sp. gr. with 97.5% H_2SeO_4 is obtained. Fabian, A. Suppl. 1. 243.)

Sp. gr. of $\text{H}_2\text{SeO}_4 + \text{Aq}$.

% H_2SeO_4	Sp. gr.	% H_2SeO_4	Sp. gr.
99.74	2.6083	90.0	2.3848
99.50	2.6051	89.0	2.3768
99.00	2.6025	88.0	2.3691
98.5	2.5993	87.0	2.3611
98.0	2.5967	86.0	2.3535
97.5	2.5935	85.0	2.3458
97.0	2.5901	84.0	2.3378
96.0	2.5888	83.0	2.3296
95.0	2.5863	82.0	2.3217
94.0	2.5825	81.0	2.3140
93.0	2.5796	80.0	2.3066
92.0	2.5722	79.0	2.2992
91.0	2.5681	78.50	2.29675

Cameron and Macdunn, Lond. R. Soc. Proc. 46. 14.)

Sp. gr. of $\text{H}_2\text{SeO}_4 + \text{Aq}$ at 20° compared with H_2O at 4° .
Wts. corrected to vacuum.

Sp. gr.	% H_2SeO_4	Sp. gr.	% H_2SeO_4	Sp. gr.	% H_2SeO_4
1.000	...	1.295	32.64	1.590	54.66
1.005	0.9	1.300	33.06	1.595	55.08
1.010	1.56	1.305	33.50	1.600	55.50
1.015	2.12	1.310	33.92	1.605	55.92
1.020	2.92	1.315	34.36	1.610	56.34
1.025	3.62	1.320	34.82	1.615	56.76
1.030	4.16	1.325	35.26	1.620	57.18
1.035	4.70	1.330	35.72	1.625	57.60
1.040	5.32	1.335	36.10	1.630	58.02
1.045	6.08	1.340	36.43	1.635	58.44
1.050	6.66	1.345	36.88	1.640	58.86
1.055	7.34	1.350	37.34	1.645	59.28
1.060	7.92	1.355	37.80	1.650	59.70
1.065	8.56	1.360	38.24	1.655	60.12
1.070	9.20	1.365	38.66	1.660	60.54
1.075	9.82	1.370	39.10	1.665	60.96
1.080	10.44	1.375	39.50	1.670	61.38
1.085	11.02	1.380	39.98	1.675	61.80
1.090	11.62	1.385	40.06	1.680	62.22
1.095	12.20	1.390	40.66	1.685	62.64
1.100	12.88	1.395	41.10	1.690	63.06
1.105	13.58	1.400	41.56	1.695	63.48
1.110	14.14	1.405	41.96	1.700	63.90
1.115	14.66	1.410	42.36	1.705	64.32
1.120	15.20	1.415	42.78	1.710	64.74
1.125	15.74	1.420	43.16	1.715	65.16
1.130	16.32	1.425	43.56	1.720	65.58
1.135	16.86	1.430	43.94	1.725	66.00
1.140	17.38	1.435	44.32	1.730	66.42
1.145	17.90	1.440	44.52	1.735	66.84
1.150	18.44	1.445	45.00	1.740	67.26
1.155	18.92	1.450	45.32	1.745	67.68
1.160	19.48	1.455	45.68	1.750	68.10
1.165	20.02	1.460	46.04	1.755	68.52
1.170	20.58	1.465	46.36	1.760	68.94
1.175	21.08	1.470	46.70	1.765	69.36
1.180	21.60	1.475	47.01	1.770	69.78
1.185	22.22	1.480	47.32	1.775	70.20
1.190	22.66	1.485	47.66	1.780	70.62
1.195	23.18	1.490	47.98	1.785	71.04
1.200	23.70	1.495	48.28	1.790	71.46
1.205	24.26	1.500	48.54	1.795	71.88
1.210	24.84	1.505	48.92	1.800	72.30
1.215	25.30	1.510	49.30	1.805	72.72
1.220	25.84	1.515	49.68	1.810	73.14
1.225	26.30	1.520	50.02	1.815	73.56
1.230	26.84	1.525	50.34	1.820	73.98
1.235	27.28	1.530	50.68	1.825	74.40
1.240	27.70	1.535	51.04	1.830	74.82
1.245	28.18	1.540	51.38	1.835	75.24
1.250	28.58	1.545	51.66	1.840	75.66
1.255	29.03	1.550	51.96	1.845	76.08
1.260	29.44	1.555	52.28	1.850	76.50
1.265	29.82	1.560	52.56	1.855	76.92
1.270	30.26	1.565	52.88	1.860	77.34
1.275	30.76	1.570	53.28	1.865	77.76
1.280	31.26	1.575	53.56	1.870	78.18
1.285	31.74	1.580	53.94	1.875	78.60
1.290	32.18	1.585	54.30	1.880	79.02

$\text{O}_4 + \text{Aq}$ —Concluded.

gr.	H_2SeO_4	Sp. gr.	H_2SeO_4
25	80.25	2.365	89.14
30	80.42	2.370	89.30
35	80.68	2.375	89.48
40	80.74	2.380	89.60
45	80.96	2.385	89.72
50	81.14	2.390	89.84
55	81.36	2.395	89.96
60	81.60	2.400	90.10
65	81.80	2.405	90.20
70	82.02	2.410	90.30
75	82.22	2.415	90.46
80	82.44	2.420	90.74
85	82.64	2.425	91.00
90	82.78	2.430	91.24
95	82.96	2.435	91.46
00	83.10	2.440	91.70
05	83.24	2.445	92.00
10	83.44	2.450	92.28
15	83.62	2.455	92.56
20	83.78	2.460	92.85
25	83.96	2.465	93.02
30	84.14	2.470	93.20
35	84.30	2.475	93.36
40	84.48	2.480	93.68
45	84.60	2.485	94.02
50	84.82	2.490	94.32
55	85.02	2.495	94.48
60	85.26	2.500	94.64
65	85.44	2.505	94.80
70	85.60	2.510	94.96
75	85.78	2.515	95.32
80	85.96	2.520	95.58
85	86.16	2.525	95.86
90	86.38	2.530	96.10
95	86.60	2.535	96.41
00	86.82	2.540	96.68
05	87.04	2.545	96.92
10	87.26	2.550	97.12
15	87.46	2.555	97.30
20	87.66	2.560	97.48
25	87.84	2.565	97.68
30	88.00	2.570	97.94
35	88.18	2.575	98.20
40	88.34	2.580	98.46
45	88.48	2.585	98.70
50	88.66	2.590	99.04
55	88.82	2.595	99.36
60	88.98		

ver. J. phys. Chem. 1909,
2, 509.)

aming H_2SO_4
 NH_4 . (Franklin, Am. Ch

hol.
on and Macallan, C. N.

H_2O (?). (C and M)
ann and Hofmeier, M.

Selenates.

All the neutral and acid salts of H_2SeO_4 are sol. in H_2O , except BaSeO_4 , SrSeO_4 , CaSeO_4 , and PbSeO_4 , which are nearly or quite insol. in H_2O or $\text{HNO}_3 + \text{Aq}$.

Aluminum selenate, $\text{Al}_2(\text{SeO}_4)_3$.

Resembles in every way aluminum sulphate. (Berselius.)

Aluminum ammonium selenate,



More sol. in H_2O than the corresponding sulphate. (Wohlwill, A. 114. 191.)

Aluminum caesium selenate, $\text{Al}_2\text{Cs}_2(\text{SeO}_4)_4 + 24\text{H}_2\text{O}$.

(Peterson, B. 9. 1563.)

Much more sol. in H_2O than the corresponding sulphate. (Fabre, C. R. 106. 114.)

Aluminum potassium selenate, $\text{Al}_2\text{K}_2(\text{SeO}_4)_4 + 24\text{H}_2\text{O}$.

More sol. in H_2O than common alum. (Weber, Pogg. 108. 615.)

Aluminum rubidium selenate, $\text{Al}_2\text{Rb}_2(\text{SeO}_4)_4 + 24\text{H}_2\text{O}$.

(Peterson, B. 9. 1563.)

Much more sol. in H_2O than the corresponding sulphate. (Fabre, C. R. 106. 114.)

Aluminum sodium selenate, $\text{Al}_2\text{Na}_2(\text{SeO}_4)_4 + 24\text{H}_2\text{O}$.

Sl. efflorescent. Very sol. in H_2O . (Wohlwill, A. 114. 191.)

Aluminum thallium sulphate, $\text{Al}_2\text{Tl}_2(\text{SeO}_4)_4 + 24\text{H}_2\text{O}$.

Sol. in H_2O . (Fabre, C. R. 106. 114.)

Aluminum selenate potassium sulphate, $\text{Al}_2(\text{SeO}_4)_3, \text{K}_2\text{SO}_4 + 24\text{H}_2\text{O}$.

Sol. in H_2O . (v. Gerichten, A. 168. 222.)

Ammonium selenate, $(\text{NH}_4)_2\text{SeO}_4$.

Easily sol. in H_2O .

100 g. H_2O dissolve 117 g. $(\text{NH}_4)_2\text{SeO}_4$ at 7° ; 164 g. at 59° ; 197 g. at 100° . (Tutton, Proc. Roy. Soc. 1907, 79, A. 351.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 826.)

Ammonium hydrogen selenate, NH_4HSeO_4 .

Sol. in H_2O . (Topsoë.)

Ammonium cadmium selenate, $(\text{NH}_4)_2\text{SeO}_4, \text{CdSeO}_4 + 2\text{H}_2\text{O}$.

Sol. in H_2O (Topsoë, W. A. B. 88, 2. 2.)
+ $6\text{H}_2\text{O}$ Efflorescent. Very easily sol. in H_2O . (Topsoë.)

Ammonium cerous selenate,
 $(\text{NH}_4)_2\text{Ce}_2(\text{SeO}_4)_4 + 9\text{H}_2\text{O}$.

Easily sol. in H_2O . (Jolin.)

Ammonium chromium selenate,
 $(\text{NH}_4)_2\text{Cr}_2(\text{SeO}_4)_4 + 24\text{H}_2\text{O}$.

Sol. in H_2O . (Fabre, C. R. 105. 114.)

Ammonium cobaltous selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{CoSeO}_4 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Topsoë.)

Ammonium cupric selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{CuSeO}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Topsoë.)

Ammonium didymium selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{Di}_2(\text{SeO}_4)_3 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Cleve.)

+10 H_2O . (Cleve, Bull. Soc. (2) 43. 363.)

Ammonium erbium selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{Er}_2(\text{SeO}_4)_3 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O . (Cleve.)

Ammonium ferrous selenate, $(\text{NH}_4)_2\text{Fe}(\text{SeO}_4)_2$,
+6 H_2O .

Easily sol. in H_2O . (Topsoë.)

+2 H_2O .

Ammonium lanthanum selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{La}_2(\text{SeO}_4)_3 + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve.)

Ammonium magnesium selenate,
 $(\text{NH}_4)_2\text{Mg}(\text{SeO}_4)_2 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Topsoë.)

Ammonium manganous selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{MnSeO}_4 + 6\text{H}_2\text{O}$.

Not deliquescent. Easily sol. in H_2O .
(Topsoë.)

Ammonium nickel selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{NiSeO}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Topsoë.)

Ammonium samarium selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{Sm}_2(\text{SeO}_4)_3 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Cleve.)

Ammonium thallium selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{Tl}_2(\text{SeO}_4)_3 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Fortini, C. C. 1903, II. 706.)

Ammonium uranyl selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $(\text{UO}_2)_2\text{SeO}_4 + 2\text{H}_2\text{O}$.

Easily sol. in H_2O . (Sendtner.)

Ammonium yttrium selenate, $(\text{NH}_4)_2\text{SeO}_4$,
 $\text{Y}_2(\text{SeO}_4)_3 + 6\text{H}_2\text{O}$.

Very sol. in H_2O . (Cleve.)

Ammonium zinc selenate,
 $(\text{NH}_4)_2\text{SeO}_4$, $\text{ZnSeO}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Topsoë.)

Antimony selenate.

Insol. in H_2O . Not very sol. in acid
in H_2SeO_4 . (Cameron and Macallan.)

Barium selenate, BaSeO_4 .

Somewhat more sol. in H_2O and d
than BaSO_4 . (Rose.) 100 ccm. H_2O
11.8 mg. in the cold, and 13.8 mg.
(Petersson, Z. anal. 12. 287.)

Not decomp. by H_2SO_4 . Insol. in
Aq (Berzelius), but decomp. by sol
alkali carbonates at ordinary temp.

Very slowly decomp. by $\text{HCl} + \text{Aq}$.
Pogg. 95. 426.)

Bismuth selenate.

Insol. in, and not decomp. by cold
 H_2O . (Cameron and Macallan.)

Cæsium selenate, Cs_2SeO_4 .

Sol. in H_2O . (Petersson, B. 9. 15
100 g. H_2O at 12° dissolve 244.8 g.
(Tutton, Chem. Soc. 1897, 71. 850.)

Sp. gr. of $\text{Cs}_2\text{SeO}_4 + \text{Aq}$ at 20°
with H_2O at 4°, containing:

% Cs_2SeO_4	45.94	51
Sp. gr.	1.5841	1.5

(Tutton.)

Cæsium hydrogen selenate, CaHSel

Ppt. Very hygroscopic. (Norris,
J. 1901, 26. 322.)

Cæsium chromic selenate, Cs_2Cr_2 ,
24 H_2O .

Sol. in H_2O . (Fabre, C. R. 105.

Cæsium cobaltous selenate, Cs_2Co ,
6 H_2O .

Sol. in H_2O . (Topsoë.)

Cæsium indium selenate, CaI ,
12 H_2O .

Efflorescent; sol. in H_2O . (M.
Am. Chem. Soc. 1908, 30. 215.)

Cæsium iron (ferric) selenate, Ca_2I ,
+24 H_2O .

Sl. sol. in H_2O . (Roncogliolo,
it. 1905, 35. (2) 553.)

Cæsium magnesium selenate,
 $\text{MgSeO}_4 + 6\text{H}_2\text{O}$.

(Tutton, Chem. Soc. 1905, 87,

Cæsium zinc selenate, $\text{Ca}_2\text{Zn}(\text{SeO}_4)_2$

(Tutton, Zeit. Kryst. 1900, 23. 1.

selenate, $\text{CdSeO}_4 + 2\text{H}_2\text{O}$.
in H_2O . (v. Hauer, W. A. B. 39.)

potassium selenate, $\text{CdSeO}_4, \text{K}_2\text{SO}_4$
 O .
 H_2O ; can be recrystallized without
v. Hauer, W. A. B. 54. 209.)

selenate, $\text{CaSeO}_4 + 2\text{H}_2\text{O}$.
in hot than in cold H_2O . (v.
or. 80. 214.)
 $\text{eO}_4 + \text{Aq}$ contains at:
5° 20° 37° 67°
3 7.6 6.8 5.1% CaSeO_4 .
rd, A. ch. 1894, (7) 2. 551.)

selenate, $\text{Ce}_2(\text{SO}_4)_3 + 6\text{H}_2\text{O}, 9\text{H}_2\text{O}$, or
.
in cold than hot H_2O . (Jolin.)
Very easily in cold, sl. sol. in hot
golani, C. C. 1908, I. 1606.)
bove 100°.
Stable at 92–100°.
Stable at 80–92°.
Stable at 50–78°.
Stable at 34–40°.
Stable at 12–28°.
Stable at 0–12°. (Cingolani, C.
58.)

ity of $\text{Ce}_2(\text{SeO}_4)_3$ in H_2O at t° .
culated as anhydrous $\text{Ce}_2(\text{SO}_4)_3$
issolved in 100 cc. H_2O .)

	Using $\text{Ce}_2(\text{SeO}_4)_3$ + 4 H_2O	t°	Using $\text{Ce}_2(\text{SeO}_4)_3$ + 11 H_2O	Using $\text{Ce}_2(\text{SeO}_4)_3$ + 4 H_2O
	39.55	60°	13.68	
		60.8		13.12
	36.9	78.2	5.52	
4		80.5		4.56
	33.2	91	2.02	
5		95.4	1.53	
6		98		1.785
	31.89	100		2.513

(Cingolani, l. c.)

potassium selenate, $\text{Ce}_2(\text{SeO}_4)_3$
 O_4 .
in H_2O than the corresponding
(Jolin.)

ium selenate, $\text{Ce}_2(\text{SeO}_4)_3, \text{Na}_2\text{SeO}_4$
 O .
in H_2O . (Jolin.)

tassium selenate, $\text{Cr}_2\text{K}_2(\text{SeO}_4)_4 +$
s the sulphate in every particular.

Chromic rubidium selenate, $\text{Cr}_2\text{Rb}_2(\text{SeO}_4)_4 +$
 $24\text{H}_2\text{O}$.
Sol. in H_2O .

Chromic sodium selenate, $\text{Cr}_2\text{Na}_2(\text{SeO}_4)_4 +$
 $24\text{H}_2\text{O}$.
Sol. in H_2O . (Fabre, C. R. 105. 114.)

Chromic thallous selenate, $\text{Cr}_2\text{Tl}_2(\text{SeO}_4)_4 +$
 $24\text{H}_2\text{O}$.
Sol. in H_2O . (Fabre, C. R. 105. 114.)

Chromic selenate potassium sulphate,
 $\text{Cr}_2(\text{SeO}_4)_3, \text{K}_2\text{SO}_4 + 24\text{H}_2\text{O}$.
Sol. in H_2O . (v. Gerichten.)

Cobaltous selenate, basic, $4\text{CoO}, 3\text{SeO}_3 +$
 H_2O .
Insol. in H_2O ; sol. in acids. (Bogdan, Bull.
Soc. (3) 9. 586.)
 $\text{Co}_2(\text{OH})_2(\text{SeO}_4)_3$. Insol. in H_2O . Sol. in
acids. (Bogdan, C. C. 1895. 630.)

Cobaltous selenate, $\text{CoSeO}_4 + 5\text{H}_2\text{O}$.
Easily sol. in H_2O . (Topsoë.)
+ 6 H_2O . Easily sol. in H_2O . (Topsoë.)
+ 7 H_2O . Efflorescent. Extremely sol. in
 H_2O . (Topsoë.)
+ 18 H_2O . Very unstable. (Copaux, A.
ch. 1905, (8) 6. 553.)

Cobaltous potassium selenate, $\text{CoSeO}_4,$
 $\text{K}_2\text{SeO}_4 + 6\text{H}_2\text{O}$.
More sol. in H_2O than corresponding sul-
phate. (v. Hauer, W. A. B. 39. 837.)

Cobaltous rubidium selenate, $\text{CoRb}_2(\text{SeO}_4)_3$
+ 6 H_2O .
Sol. in H_2O . (Topsoë.)

Cobaltous thallous selenate, $\text{CoTl}_2(\text{SeO}_4)_3 +$
6 H_2O .
Sol. in H_2O . (Topsoë.)

Cupric selenate, basic, $3\text{CuO}, 2\text{SeO}_3 + 4\text{H}_2\text{O}$.
Insol. in H_2O ; sol. in acids. (Bogdan, Bull.
Soc. (3) 9. 588.)
+ 5 H_2O . Sl. sol. in cold H_2O . (Metzner,
C. R. 1898, 127. 55.)

Cupric selenate, $\text{CuSeO}_4 + 5\text{H}_2\text{O}$.

Solubility in H_2O :—

257 g. salt in 1 l. sat. solution at 15°.

346 " " " 1 l. " " 35°.

435 " " " 1 l. " " 55°.

Aq. solution decomp. at 70°. (Metzner,
C. R. 1898, 127. 55.)

+ H_2O , and + 2 H_2O . (Metzner.)

Cupric hydrazine selenate, $\text{N}_2\text{H}_4.\text{H}_2\text{SeO}_4,$
 $\text{CuSeO}_4 + \frac{1}{2}\text{H}_2\text{O}$.

Decomp. in aq. solution. (Rimini, C. C.
1907, I. 86.)

- Cupric magnesium selenate**, $\text{CuMg}_2(\text{SeO}_4)_4 + 28\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill.)
- Cupric nickel selenate**, $\text{CuSeO}_4, \text{NiSeO}_4 + 14\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill.)
- Cupric potassium selenate**, $\text{CuSeO}_4, \text{K}_2\text{SeO}_4 + 6\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Topsoë.)
- Cupric zinc selenate**, $\text{CuZn}_2(\text{SeO}_4)_4 + 28\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill.)
- Cupric selenate ferrous sulphate**, $2\text{CuSeO}_4, 3\text{FeSO}_4 + 35\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill.)
- Cupric selenate magnesium sulphate**, $\text{CuSeO}_4, 3\text{MgSO}_4 + 28\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill.)
- Cupric selenate zinc sulphate**, $\text{CuSeO}_4, 3\text{ZnSO}_4 + 28\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill.)
- Didymium selenate**, $\text{Di}_2(\text{SeO}_4)_3 + 5\text{H}_2\text{O}$, and $6\text{H}_2\text{O}$.
Sol. in H_2O .
+ $8\text{H}_2\text{O}$. Easily sol. in H_2O . (Cleve.)
+ $10\text{H}_2\text{O}$. Sol. in H_2O . (Cleve.)
- Didymium potassium selenate**, $\text{Di}_2(\text{SeO}_4)_3, \text{K}_2\text{SeO}_4 + 9\text{H}_2\text{O}$.
Not deliquescent. Easily sol. in H_2O . (Cleve.)
- Didymium sodium selenate**, $\text{Di}_2(\text{SeO}_4)_3, \text{Na}_2\text{SeO}_4 + 4\text{H}_2\text{O}$.
Easily sol. in H_2O . (Cleve.)
- Dysprosium selenate**, $\text{Dy}_2(\text{SeO}_4)_3 + 8\text{H}_2\text{O}$.
Easily sol. in H_2O ; insol. in alcohol. (Jantsch, B. 1911, 44. 1275.)
- Erbium selenate**, $\text{Er}_2(\text{SeO}_4)_3 + 8\text{H}_2\text{O}$, and $9\text{H}_2\text{O}$.
Easily sol. in H_2O . (Topsoë.)
- Erbium potassium selenate**, $\text{Er}_2(\text{SeO}_4)_3, \text{K}_2\text{SeO}_4 + 8\text{H}_2\text{O}$.
Easily sol. in H_2O . (Cleve.)
- Gadolinium selenate**, $\text{Gd}_2(\text{SeO}_4)_3 + 10\text{H}_2\text{O}$.
Decomp. in the air. (Benedicks, Z. anorg. 1900, 22. 410.)
- Gadolinium potassium selenate**, $\text{Gd}_2(\text{SeO}_4)_3, 3\text{K}_2\text{SeO}_4 + 4\text{H}_2\text{O}$.
Sol. in H_2O . (Benedicks, Z. anorg. 1900, 22. 412.)
- Glucinum selenate**, $\text{GlSeO}_4 + 4\text{H}_2\text{O}$.
Very sol. in H_2O . (Atterberg.)
- Gold (auric) selenate**, $\text{Au}_2(\text{SeO}_4)_3$.
Insol. in H_2O . Sol. in hot conc. H Aq. Somewhat sol. in H_2SO_4 and Aq. Decomp. by $\text{HCl} + \text{Aq}$. (Lenba Chem. Soc. 1902, 24. 355.)
- Indium selenate**, $\text{In}_2(\text{SeO}_4)_3 + 10\text{H}_2\text{O}$.
Hydroscopic; easily sol. in H_2O . (J. Am. Chem. Soc. 1908, 30. 214.)
- Iron (ferrous) selenate**, $\text{FeSeO}_4 + 5\text{H}_2\text{O}$.
Sol. in H_2O . (Wohlwill, A. 114. 11) + $7\text{H}_2\text{O}$. Efflorescent, and sol. (Topsoë.)
- Iron (ferrous) potassium selenate**, $\text{K}_2\text{SeO}_4 + 6\text{H}_2\text{O}$.
Easily sol. in H_2O . Solution decom what on standing. (Topsoë.)
- Iron (ferric) rubidium selenate**, $\text{Rb}_2\text{Fe}_2(\text{SeO}_4)_6 + 24\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Roncogliolo, Ga 1905, 35. (2) 553.)
- Iron (ferric) selenate potassium**, $\text{Fe}_2(\text{SeO}_4)_3, \text{K}_2\text{SO}_4 + 24\text{H}_2\text{O}$.
Sol. in H_2O . (v. Gerichten.)
- Lanthanum selenate**, $\text{La}_2(\text{SeO}_4)_3 + 10\text{H}_2\text{O}$.
Easily sol. in cold H_2O . (Cleve.) + $12\text{H}_2\text{O}$. (Frerichs and Smith 355.)
- Lanthanum potassium selenate**, I $\text{K}_2\text{SeO}_4 + 9\text{H}_2\text{O}$.
Quite sol. in H_2O . (Cleve.)
- Lanthanum sodium selenate**, I $\text{Na}_2\text{SeO}_4 + 4\text{H}_2\text{O}$.
Easily sol. in H_2O . (Cleve.)
- Lead selenate, basic**, $2\text{PbO}, \text{SeO}_2$.
Decomp. by acids with sepa PbSeO_4 .
 $3\text{PbO}, \text{PbSeO}_4 + \text{H}_2\text{O}$. Ppt. (Z. anorg. 1904, 38. 443.)
- Lead selenate**, PbSeO_4 .
Insol. in H_2O or $\text{HNO}_3 + \text{Aq}$. (W. A. B. 47. 256.)
Min. *Kerstenite*.
- Lithium selenate**, $\text{Li}_2\text{SeO}_4 + \text{H}_2\text{O}$.
Not deliquescent. Easily sol. (Topsoë.)

- selenate, $\text{MgSeO}_4 + 6\text{H}_2\text{O}$.**
 resembles closely that of MgSO_4 .
- potassium selenate, $\text{MgK}_2(\text{SeO}_4)_2$.**
 in H_2O . (Topsoë.)
- rubidium selenate, $\text{MgSeO}_4 + 6\text{H}_2\text{O}$.**
 hem. Soc. 1905, 87. 1163.)
- selenate, $\text{MnSeO}_4 + 2\text{H}_2\text{O}$.**
 in H_2O . (Topsoë.)
 Easily sol. in H_2O . Solution
 warming or standing. (Topsoë.)
- potassium selenate, K_2SeO_4 .**
 uescent. Easily sol. in H_2O .
- selenate, $6\text{Hg}_2\text{O}, 5\text{SeO}_3$.**
 l. in H_2O . Sl. attacked by boil-
 Insol. in $\text{HCl} + \text{Aq}$. (Köhler,
 i.)
 Very sl. sol. in H_2O ; insol. in
 Cameron and Davy, C. N. 44.
- mate, basic, $6\text{HgO}, 2\text{SeO}_3 + \text{H}_2\text{O}$.**
 H_2O , or cold $\text{HNO}_3 + \text{Aq}$. Sol. in
 $\text{HCl} + \text{Aq}$. (Köhler.)
 HgO . Sol. in 10,330 pts. H_2O .
 id Davy.)
- enate, $\text{HgSeO}_4 + \text{H}_2\text{O}$.**
 by H_2O with formation of basic
 r.)
 SeO_4 , H_2SO_4 , HNO_3 , or $\text{HCl} + \text{Aq}$,
 by H_2O to 2HgO , HgSeO_4 .
 id Davy, C. N. 44. 63.)
- ate, $\text{NiSeO}_4 + 6\text{H}_2\text{O}$.**
 y sol. in H_2O . (v. Hauer, W. A.
- sium selenate, $\text{NiSeO}_4, \text{K}_2\text{SeO}_4 +$
 O . (Topsoë.)**
- ium selenate, $\text{NiSeO}_4, \text{Ti}_2\text{SeO}_4 +$
 O . (Petersson.)**
- lenate.**
 oiling H_2O . Sol. in $\text{HCl} + \text{Aq}$.
 xhol. (Cameron and Macallan,
 o. Proc. 46. 13.)
- Potassium selenate, K_2SeO_4 .**
 Nearly equally sol. in cold and hot H_2O .
 (Mitscherlich, Pogg. 9. 623.)
 100 g. H_2O dissolve 110.5 g. K_2SeO_4 at 0° ;
 112.8 g. at 20° ; 122.2 g. at 100° . (Étard,
 C. R. 1888, 106. 741.)
 Sat. $\text{K}_2\text{SeO}_4 + \text{Aq}$ contains at:
- | | | |
|-------------|------------|----------------------------------|
| -20° | -5° | $+5^\circ$ |
| 51.5 | 51.7 | 52.0% K_2SeO_4 , |
-
- | | |
|------------|----------------------------------|
| 18° | 97° |
| 52.6 | 54.9% K_2SeO_4 . |
- (Étard, A. ch. 1894, (7) 2. 550.)
- 100 g. H_2O at 12° dissolve 115.0 g. K_2SeO_4 .
 (Tutton, Chem. Soc. 1897, 71. 850.)
 Sp. gr. of $\text{K}_2\text{SeO}_4 + \text{Aq}$ at 20° compared
 with H_2O at 4° , containing:
- | | | | |
|----------------------------|--------|--------|--------|
| % K_2SeO_4 | 35.76 | 41.79 | 50.00 |
| Sp. gr. | 1.3591 | 1.4385 | 1.5590 |
- (Tutton, Chem. Soc. 1897, 71. 851.)
- Potassium hydrogen selenate, KHSeO_4 .**
 Sol. in H_2O .
- Potassium praseodymium selenate, $3\text{K}_2\text{SeO}_4$,
 $\text{Pr}_2(\text{SeO}_4)_3 + 4\text{H}_2\text{O}$.**
 Sl. sol. in H_2O . (von Scheele, Z. anorg.
 1898, 18. 361.)
- Potassium samarium selenate, K_2SeO_4 ,
 $\text{Sm}_2(\text{SeO}_4)_3 + 6\text{H}_2\text{O}$.**
 Easily sol. in H_2O . (Cleve, Bull. Soc. (2)
 43. 166.)
- Potassium sodium selenate, $3\text{K}_2\text{SeO}_4$,
 Na_2SeO_4 .**
 Sol. in H_2O . (Topsoë.)
- Potassium thallium selenate, K_2SeO_4 ,
 $\text{Tl}_2(\text{SeO}_4)_3 + 8\text{H}_2\text{O}$.**
 Very sol. in dil. acids. (Fortini, C. C. 1903,
 II. 706.)
- Potassium uranyl selenate, K_2SeO_4 ,
 $(\text{UO}_2)\text{SeO}_4 + 2\text{H}_2\text{O}$.**
 Sl. sol. in cold, easily in hot H_2O . (Sendt-
 ner.)
- Potassium yttrium selenate, K_2SeO_4 ,
 $\text{Y}_2(\text{SeO}_4)_3 + 6\text{H}_2\text{O}$.**
 Very sol. in H_2O . (Cleve.)
- Potassium zinc selenate, $\text{K}_2\text{SeO}_4, \text{ZnSeO}_4 +$
 $2\text{H}_2\text{O}$.**
 Sol. in H_2O . (Topsoë.)
 $+6\text{H}_2\text{O}$. Sol. in H_2O . (Topsoë.)
- Potassium selenate aluminum sulphate,
 $\text{K}_2\text{SeO}_4, \text{Al}_2(\text{SO}_4)_3 + 24\text{H}_2\text{O}$.**
 Sol. in H_2O . (v. Gerichten.)

Potassium selenate chromic sulphate,
 $K_2SeO_4, Cr_2(SO_4)_3 + 24H_2O$.
Sol. in H_2O . (v. Gerichten.)

Potassium selenate ferric sulphate, $K_2SeO_4, Fe_2(SO_4)_3 + 24H_2O$.
Sol. in H_2O . (v. Gerichten.)

Potassium selenate manganous sulphate,
 $K_2SeO_4, MnSO_4 + 6H_2O$.
Sol. in H_2O . (v. Gerichten, A. 168. 225.)

Potassium selenate manganic sulphate,
 $K_2SeO_4, Mn_2(SeO_4)_3 + 24H_2O$.
Sol. in H_2O . (v. Gerichten.)

Praseodymium selenate, $Pr_2(SeO_4)_3$.
Sol. in H_2O . (von Schule, Z. anorg. 1898, 18. 360.)
+8 H_2O . Sl. sol. in H_2O ; sol. in H_2SO_4 . (von Schule.)

Rubidium selenate, Rb_2SeO_4 .
Sol. in H_2O . (Petersson.)
100 g. H_2O at 12° dissolve 158.9 g. Rb_2SeO_4 . (Tutton, Chem. Soc. 1897, 71. 850.)
Sp. gr. of $Rb_2SeO_4 + Aq$ at 20° compared with H_2O at 4°, containing:
% Rb_2SeO_4 40.60 47.07
Sp. gr. 1.4688 1.5806
(Tutton.)

Rubidium hydrogen selenate, $RbHSeO_4$.
Sol. in equal pts. H_2O ; very hygroscopic. (Norris, Am. Ch. J. 1901, 26. 321.)

Rubidium zinc selenate, $Rb_2Zn(SeO_4)_2 + 6H_2O$.
(Tutton, Zeit. Kryst. 1900, 33. 8.)

Samarium selenate, $Sm_2(SeO_4)_3 + 8H_2O$.
More sol. in H_2O than $Sm_2(SO_4)_3$.
+12 H_2O . Efflorescent. (Cleve.)

Scandium selenate, $Sc_2(SeO_4)_3 + 2H_2O$, and +8 H_2O .
(Crookes, Roy. Soc. Proc. 1908, 80, A. 518.)

Silver selenate, Ag_2SeO_4 .
As Ag_2SO_4 . (Mitscherlich, Pogg. 12. 138.)

Silver selenate ammonia, $Ag_2SeO_4, 4NH_3$.
Easily sol. in H_2O or $NH_4OH + Aq$ without decomp. (Mitscherlich, Pogg. 12. 141.)

Sodium selenate, Na_2SeO_4 .
Very sol. in H_2O , forming supersat. solutions. Cryst. also with 10 H_2O , which effloresce. Maximum point of solubility is at 33°. (Mitscherlich.)

Solubility in H_2O at t°.			
t°	% Na_2SeO_4	Mols. H_2O to 1 mol. Na_2SeO_4	M. drs 100
35.2	45.47	12.50	
39.5	45.26	12.70	
50	44.49	13.10	
75	42.83	14.00	
100	42.14	14.42	

(Funk, B. 1900, 33. 3097.)

+10 H_2O . Solubility in H_2O at			
t°	% Na_2SeO_4	Mols. H_2O to 1 mol. Na_2SeO_4	M. drs 100
0	11.74	79.08	
15	25.01	31.48	
25.2	36.91	17.95	
27	39.18	16.30	
30	44.05	13.33	

(Funk.)

Sp. gr. of sat. solution at 1°
(Funk.)

Sodium selenate vanadate.
See Selenovanadate, sodium.

Strontium selenate, $SrSeO_4$.
Insol. in H_2O or $HNO_3 + Aq$; d long boiling with $HCl + Aq$.

Tellurium selenate, $2TeO_3, SeO_2$.
As sulphate. (Metzner, A. ch 15. 203.)

Thalious selenate, Tl_2SeO_4 .
Sl. sol. in cold, much more in Insol. in alcohol and ether. (Kuh 100 g. H_2O dissolve 2.13 g. at at 12°; 10.86 g. at 100°. (Tut Roy. Soc. 1907, 79. A, 351.)
2.8 g. are sol. in 100 g. H_2O at at 80°. (Glauser, Z. anorg. 1910,

Thalious hydrogen selenate, $3H_2O$.
(Oettinger.)

Thalious zinc selenate, $Tl_2SeO_4, 6H_2O$.
Easily sol. in H_2O , but less th responding sulphate. (Werther, 1865. 60.)

Thorium selenate, $Th(SeO_4)_4 + 9H_2O$
100 pts. H_2O dissolve 0.498 pt at 0°, and 1.972 pts. at 100°. (Cl

tannic) selenate, basic, $\text{SnO}(\text{SeO}_4) + \text{H}_2\text{O}$.
 quiescent. Sol. in H_2O . (Ditte, C. R. 11.)

selenate, $(\text{UO}_2)\text{SeO}_4, \text{H}_2\text{SeO}_4 + \text{H}_2\text{O}$.

deliquescent.

$(\text{UO}_2)\text{SeO}_4, \text{H}_2\text{SeO}_4 + 12\text{H}_2\text{O}$. Efflores-
 Sol. in H_2O . (Sendtner, A. 195. 325.)

ium selenate, $\text{Yb}_2(\text{SeO}_4)_3$.

hydrous.

H_2O (?), $+8\text{H}_2\text{O}$. Ppt. (Cleve, Z. 1902, 32. 145.)

n selenate, $\text{Y}_2(\text{SeO}_4)_3$.

hydrous. Sol. in H_2O with hissing and on of heat. (Popp.)

I_2O . Easily sol. in H_2O . (Cleve.)

I_2O . Efflorescent.

selenate, $\text{ZnSeO}_4 + 5\text{H}_2\text{O}$.

in H_2O . (Topsoë.)

I_2O . Sol. in H_2O . (Topsoë.)

I_2O . Sol. in H_2O .

ous acid, H_2SeO_3 .

quiescent in moist, efflorescent in dry very sol. in cold, and in nearly every tion in hot H_2O . Easily sol. in alcohol. (lius.)

of H_2SeO_3 and of $\text{H}_2\text{SeO}_3 + \text{Aq}$ at t° .
 Two series of experiments.

	t°	Sp. gr. at t°
$\text{H}_2\text{SeO}_3 + \text{Aq}$ (A)	18.0	1.4386
sol. A + 0.5 vol. H_2O	18.0	1.3179
" + 1.0 "	17.7	1.2337
" + 1.5 "	16.6	1.2045
" + 2.0 "	14.0	1.1984
" + 2.5 "	17.0	1.1712
" + 3.0 "	19.2	1.1600
$\text{H}_2\text{SeO}_3 + \text{Aq}$ (B)	15.8	1.4698
sol. B + 0.5 vol. H_2O	16.5	1.3191
" + 1.0 "	13.0	1.2515
" + 1.5 "	14.2	1.2074
" + 2.0 "	17.0	1.1992
" + 2.5 "	16.5	1.1793
" + 3.0 "	14.2	1.1678

le Coninck, C. C. 1905, I. 1693.)

also Selenium dioxide.

l. in liquid NH_3 . (Gore, Am. Ch. J. 10. 830.)

es.

li selenites are sol. in H_2O . The other selenites are insol. in H_2O , but sol. in $+ \text{Aq}$, Pb, and Ag salts slowly. The

neutral salts are insol. in $\text{HCl} + \text{Aq}$. The acid salts of the heavy metals are sol. in H_2O .

Aluminum selenite, basic, $4\text{Al}_2\text{O}_3, 9\text{SeO}_2 + 36\text{H}_2\text{O}$.

Precipitate. (Nilson, Upsala, 1875.)

Aluminum selenite, $\text{Al}_2(\text{SeO}_3)_3$.

Precipitate. (Berzelius.)

$+7\text{H}_2\text{O}$. Sl. sol. in H_2O . (Nilson.) Sol. in $\text{H}_2\text{SeO}_3 + \text{Aq}$.

$+3\text{H}_2\text{O}$. Insol. in H_2O ; sol. in acids. (Boutzoureano, A. ch. (6) 17. 289.)

Aluminum selenite, acid, $\text{Al}_2\text{O}_3, 4\text{SeO}_2 + 3\text{H}_2\text{O}$.

(Boutzoureano.)

$2\text{Al}_2\text{O}_3, 9\text{SeO}_2 + 12\text{H}_2\text{O}$. Sol. in H_2O . (Nilson.)

$\text{Al}_2\text{O}_3, 6\text{SeO}_2$. Very sol. in H_2O . (Berzelius.)

$+5\text{H}_2\text{O}$. (Nilson.)

$+2\text{H}_2\text{O}$. (Boutzoureano.)

Ammonium selenite, $(\text{NH}_4)_2\text{SeO}_3$.

Deliquescent. Very sol. in H_2O .

Precipitated from aqueous solution by alcohol. Insol. in ether. (Muspratt, A. 70. 275.)

Ammonium hydrogen selenite, NH_4HSeO_3 .

Not deliquescent. Sol. in H_2O . (Berzelius.)

Ammonium trihydrogen selenite,

$\text{NH}_4\text{H}_3(\text{SeO}_3)_2$.

Deliquescent. (Berzelius.)

Ammonium vanadium selenite.

See Vanadioselenite, ammonium.

Ammonium uranyl selenite, $(\text{NH}_4)_2\text{SeO}_3, (\text{UO}_2)\text{SeO}_3$.

Completely insol. in H_2O . (Sendtner.)

Antimony selenite, $\text{Sb}_2(\text{SeO}_3)_3, \text{SeO}_2$.

(Nilson, Bull. Soc. (2) 23. 494.)

Barium selenite, BaSeO_3 .

Sl. sol. in H_2O . Sol. in $\text{H}_2\text{SeO}_3 + \text{Aq}$. So in acids. (Nilson.)

$+ \text{H}_2\text{O}$. (Nilson.)

Barium pyroselenite, BaSe_2O_5 .

Very sl. sol. in cold, more in warm H_2O . (Berzelius.)

Bismuth selenite, $\text{Bi}_2(\text{SeO}_3)_3, \text{H}_2\text{SeO}_3$.

(Nilson.)

$\text{Bi}_2(\text{SeO}_3)_3$. (Nilson.)

Cadmium selenite, CdSeO_3 .

Insol. in H_2O . Sol. in $\text{H}_2\text{SeO}_3 + \text{Aq}$. (Muspratt, Chem. Soc. 2. 65.)

2CdO , $3\text{SeO}_2 + \text{H}_2\text{O}$. Insol. in H_2O ; sol. in acids. (Boutzoureano.)
 $+ \frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O ; sol. in dil. acids. (Boutzoureano.)

Cadmium selenite ammonia, $\text{CdSeO}_3 \cdot \text{NH}_3$.
 Insol. in cold or hot H_2O . (Boutzoureano, A. ch. (6) 17. 289.)

Calcium selenite, $\text{CaSeO}_3 + \frac{1}{2}\text{H}_2\text{O}$.
 Very sl. sol. in H_2O . (Berzelius.) More sol. in $\text{H}_2\text{SeO}_3 + \text{Aq}$.
 $+ 2\text{H}_2\text{O}$. (Nilson.)

Calcium hydrogen selenite, $\text{CaH}_2(\text{SeO}_3)_2 + \text{H}_2\text{O}$.
 Quite sol. in H_2O . (Nilson.)
 $\text{Ca}_2\text{H}_2\text{Se}_4\text{O}_{11}$. Easily sol. in H_2O . (Nilson.)

Cerous selenite, basic, $2\text{Ce}_2\text{O}_3$, $5\text{SeO}_2 + 30\text{H}_2\text{O}$.
 Precipitate. (Nilson.)

Cerous selenite, $\text{Ce}_2(\text{SeO}_3)_3 + 3\text{H}_2\text{O}$.
 Insol. in H_2O . Sol. in much selenious acid. (Jolin.)
 $+ 12\text{H}_2\text{O}$. (Nilson.)

Cerous selenite, acid, Ce_2O_3 , $4\text{SeO}_2 + 5$, or $6 \text{H}_2\text{O}$.
 Insol. in H_2O , but sol. in selenious, and other acids. (Jolin.)
 Ce_2O_3 , $6\text{SeO}_2 + 5\text{H}_2\text{O}$. Not decomp. by H_2O . (Nilson.)

Ceric selenite, $\text{Ce}(\text{SeO}_3)_2$.
 Insol. in H_2O .
 Sl. sol. in conc. HNO_3 . Sol. in dil. acids. Sol. in $\text{H}_2\text{O}_2 + \text{Aq}$. (Barbieri, B. 1910, 43. 2215.)

Chromium selenite, basic, $4\text{Cr}_2\text{O}_3$, $9\text{SeO}_2 + 64\text{H}_2\text{O}$.
 Precipitate. (Nilson.)

Chromic selenite, $\text{Cr}_2(\text{SeO}_3)_3 + 3\text{H}_2\text{O}$.
 (Boutzoureano.)
 $+ 15\text{H}_2\text{O}$. (Nilson.)
 Very sl. sol. or insol. in H_2O ; sl. sol. in $\text{H}_2\text{SeO}_3 + \text{Aq}$; sol. in hot conc. $\text{HCl} + \text{Aq}$. (Taquet, C. R. 96. 107.)

Chromic selenite, acid, Cr_2O_3 , $4\text{SeO}_2 + 13\text{H}_2\text{O}$.
 Slowly sol. in $\text{HCl} + \text{Aq}$. Insol. in H_2O . (Nilson.)
 Cr_2O_3 , $5\text{SeO}_2 + 9\text{H}_2\text{O}$. Insol. in H_2O . (Nilson.)

Chromic diselenite.
 Insol. in H_2O ; sol. in acids. (Taquet, C. R. 97. 1435.)

Cobaltous selenite, CoSeO_3 .
 Insol. in H_2O . (Berzelius.)
 $+ \frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O ; sol. in (Boutzoureano, A. ch. (6) 17. 289.)

Cobaltous hydrogen selenite, CoH_2 .
 Sol. in H_2O . (Berzelius.)
 $+ 2\text{H}_2\text{O}$. Sol. in H_2O with (Boutzoureano.)

Cuprous selenite.
 Insol. in H_2O . Sol. in NH_4 (Berzelius.)

Cupric selenite, basic, 2CuO , SeO_2 .
 Insol. in H_2O ; sol. in NH_4OH (Boutzoureano.)
 Sol. in acids.

Cupric selenite, $\text{CuSeO}_3 + \frac{1}{2}\text{H}_2\text{O}$.
 Insol. in H_2O or $\text{H}_2\text{SeO}_3 + \text{Aq}$. $+ \text{H}_2\text{O}$, and $2\text{H}_2\text{O}$. (Boutzoureano.)
 $+ 2\text{H}_2\text{O}$. Min. *Chalcomenite*. Insol. in $\text{H}_2\text{SeO}_3 + \text{Aq}$. (Friedel and Sar Kryst. 1881, 6. 300.)

Cupric selenite, acid, CuO , 2SeO_2 , $\text{CuH}_2(\text{SeO}_3)_2$.
 Insol. in H_2O . Sol. in acids. $+ 2\text{H}_2\text{O}$. As above. (Boutzoureano.)
 $+ 4\text{H}_2\text{O}$. As above. (B.)

Cupric selenite ammonia, $\text{CuSeO}_3 \cdot \text{H}_2\text{O}$.
 Decomp. by H_2O . (Boutzoureano, (6) 17. 289.)

Didymium selenite, basic, $3\text{Di}_2\text{O}_3$, $28\text{H}_2\text{O}$.
 Precipitate. (Nilson.)
 $+ 21\text{H}_2\text{O}$. Insol. in H_2O . (C. Soc. (2) 43. 363.)

Didymium selenite, $\text{Di}_2(\text{SeO}_3)_3 + 4\text{H}_2\text{O}$.
 Precipitate. (Smith.)

Didymium selenite, acid, Di_2O_3 , $5\text{H}_2\text{O}$.
 Precipitate. (Cleve.)
 Composition is $\text{Di}_2(\text{SeO}_3)_3 + 6\text{H}_2\text{O} + 9\text{H}_2\text{O}$. (Nilson.)
 $2\text{Di}_2\text{O}_3$, $9\text{SeO}_2 + 18\text{H}_2\text{O}$. (Nilson.)

Erbium selenite, $\text{Er}_2(\text{SeO}_3)_3 + 5\text{H}_2\text{O}$.
 Precipitate. (Nilson.)

Erbium hydrogen selenite, Er_2H_2 , $4\text{H}_2\text{O}$.
 Decomp. by hot H_2O .

1 hydrogen selenite,
 $(\text{O}_2)_3, \text{H}_2\text{SeO}_3 + 6\text{H}_2\text{O}.$

Benedicks, Z. anorg. 1900, 22.

selenite, basic, $5\text{GlO}, 2\text{SeO}_2 +$

te. (Nilson.) According to Atter-
 $\text{GlO}, 3\text{SeO}_2 + 14\text{H}_2\text{O}.$

$\text{O}_2 + 4\text{H}_2\text{O}.$ (Atterberg, Bull. Soc.

$2\text{SeO}_2 + 6\text{H}_2\text{O}.$ Insol. in $\text{H}_2\text{O}.$

selenite, $\text{GlSeO}_4 + 2\text{H}_2\text{O}.$

little H_2O , decomp. by excess.

selenite, acid.

$), 5\text{SeO}_2 + 3\text{H}_2\text{O}; (b) \text{GlO}, 2\text{SeO}_2,$
 $) 3\text{GlO}, 7\text{SeO}_2 + 5\text{H}_2\text{O}; (d) \text{GlO},$
 $\text{I}_2\text{O}.$ All are very sl. sol. in cold
 $\text{I}_2\text{O}.$ $a, b,$ and c are sol. in warm
 ining $\text{HCl}; d$ is sol. only in boiling
 Aq. (Nilson.)

lenite, basic, $\text{In}_2\text{Se}_2\text{O}_{10} + 64\text{H}_2\text{O}.$

lenite, $\text{In}_2(\text{SO}_3)_3 + 6\text{H}_2\text{O}.$

in $\text{H}_2\text{O}.$ (Nilson.)

hydrogen selenite, $\text{In}_2(\text{SeO}_3)_3,$
 $\text{O}_3 + 4\text{H}_2\text{O}.$

$\text{H}_2\text{O}.$ (Nilson.)

$), 3\text{H}_2\text{SeO}_3 + 12\text{H}_2\text{O}.$ Sol. in
 lson.)

ous) selenite.

sl. in $\text{HCl} + \text{Aq}$ with partial separa-
 (Berzelius.)

ous) hydrogen selenite.

in $\text{H}_2\text{O}.$ (Berzelius.)

c) selenite, basic, $2\text{Fe}_2\text{O}_3, 3\text{SeO}_2 +$

$\text{H}_2\text{O}.$ (Berzelius.)

$2\text{SeO}_2.$ Insol. in H_2O , easily sol. in
 boutzoureano, A. ch. (6) 17. 289.)

$8\text{SeO}_2 + 28\text{H}_2\text{O}.$ Insol. in $\text{H}_2\text{O}.$

ic) selenite, $\text{Fe}_2(\text{SeO}_3)_3 + 4\text{H}_2\text{O}.$

$1 \text{H}_2\text{O}.$ (Muspratt, Chem. Soc. 2.

Insol. in $\text{H}_2\text{O}.$ (Boutzoureano,
 17. 289.)

Insol. in $\text{H}_2\text{O}.$ (B.)

$).$ Insol. in $\text{H}_2\text{O}.$ (B.)

Iron (ferric) selenite, acid, $\text{Fe}_2\text{O}_3, 6\text{SeO}_2 +$
 $x\text{H}_2\text{O}.$

Insol. in $\text{H}_2\text{O}.$ Sol. in $\text{HCl} + \text{Aq}.$ (Ber-
 zelius.)

$+ 2\text{H}_2\text{O}.$ (Boutzoureano, A. ch. (6) 17.
 289.)

$\text{Fe}_2\text{O}_3, 4\text{SeO}_2 + \text{H}_2\text{O}.$ Insol. in $\text{H}_2\text{O};$ sol. in
 acids. (Boutzoureano.)

Lanthanum selenite, basic, $3\text{La}_2\text{O}_3, 8\text{SeO}_2 +$
 $28\text{H}_2\text{O}.$

Precipitate. (Nilson.)

Lanthanum selenite, $\text{La}_2(\text{SeO}_3)_3 + 9\text{H}_2\text{O},$ or
 $12\text{H}_2\text{O}.$

Insol. in $\text{H}_2\text{O}.$ (Nilson.)

Lanthanum selenite, acid, $\text{La}_2\text{H}_4(\text{SeO}_3)_3 +$
 $4\text{H}_2\text{O}.$

(Nilson.)

$\text{La}_2\text{H}_3(\text{SeO}_3)_3 + 2\text{H}_2\text{O}.$ (Cleve.)

Lead selenite, $\text{PbSeO}_3.$

Scarcely sol. in H_2O , even when it contains
 $\text{H}_2\text{SeO}_3.$ Sl. sol. in $\text{HNO}_3 + \text{Aq}.$ (Berzelius.)

Lithium selenite, $\text{Li}_2\text{SeO}_3 + \text{H}_2\text{O}.$

Difficultly sol. in $\text{H}_2\text{O}.$ (Nilson, Bull. Soc.
 (2) 21. 253.)

Lithium hydrogen selenite, $\text{LiHSO}_3.$

Very sol. in $\text{H}_2\text{O}.$ (Nilson.)

Lithium trihydrogen selenite, $\text{LiH}_3(\text{SeO}_3)_2.$

Not deliquescent. Sol. in $\text{H}_2\text{O}.$ (Nilson.)

Lithium vanadium selenite.

See Vanadioselenite, lithium.

Magnesium selenite, $\text{MgSeO}_3 + 2\text{H}_2\text{O}.$

Insol. in $\text{H}_2\text{O};$ sol. in dil. acids, especially
 if warm, also in $\text{H}_2\text{SeO}_3 + \text{Aq}.$ (Boutzour-
 eano, A. ch. (6) 18. 302.)

$+ 3\text{H}_2\text{O}.$ Very sl. sol. in hot $\text{H}_2\text{O}.$ (Ber-
 zelius.)

$+ 6\text{H}_2\text{O}.$ As the $2\text{H}_2\text{O}$ salt. (Boutzour-
 eano.)

$+ 7\text{H}_2\text{O}.$ Sl. sol. in $\text{H}_2\text{O}.$ Easily sol. in
 acetic, and mineral acids. (Hilger, Z. anal.
 13. 132.)

Magnesium hydrogen selenite, $\text{MgH}_2(\text{SeO}_3)_2,$
 $+ 3\text{H}_2\text{O}.$

Very deliquescent. Easily sol. in $\text{H}_2\text{O}.$
 (Nilson.)

Insol. in alcohol. (Muspratt.)

$\text{MgO}, 2\text{SeO}_2.$ Insol. in $\text{H}_2\text{O};$ sol. in acids.
 (Boutzoureano.)

Magnesium tetrahydrogen selenite,

$\text{MgH}_4(\text{SeO}_3)_3,$ and $+ 3\text{H}_2\text{O}.$

Sol. in $\text{H}_2\text{O}.$ (Nilson.)

Manganous selenite, $MnSeO_3 + H_2O$.
Precipitate. (Nilson.)
 $+ 2H_2O$. Insol. in H_2O . (Berzelius.)
Sol. in cold $HCl + Aq$. (Muspratt.)
 $+ \frac{1}{2}H_2O$. Insol. in H_2O ; sol. in dil. acids. (Boutzoureano.)

Manganous selenite, acid, $MnSe_2O_5$.
Sol. in H_2O . (Berzelius; Nilson.)
 $MnO, 2SeO_2 + H_2O = MnH_2(SeO_3)_2$.
(Boutzoureano, A. ch. (6) 17. 289.)
 $+ 5H_2O$. Decomp. by H_2O to $MnSeO_3$. (Boutzoureano.)

Manganic selenite, basic, $Mn_2O_3, 2SeO_2$.
Insol. in H_2O , cold H_2SO_4 , or $HNO_3 + Aq$; insol. in hot dil. H_2SO_4 or $HNO_3 + Aq$. (Laugier, C. R. 104. 1508.)
Sol. in warm $HCl + Aq$ with decomp.

Manganic selenite, $Mn_2(SeO_3)_3 + 5H_2O$.
(Laugier.)

Manganic selenite, acid, $Mn_2O_3, 4SeO_2$.
Insol. in H_2O , cold H_2SO_4 , and $HNO_3 + Aq$. Insol. in dil. hot H_2SO_4 , and $HNO_3 + Aq$. Sol. in cold $HCl + Aq$; and in $H_2SO_3 + Aq$ with separation of Se. (Laugier, C. R. 104. 1508.)

Mercurous selenite, basic, $3Hg_2O, 2SeO_2 + 5H_2O$.
(Boutzoureano.)

Mercurous selenite, Hg_2SeO_3 .
Insol. in H_2O or $H_2SeO_3 + Aq$. Sol. in hot $HNO_3 + Aq$. (Köhler, Pogg. 89. 146.)
Sl. sol. in $HCl + Aq$, and $KOH + Aq$. (Berzelius.)

Mercurous selenite, acid, $3Hg_2O, 4SeO_2$.
Insol. in H_2O or $H_2SeO_3 + Aq$. Sl. sol. in boiling $HNO_3 + Aq$. (Köhler.)

Mercuric selenite, basic, $7HgO, 4SeO_2$.
Insol. in H_2O . Sl. sol. in $HNO_3 + Aq$. Easily sol. in $HCl + Aq$. (Köhler, Pogg. 89. 146.)

Mercuric selenite, $HgSeO_3$.
Insol. in H_2O . (Berzelius.) Nearly insol. in $HNO_3 + Aq$. Sol. in $K_2SeO_3 + Aq$. (Divers, Chem. Soc. 48. 585.)
Insol. in dil. $HNO_3 + Aq$; sol. in $HCl + Aq$. (Rosenheim and Pritze, Z. anorg. 1909, 63. 278.)
Solubility in $Na_2SeO_3 + Aq$ at 25° .

$Na_2SeO_3 + Aq$ Normality	% $HgSeO_3$
2.0	2.73
1.0	1.39
0.5	0.70
0.25	0.53
0.125	0.32
0.0625	0.18

(Rosenheim and Pritze, Z. anorg. 1909, 63. 281.)

$HgSeO_3, H_2SeO_3$. Easily sol. in H_2O ; sl. sol. in alcohol. (Berzelius.)
See also selenium dioxide.

Mercuric sodium selenite, $HgSeO_3, Na_2$
Decomp. by H_2O and alkalis with of $HgSeO_3$. (Rosenheim and Pritze anorg. 1909, 63. 279.)

Mercuric selenite sodium chloride, $HgSeO_3, NaCl + 2H_2O$.
Decomp. by H_2O . (Rosenheim and P Z. anorg. 1909, 63. 280.)

Nickel selenite, $NiSeO_3 + H_2O$.
Insol. in H_2O ; sol. in $H_2SeO_3 + Aq$. (pratt, Chem. Soc. 2. 52.)
 $+ \frac{1}{2}H_2O$. Insol. in H_2O . (Boutzou A. ch. (6) 17. 28.)

Nickel selenite, acid.
Sol. in H_2O . (Berzelius.)

Potassium selenite, $K_2SeO_3 + H_2O$.
Very deliquescent. Sol. in nearly all portions in H_2O . Insol. in alcohol, & separates it as oil from aqueous soln (Muspratt, Chem. Soc. 2. 52.)

Potassium hydrogen selenite, $KHSeO_3$.
Very deliquescent. Very sl. sol. in alk (Muspratt, Chem. Soc. 2. 52.)

Potassium trihydrogen selenite, $KH_3(Se$
Very deliquescent. Pptd. from H_2C alcohol. (Muspratt.)
Not deliquescent. (Nilson.)

Potassium hydrogen pyroselenite, $KHS + H_2O$.
(Muthmann, B. 1893, 26. 1015.)

Potassium uranyl selenite, $K_2SeO_3, (UO_2)SeO_3$.
Absolutely insol. in H_2O . (Sendtner.)

Praseodymium hydrogen selenite, $Pr_2(SeO_3)_3, H_2SeO_3 + 3H_2O$.
Sol. in H_2O . (von Scheele, Z. an 1898, 18. 362.)

Samarium selenite, basic, $3Sm_2O_3, 8SeO_2 + 7H_2O$.
Precipitate. (Cleve.)

Samarium selenite, acid, $Sm_2O_3, 4SeO_2 + 5H_2O$.
Precipitate. (Cleve.)

Scandium selenite, $Sc_2(SeO_3)_3 + H_2O$.
Insol. precipitate.

m hydrogen selenite, $\text{Sc}_2(\text{SeO}_3)_2 \cdot \text{SeO}_3$.

in H_2O . Not attacked by cold dil. but easily if warmed.

selenite, Ag_2SeO_3 .

sl. sol. in cold, somewhat more sol. in O . Easily sol. in hot $\text{HNO}_3 + \text{Aq}$, which it is precipitated by H_2O . (us.)

in $\text{K}_2\text{SeO}_3 + \text{Aq}$; sl. sol. in dil $\text{HNO}_3 + \text{H}_2\text{O}$. (Waters, Chem. Soc. 49. 585.)

selenite ammonia, $\text{Ag}_2\text{SeO}_3 \cdot \text{NH}_3$.

in boiling H_2O . (Boutzoureano, A. 17. 289.)

selenite, Na_2SeO_3 .

sol. in H_2O . Insol. in alcohol. (Berzelius.)

selenite, acid, NaHSeO_3 .

anhydrous. Sol. in H_2O .

Na_2O . Sol. in H_2O . (Sacc, A. ch. (3)

) $(\text{SeO}_3)_2$. Not deliquescent. Sol. in H_2O .

vanadium selenite.

vanadioselenite, sodium.

m selenite, $\text{SrSeO}_3 + 7\text{H}_2\text{O}$.

precipitate. Insol. in H_2O . Sol. in HNO_3 . (Muspratt.)

m hydrogen selenite, $\text{SrH}_2(\text{SeO}_3)_2$.

sol. in hot or cold H_2O . (Nilson.)

by insol. in hot or cold H_2O . (Berzelius.)

selenite, Tl_2SeO_3 .

sol. in H_2O . Insol. in alcohol and HCl . (Kuhlmann, Bull. Soc. (2) 1. 330.)

s hydrogen selenite, TlHSeO_3 .

sol. in H_2O than the above comp. (ann.)

selenite, $\text{Th}_2(\text{SeO}_3)_2$.

in H_2O . Sol. in dil. HNO_3 .

decomp. by HCl and H_2SO_4 . (Mann, anorg. 1909, 62. 177.)

selenite, $\text{Th}(\text{SeO}_3)_2 + \text{H}_2\text{O}$, or $8\text{H}_2\text{O}$.

in H_2O ; easily sol. in $\text{HCl} + \text{Aq}$.

)

selenite, acid, 2ThO_2 , $7\text{SeO}_3 +$

H_2O .

$5\text{SeO}_3 + 8\text{H}_2\text{O}$. (Nilson.)

Tin (stannic) selenite.

Insol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$, from which it is pptd. by H_2O . (Berzelius.)

Uranic selenite, $\text{U}_2\text{O}_3 \cdot \text{SeO}_3$.

Insol. in H_2O . (Boutzoureano.)

$+ 2\text{H}_2\text{O}$. (B.)

Uranic selenite, acid, $2\text{U}_2\text{O}_3 \cdot 3\text{SeO}_3 + 7\text{H}_2\text{O}$.

Insol. in H_2O . (Boutzoureano, A. ch. (6) 17. 289.)

Uranyl selenite, $(\text{UO}_2)_2\text{SeO}_3 + 2\text{H}_2\text{O}$.

Precipitate. (Nilson.)

Uranyl selenite, acid, $3\text{UO}_3 \cdot 5\text{SeO}_3 + 7\text{H}_2\text{O}$, or $9\text{H}_2\text{O}$.

Insol. in H_2O .

$\text{UO}_3 \cdot 2\text{SeO}_3 + \text{H}_2\text{O} = (\text{UO}_2)_2\text{H}_2(\text{SeO}_3)_2$.

Absolutely insol. in H_2O and $\text{H}_2\text{SeO}_3 + \text{Aq}$. (Sendtner, A. 195. 325.)

Vanadium selenite.

See Vanadioselenious acid.

Ytterbium selenite, $\text{Yb}_2(\text{SeO}_3)_3$.

Insol. precipitate.

Ytterbium hydrogen selenite, $\text{Yb}_2\text{H}_2(\text{SeO}_3)_4 + 4\text{H}_2\text{O}$.

Insol. in H_2O .

Yttrium selenite, $\text{Y}_2(\text{SeO}_3)_3 + 12\text{H}_2\text{O}$.

Insol. in H_2O or $\text{H}_2\text{SeO}_3 + \text{Aq}$. (Berzelius.)

Sol. in hot $\text{H}_2\text{SeO}_3 + \text{Aq}$. (Nilson.)

Yttrium hydrogen selenite, $\text{Y}_2\text{H}_2(\text{SeO}_3)_4 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O . Easily sol. in HCl or $\text{HNO}_3 + \text{Aq}$. (Cleve.)

Zinc selenite, ZnSeO_3 .

Insol. in H_2O ; sol. in acids. (Boutzoureano, A. ch. (6) 18. 289.)

$+ 2\text{H}_2\text{O}$. Insol. in H_2O . Sol. in H_2SeO_3 , or $\text{HNO}_3 + \text{Aq}$. (Muspratt, Chem. Soc. 2. 52.)

Zinc hydrogen selenite, $\text{ZnH}_2(\text{SeO}_3)_2$.

Easily sol. in H_2O . (Berzelius.)

$+ 2\text{H}_2\text{O}$. Sol. in cold H_2O . (Boutzoureano.)

$\text{ZnO} \cdot 4\text{SeO}_3 + 3\text{H}_2\text{O}$. Easily sol. in H_2O . (Wöhler, A. 63. 279.)

Zinc selenite ammonia, $\text{ZnSeO}_3 \cdot \text{NH}_3$.

Insol. in cold or hot H_2O . (Boutzoureano, A. ch. (6) 17. 289.)

Zirconium selenite, basic, $4\text{ZrO}_2 \cdot 3\text{SeO}_3 + 18\text{H}_2\text{O}$.

Precipitate. Sl. sol. in $\text{HCl} + \text{Aq}$. (Nilson.)

Zirconium selenite, $Zr(SeO_3)_2$.

Absolutely insol. in H_2O ; difficultly sol. in boiling $HCl + Aq.$ (Nilson.)
 $+ H_2O.$ (Nilson.)

Selenium, Se.

Insol. in H_2O . Schultz (J. pr. (2) 32. 390) has obtained a soluble colloidal modification which can be isolated by dialysis.

Insol. in $HCl + Aq.$ Decomp. by $HNO_3 + Aq.$ Sol. in fuming H_2SO_4 . (Schultz-Sellac, B. 4. 113.)

1000 pts. CS_2 dissolve 1 pt. cryst. Se at boiling-point (46.6°), and 0.16 pt. at 0° (Mitscherlich, J. B. 1855. 314.) Solubility of Se in CS_2 is variable—1 pt. Se is sol. in 1376–2464–3746 pts. CS_2 at 20° (Rammelsberg, B. 7. 669). Cryst. Se, which is sol. in CS_2 , becomes insol. in CS_2 after heating to 110° , but after fusion is again sol. (Otto.)

Four modifications.—(1) Amorphous red; (2) crystalline red; (3) granular gray; (4) laminated. 1 and 2 are sol. in CS_2 , 3 and 4 are insol. in CS_2 . All forms are sol. in $SeCl_2$, from which crystallizes a black modification, insol. in CS_2 . CCl_4 with trace of CS_2 dissolves red Se slightly, black Se not at all. $Se(C_2H_5)_2$ dissolves all modifications in small but apparently equal quantities. (Rathke, A. 152. 181.)

According to Saunders (J. phys. Chem. 1900, 4. 428) selenium exists in three modifications.

1. Liquid, including vitreous, amorphous, and colloidal selenium.

a. Vitreous.

Sol. in liquid NH_3 at 25° . (Franklin, Am. Ch. J. 1898, 20. 820.)

Insol. in liquid NH_3 between -30° and $+10^\circ$. Franklin's results are due to impure selenium and not completely dry NH_3 . (Hugot, A. Ch. 1900, (7) 21. 5.)

Almost insol. in CS_2 . (Schützenberger Chimie générale 1. 438.)

Action of light increases solubility in CS_2 . (Saunders, J. phys. Chem. 1900, 4. 456.)

Solubility in methylene iodide at 12° is 1.3 pts in 100. (Retgers, Z. anorg. 1893, 3. 343.)

Sol. in CSe_2 , ethyl selenide, and in ethyl sulphide.

Very sol. in Se_2Cl_2 . (Rathke, A. 1869, 152. 181.)

b. Amorphous.

Completely sol. in CS_2 at ord. temp. if Se has not been heated. If heated or extracted with warm CS_2 it becomes partly insol. (Peterson, Z. phys. Chem. 1891, 81. 612.)

Passes into red crystalline form in solution in CS_2 , C_6H_6 , isobutyric acid, acetophenone, acetone, $CHCl_3$, thiophene, toluene, benzonitrile, ethyl acetate, and alcohol. (Saunders, J. phys. Chem. 1900, 4. 463.)

Solution in quinoline, aniline, pyridine,

etc., cause conversion into gray form.

c. Colloidal.

Forms colloidal solution with H_2O .

A colloidal solution of Se in H_2O obtained. It is not decomp. by boiling is decomp. by electrolytes with separated selenium. (Gutbier, Z. anorg. 1910, 106.)

2. Red crystalline.

Sol. in CS_2 .

3. Gray, crystalline or metallic.

Sol. in selenium chloride and others as vitreous Se. (Rathke, A. 152. 181.)

Sl. sol. in CS_2 , toluene, nitroquinoline, aniline, and KOH . Pp. conc. $KOH + Aq.$ in long needles at 219° . (Coste, C. R. 1909, 149. 674.)

Sol. in many organic substances at temp. as quinoline, ethyl benzoate and naphthalene. (Saunders, J. ph. 1900, 4. 469.)

Completely insol. in CS_2 . (Saunders, J. phys. Chem. 1900, 4. 474.)

Solubility of the two modifications of crystalline Se in CS_2 .

100 cc. boiling CS_2 dissolve mg.

I		II	
Mg. Se		Mg. Se	
3.2		4.1	
2.8		4.0	
3.6		2.9	
3.3		2.8	
2.2		2.9	
...		4.0	

I. Se heated 1 hr. at 140° . Modification A.

II. Modification B.

III. Se heated 48 hrs. at 140° . Modification B.

(Marc, Z. anorg. 1907, 53. 3)

Se_2Br_2 dissolves 22% Se. (Pogg. 128. 327.)

Red Se is sol. in $(NH_4)_2SO_4 + Aq.$ (mann, A. 116. 122.)

Sol. in alkalis and Mg sulphite. 365 pts. $K_2SO_4 + Aq.$ dissolve 10 pts. Se. 360 pts. $MgSO_4 \cdot 3H_2O + Aq.$ dissolve 10 pts. Se.

Insol. in $BaSO_4 + Aq.$ (Rathke, A. 1869, 152. 181.)

Sol. in $KCN + Aq.$ with formation of $KSeCN$. (Franklin, Am. Ch. J. 1898, 20. 830.)

100 pts. methylene iodide dissolve 1 pt. Se at 12° . (Retgers, Z. anorg. 1893, 3. 343.)

Sol. in quinoline, but reacts with it with evolution of H_2 . (Beckmann, Z. anorg. 1906, 51. 236.)

bromide, Se_2Br_2 .

H_2O , but gradually decomp. mp. by absolute alcohol and $\text{C}_2\text{H}_5\text{I}$, but soon decomposed. CS_2 ; less sol. in CHCl_3 and (Schneider, Pogg. 128. 327.)

bromide, SeBr_4 .

with decomp. Decomp. by $\text{HCl} + \text{Aq}$; sl. sol. in CS_2 , $\text{C}_2\text{H}_5\text{Br}$. (Schneider, Pogg.

 $\text{C}_2\text{H}_5\text{I}$.**tribromotrichloride, SeCl_2Br_3 .**

CS_2 . (Evans and Ramsay, 62.)

tribromide sulphur trioxide, Se_2O_3 .

H_2O . (Prandtl, Z. anorg.

tribromochloride, SeClBr_3 .**tribromochlorotribromide.****tribromochloride, Se_2Cl_2 .**

decomp. by H_2O . Dissolves in H_2O on heating (l. 181). Insol. in conc. H_2SO_4 ; fuming H_2SO_4 . Sol. in CHCl_3 , Gradually decomp. by H_2O , ther. (Divers and Shimose, sol. in CS_2 . (Evans and Ramsay, 45. 62.)

tribromochloride, SeCl_4 .

on moist air. Decomp. with H_2O , A. ch. 9. 225.) Insol. in H_2O in hot POCl_3 , from which on cooling. (Michaelis, Zeit. 460.) Very sl. sol. in CS_2 . (Ramsay, Chem. Soc. 45. 62.)

tribromochloride, SeCl_2Br_2 .

(Ramsay, Chem. Soc. 45. 62.)

tribromochloride, SeClBr_3 .

in CS_2 . (Evans and Ramsay.)

tribromochloride, SeCl_3Br .**tribromochlorotrichloride.****de.**

$\text{HF} + \text{Aq}$. Decomp. immediately (Knox.)

iodide, Se_2I_2 .

H_2O . All solvents of iodine that element. (Schneider,

Selenium tetraiodide, SeI_4 .

Slowly decomp. by much H_2O . Iodine is dissolved out by all solvents of that element. (Schneider, Pogg. 129. 627.)

Selenium nitride.

See Nitrogen selenide.

Selenium monoxide, SeO (?).

Sl. sol. in H_2O . (Berzelius.)

Does not exist. (Sacc.)

Selenium dioxide, SeO_2 .

Deliquescent. Easily sol. in H_2O and alcohol. Sol. in glacial $\text{HC}_2\text{H}_3\text{O}_2$. (Hinsberg, A. 260. 40)

Solubility in H_2O between -3° and $+36^\circ$ $= 45.0 + 0.7692t$. (Étard, C. R. 1888, 106. 742.)

1 pt. is sol. in 2.67 pts. H_2O at 11.3°

1 pt. " " " 2.60 " " " 14°

1 pt. " " " 2.54 " " " 15.6°

(de Coninck, C. R. 1906, 142. 571.)

Sp. gr. of $\text{SeO}_2 + \text{Aq}$ at t° .

t°	% SeO_2	Sp. gr.
15.1	1	0.9923
15.3	2	1.0068
13.0	3	1.0200
13.0	4	1.0302
14.5	5	1.0346
14.8	6	1.0402
14.1	7	1.0535
15.0	8	1.0571
15.6	9	1.0719
15.2	10	1.0743

(de Coninck, C. R. 1906, 142. 571.)

See also Selenious acid.

1 pt. SeO_2 is sol. in 9.84 pts. alcohol (93°) at 14°

1 pt. SeO_2 is sol. in 15.0 pts. methyl alcohol at 11.8° .

1 pt. SeO_2 is sol. in 23.0 pts. acetone at 15.3°

1 pt. SeO_2 is sol. in 90.0 pts. acetic acid at 12.9° (de Coninck, C. R. 1906, 142. 572.)

Traces dissolve in acetic anhydride. Sol. in phenyl mercaptan. (Hinsberg, A. 1890, 260. 40.)

Insol. in pure C_6H_6 . (Clausnizer, A. 1879, 196. 271.)

See Selenious acid.

Selenium trioxide, SeO_3 .

Not obtained in a pure state. (Cameron and Macallan.)

See Selenic acid.

Selenium dioxide hydrobromic acid, SeO_2 , 4HBr .

Decomp. at 55° (Ditte, A. ch. (5) 10. 82.)

SeO_2 , 5HBr . Decomp. at 65° (Ditte, A. ch. (5) 10. 82.)

Selenium dioxide hydrochloric acid, SeO_2 , 2HCl .

Decomp. at 26° .

SeO_2 , 4HCl . Decomp. at 0° . Sol. in H_2O without evolution of gas. (Ditte, A. ch. (5) 10. 82.)

Selenium dioxide sulphur trioxide, SeO_2 , SO_3 .

Decomp. violently by H_2O . (Weber, B. 19. 3185.)

Composition may be $(\text{SeO})\text{SO}_3$ (?).

Selenium oxy-compounds.

See Selenyl compounds.

Selenium diphosphide, P_2Se .

See Phosphorus monoselenide.

Selenium tetraphosphide, P_4Se .

See Phosphorus semiselenide.

Selenium monosulphide, SeS .

Insol. in H_2O and ether. Sol. in CS_2 . Decomp. by alcohol. (Ditte, C. R. 73. 625, 660.)

Other compounds of Se and S are probably mixtures of the two elements.

Selenium disulphide, SeS_2 .

Compound of this formula is a mixture of SeS and S. (Ditte, C. R. 73. 625, 660.)

Selenium sulphoxide, SeSO_2 .

Decomp. by H_2O . Sol. in fuming H_2SO_4 , conc. H_2SO_4 . Sol. in H_2SO_4 of 1.806 sp. gr. without decomp. (Weber, Pogg. 156. 531.)

Decomp. by H_2O ; sol. in H_2SO_4 . (Divers and Shimosé, B. 17. 858.)

Seleniuretted hydrogen, H_2Se .

See Hydrogen selenide.

Selenoarsenic acid.**Potassium selenoarsenate, $\text{KAsSe}_3 + 2\text{H}_2\text{O}$.**

Only sl. sol. in cold H_2O ; sol. in warm H_2O with decomp.; more stable in $\text{KOH} + \text{Aq}$. (Clever, Z. anorg. 1895, 10. 132.)

Sodium selenoarsenate, $\text{Na}_3\text{AsSe}_4 + 9\text{H}_2\text{O}$.

Very sol. in H_2O ; very unstable. (Szary, B. 1895, 28. 2658.)

Selenoarsenious acid.**Sodium selenoarsenite, $\text{Na}_3\text{AsSe}_3 + 9\text{H}_2\text{O}$.**

Moderately sol. in H_2O . (Clever and Muthmann, Z. anorg. 1895, 10. 130.)

Selenobismuthous acid.**Potassium metaselenobismuthite, Bi_2Se_3 , K_2Se or KBiSe_3 .**

Insol. in cold dil. $\text{HCl} + \text{Aq}$. Sol. on warming, with evolution of H_2Se . (Hilger and van Scherpenberg, Mitt. Pharm. II. 4.)

Selenocyanhydric acid, HSeCN .

Known only in aqueous solution.

Ammonium selenocyanide, NH_4SeCN .

Very deliquescent, and sol. in H_2O .

Barium —, $\text{Ba}(\text{SeSCN})_2$.

Very sol. in H_2O .

Lead —, $\text{Pb}(\text{SeCN})_2$.

Sl. sol. in cold, sol. with sl. decomp. in boiling H_2O . Insol. in alcohol.

Mercurous —, $\text{Hg}_2(\text{SeCN})_2$.

Ppt.

Mercuric —, $\text{Hg}(\text{SeCN})_2$.

Sl. sol. in cold H_2O . Easily sol. in MCl , MSCN , or $\text{MSeCN} + \text{Aq}$; also sol. in hot $\text{HgCl}_2 + \text{Aq}$. (Cameron and Davy, C. N. 44. 63.)

Decomp. by hot H_2O . (Rosenheim, Z. anorg. 1909, 63. 276.)

Mercuric potassium —, $\text{Hg}(\text{SeCN})_2$, KSeCN .

Easily sol. in H_2O . Sl. sol. in cold alcohol. (Cameron and Davy, C. N. 44. 63.)

Mercuric selenocyanide chloride, $\text{Hg}(\text{SeCN})_2$, HgCl_2 .

Sol. in boiling H_2O and in abs. alcohol. Decomp. by long boiling with H_2O . (Rosenheim and Pritze, Z. anorg. 1909, 63. 276.)

Platinum potassium — (Potassium platinum-selenocyanide), $\text{K}_2\text{Pt}(\text{SeCN})_4$.

Sol. in H_2O and alcohol. (Clarke, B. 11. 1325.)

Potassium —, KSeCN .

Very deliquescent, and sol. in H_2O with absorption of heat. More sol. in H_2O than KSCN . Sol. in alcohol.

Potassium — mercuric bromide, KSeCN , HgBr_2 .

Sl. sol. in cold, more easily in hot H_2O or alcohol. (Cameron and Davy, C. N. 44. 63.)

m selenocyanide mercuric chloride, CN, HgCl_2 .
bromide.

m — mercuric iodide, KSeCN ,
in cold, easily in hot H_2O or alcohol
n and Davy.)

m — mercuric sulphocyanide,
 CN, Hg(SCN)_2 .
in cold, much more in hot H_2O or
Somewhat sol. in ether. (Cameron
y.)

—, AgSeCN .
in H_2O . Almost insol. in NH_4OH +
old dil. acids. Quickly decomp. by
acids.

—, NaSeCN .
sol. in H_2O .

nolybdic acid.

m selenomolybdate, $5\text{K}_2\text{O, } 6\text{SeO}_3$,
 MoO_3 .

y sol. in H_2O without decomp.
Am. Ch. J. 1895, 17. 177.)

entathionic acid.

selenopentathionate, $\text{Na}_2\text{S}_4\text{SeO}_6$.
solution may be boiled for some time
change. (Norris and Fay, Am. Ch.
23. 121.)

hosphoric acid.

m selenophosphate,
 $\text{H}_4\text{O, P}_2\text{O}_5, 2\text{SeO}_3 + 3\text{H}_2\text{O}$.
 H_2O with decomp. (Weinland, B.
1402.)

m selenophosphate,
 $\text{O, P}_2\text{O}_5, 2\text{SeO}_3 + 3\text{H}_2\text{O}$.
 H_2O with decomp.
 $\text{O, P}_2\text{O}_5, 5\text{SeO}_3 + 5.5\text{H}_2\text{O}$. Easily
 I_2O . (Weinland.)

n selenophosphate,
 $\text{O, P}_2\text{O}_5, 2\text{SeO}_3 + 3\text{H}_2\text{O}$.
 H_2O with decomp. (Weinland.)

ophosphorous acid.

m triselenophosphite,
 $\text{PSe}_3 + 2\frac{1}{2}\text{H}_2\text{O}$.
p. by moist air and dil. acids; sol.
 $\text{KOH} + \text{Aq.}$; sl. sol. in cold, easily sol.
 I_2O . (Muthmann, Z. anorg. 1897,

Selenosamic acid, HSeO_2NH_2 .

Known only in its salts.

Ammonium selenosamate, $(\text{NH}_4)\text{SeO}_2\text{NH}_2$.
Deliquescent. Decomp. slowly by H_2O
into $(\text{NH}_4)_2\text{SeO}_3$.

1 pt. is sol. in 116 pts. cold alcoholic am-
monia at 12° . More sol. in hot alcoholic
ammonia. Sl. attacked by cold HCl or HNO_3 .
(Cameron and Macallan, C. N. 1888, 57. 163.)

Ammonium hydrogen selenosamate,
 $\text{NH}_4\text{H}(\text{SeO}_2\text{NH}_2)_2$.
Deliquescent. Sol. in 14 pts. alcohol at
 14° . (Cameron and Macallan, Proc. Roy.
Soc. 44. 112.)

Selenostannic acid.

Ammonium selenostannate, $3\text{SnSe}_2, (\text{NH}_4)_2\text{Se}$
 $+ 3\text{H}_2\text{O}$.
Sol. in H_2O . (Ditte, C. R. 95. 641.)

Platinum potassium —, $\text{K}_2\text{Se, } 3\text{PtSe, SnSe}_2$.
Insol. in hot or cold H_2O , NH_4OH , or
 $\text{KOH} + \text{Aq.}$ Not attacked by hot $\text{HCl} + \text{Aq.}$
(Schneider, J. pr. (2) 44. 507.)

Platinum sodium —, $\text{Na}_2\text{Se, } 3\text{PtSe, SnSe}_2$.
Properties as the corresponding K salt.
(Schneider.)

Potassium —, $\text{K}_2\text{SnSe}_3 + 3\text{H}_2\text{O}$.
Easily sol. in H_2O . (Ditte, C. R. 95. 441.)

Selenosulphantimonic acid.

Sodium selenosulphantimonate, $\text{Na}_2\text{SbSeS}_2 +$
 $9\text{H}_2\text{O}$.
Sol. in H_2O . (Hofacker, A. 107. 6.)
 $\text{Na}_2\text{SbS}_{1.5}\text{Se}_{2.5} + 9\text{H}_2\text{O}$. Somewhat sol. in
 H_2O . (Pouget, A. ch. 1899, (7) 18. 564.)

Selenosulphantimonous acid.

Potassium selenosulphantimonite,
 $\text{Sb}_4\text{S}_3\text{Se}_3\text{K}_{10} + 4\text{H}_2\text{O}$.
Sol. in H_2O . (Pouget, A. ch. 1899, (7)
18. 563.)

Sodium selenosulphantimonite, $\text{Na}_2\text{SbS}_{1.5}\text{Se}_{1.5}$
 $+ 9\text{H}_2\text{O}$.
Sol. in H_2O . (Pouget, A. ch. 1899, (7) 18.
564.)

Selenosulpharsenic acid.

Potassium selenosulpharsenate, $3\text{K}_2\text{S, As}_2\text{Se}_3$
 $+ 12\text{H}_2\text{O}$.
Very unstable in the air. Very sol. in H_2O .
Fairly stable in aqueous solution. Decomp.
by acids. (Clever, Z. anorg. 1895, 10. 134.)

Sodium selenosulpharsenate, $\text{Na}_3\text{AsS}_2\text{Se} + 8\text{H}_2\text{O}$.

Decomp. by acids; stable in dry air. (Messinger, B. 1897, **30**. 801.)

$3\text{Na}_2\text{S}, \text{As}_2\text{Se}_3 + 18\text{H}_2\text{O}$. Quite sol. in H_2O ; quite stable in air. (Clever, Z. anorg. 1895, **10**. 140.)

$\text{Na}_4\text{As}_2\text{S}_3\text{Se}_2 + 16\text{H}_2\text{O}$. Sl. sol. in H_2O ; decomp. by acids. (Messinger, B. 1897, **30**. 803.)

$\text{Na}_4\text{As}_2\text{S}_7\text{Se} + 16\text{H}_2\text{O}$. Stable in dry air; easily sol. in H_2O ; decomp. by acids. (Messinger, B. 1897, **30**. 800.)

$\text{Na}_3\text{AsS}_2\text{Se}_2 + 9\text{H}_2\text{O}$. Decomp. in aq. solution by dil. acids. (Messinger, B. 1897, **30**. 802.)

$\text{Na}_2\text{AsSSe}_2 + 9\text{H}_2\text{O}$. Sol. in H_2O ; decomp. by aq. acids; hygroscopic. (Messinger.)

Selenosulphophosphorous acid.

Potassium selenosulphophosphite, $2\text{K}_2\text{S}, \text{P}_2\text{Se}_3 + 5\text{H}_2\text{O}$.

Sol. in H_2O ; decomp. by acids. (Muthmann, Z. anorg. 1897, **13**. 198.)

Selenosulphostannic acid.

Ammonium selenosulphostannate, $(\text{NH}_4)_2\text{S}, 3\text{SnSe}_2 + 3\text{H}_2\text{O}$.

Easily decomp. (Ditte, C. R. 1882, **95**. 643.)

Potassium —, $\text{K}_2\text{SnSe}_2\text{S} + 3\text{H}_2\text{O}$.

Very easily sol. in H_2O . (Ditte, C. R. **95**. 641.)

Sodium —, $\text{Na}_2\text{SnSe}_2\text{S} + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Ditte, C. R. **95**. 641.)

Selenosulphoxyarsenic acid.

Sodium selenosulphoxyarsenate, $\text{Na}_2\text{AsO}_3\text{SSe} + 10\text{H}_2\text{O}$.

Easily sol. in H_2O but solution rapidly decomp. (Messinger, B. 1897, **30**. 798.)

$\text{Na}_4\text{As}_2\text{S}_2\text{SeO}_3 + 24\text{H}_2\text{O}$. Sol. in H_2O . (Messinger.)

$\text{Na}_4\text{As}_2\text{SeS}_3\text{O}_4 + 20\text{H}_2\text{O}$. Stable in dry air. Sl. sol. in H_2O ; decomp. by dil. acids. (Messinger.)

$\text{Na}_4\text{As}_2\text{S}_3\text{Se}_2\text{O}_3 + 20\text{H}_2\text{O}$. Ppt. (Messinger.)

$\text{Na}_4\text{As}_2\text{S}_7\text{Se}_2\text{O}_3 + 36\text{H}_2\text{O}$. Decomp. by aq. acids; sol. in H_2O ; quite stable. (Messinger.)

Selenosulphur trioxide, SeSO_3 .

See Selenium sulphoxide.

Selenosulphuric acid, H_2SeSO_3 .

Known only in its salts.

Potassium selenosulphate, $\text{K}_2\text{SeSO}_4 + \text{H}_2\text{O}$.

Deliquescent in moist air; decomp. by acids. (Rathke, J. pr. **95**. 1.)

Selenotrithionic acid, $\text{H}_2\text{S}_3\text{SeO}_4$.

Known only in solution, which is dark. (Schulze, J. pr. (2) **32**. 390.)

Barium selenotrithionate.

Sol. in H_2O . (Rathke.)

Potassium —, $\text{K}_2\text{SeS}_3\text{O}_4$.

Sol. in H_2O with gradual decomp. (J. pr. **95**. 8; **97**. 56.)

Diselenotrithionic acid, $\text{H}_2\text{S}_2\text{Se}_2\text{O}_4$.

Exceedingly unstable. (Schulze.)

Selenovanadic acid.

Lithium selenovanadate, $4\text{Li}_2\text{O}, 6\text{V}_2\text{O}_5 + 30\text{H}_2\text{O}$.

Very sol. in H_2O . (Prandtl and Lustig, Z. anorg. 1907, **53**. 401.)

Potassium selenovanadate, $2\text{K}_2\text{O}, 12\text{SeO}_2 + 12\text{H}_2\text{O}$.

(Prandtl and Lustig.)

$3\text{K}_2\text{O}, 5\text{V}_2\text{O}_5, 16\text{SeO}_2 + 40\text{H}_2\text{O}$. and Lustig.)

$4\text{K}_2\text{O}, 6\text{V}_2\text{O}_5, 21\text{SeO}_2 + 37\text{H}_2\text{O}$. and Lustig.)

$5\text{K}_2\text{O}, 10\text{V}_2\text{O}_5, 26\text{SeO}_2 + 43\text{H}_2\text{O}$. and Lustig.)

Sodium selenovanadate, $4\text{Na}_2\text{O}, 6\text{V}_2\text{O}_5 + 20\text{H}_2\text{O}$.

Very sol. in H_2O . Solution decomposes gradually. (Prandtl and Lustig.)

$2\text{Na}_2\text{O}, 7\text{V}_2\text{O}_5, 10\text{SeO}_2 + 13\text{H}_2\text{O}$. and Lustig.)

$2\text{Na}_2\text{O}, 7\text{V}_2\text{O}_5, 12\text{SeO}_2 + 45\text{H}_2\text{O} + 90\text{H}_2\text{O}$. (Prandtl and Lustig.)

Selenoxyarsenic acid.

Ammonium selenoxyarsenate, $2(\text{NH}_4)_2\text{SeO}_3, \text{As}_2\text{O}_3 + 3\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Weinland, Z. anorg. 1903, **36**. 1403.)

Barium sodium selenoxyarsenate, $\text{BaNaAsO}_3\text{Se} + 9\text{H}_2\text{O}$.

Ppt. (Weinland, Z. anorg. 1897, **13**. 198.)

Potassium selenoxyarsenate, $2\text{K}_2\text{O}, \text{As}_2\text{O}_3 + 3\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Weinland and Bartling, B. 1903, **36**. 1403.)

$7\text{K}_2\text{O}, 10\text{SeO}_2, 2\text{As}_2\text{O}_3 + 11\text{H}_2\text{O}$. Sol. in H_2O . (Weinland and Bartling, B. 1903, **36**. 1403.)

$3\text{K}_2\text{O}, \text{As}_2\text{Se}_3 + 10\text{H}_2\text{O}$. Easily decomp.

Very sol. in H_2O . (Clever, Z. , 10. 126.)

selenoxyarsenate, $2Rb_2O, 2SeO_3, -3H_2O$.

H_2O with decomp. (Weinland and)

oxyarsenate, Na_3AsSeO_3 .

able in air and in aq. solution. B. 1896, 29. 1010.)

$As_2O_3 + 12H_2O$. Stable in the air sol. in H_2O with decomp. (Wein-rg. 1897, 14. 50.)

$As_2O_3 + 12H_2O$. Very sol. in H_2O ; le. (Szarvasy, B. 1895, 28. 2657.)

$3Na_2Se, As_2O_3 + 50H_2O$. Easily . Solution may be boiled for a without decomp. (Clever, Z. 10. 136.)

phosphoric acid.

triselenmonoxyphosphate, $PSe_3O + 10H_2O$.

Ephraim, B. 1910, 43. 280.)

hydrogen triselenmonoxyphos-
(NH_4) $_2H(PSe_3O)_2 + 18H_2O$.
(Ephraim.)

hydrogen diselendioxyphosphate,
 $Se_2O_2 + 14H_2O$.

in moist air. (Ephraim.)

selenoxyphosphate, $K_3PSe_3O_{11}$.

by HNO_3 . Insol. in alcohol and
(Ephraim.)

oselenotrioxophosphate,
 $O_3 + 20H_2O$.

by H_2O . (Ephraim.)

selenmonoxyphosphate, Na_3PSe_3O
)

H_2O . Decomp. in aq. solution.
in conc. $NaOH + Aq$. (Muth-
org. 1897, 13. 199.)

amide, $SeOBr_2$ (?).

; Pogg. 129. 450.)

amide sulphur trioxide, $SeOBr_2$,

Z. anorg. 1909, 62. 242.)

ride, SeO_2Cl_2 .

comp. by H_2O . (Weber, Pogg.

ur chloride.

selenyl chloride.

Selenyl stannic chloride, $2SeOCl, SnCl_4$.

Extremely deliquescent. Completely sol.
in H_2O . (Weber, B. A. B. 1865. 154.)

Selenyl titanium chloride, $2SeOCl_2, TiCl_4$.

Decomp. by H_2O with separation of an
insol. residue. Decomp. by $NH_4OH + Aq$.
(Weber, B. A. B. 1865. 154.)

Sesquiauramine.

See Sesquiauramine.

Sesquihydraurylamine, $(HOAu)_3N, NH_3$.

See Sesquihydraurylamine.

Silicic acid, SiO_2, xH_2O .

See also Silicon dioxide.

Silicic acid is sol. in 1000 pts. pure H_2O .
(Kirwan.)

When pptd. from alkali silicates + Aq by
 CO_2 , 0.021 pt. SiO_2 remains dissolved in 100
pts. H_2O . (Struckmann, A. 94. 341.)

When pptd. as above, 100 pts. H_2O dissolve
0.09 pt. SiO_2 in 3 days; 100 pts. $H_2CO_3 + Aq$
dissolve 0.078 pt. SiO_2 in 3 days. But if
heated much more dissolves, the jelly itself
becoming liquid, such jelly containing 2.49
pts. SiO_2 to 100 pts. H_2O . This solution is
not pptd. by considerable quantities of al-
cohol, but conc. $(NH_4)_2CO_3$, $NaCl$, or $CaCl_2 +$
 Aq , etc., cause gelatinization. (Maschke, J.
pr. 68. 234.)

Solubility in H_2O depends on the amt. of
 H_2O , in presence of which the silicic acid
is set free by dil. acids, CO_2 , or alkali salts +
 Aq . If H_2O is present in sufficient quantity
to retain the silicic acid, much more will
remain in solution than can be dissolved by .
digesting the gelatinous acid with H_2O after-
wards. 1 pt. SiO_2 can thus be held in solution
by 500 pts. H_2O . Presence of NH_4OH ,
 $(NH_4)_2CO_3$, or NH_4Cl (in solutions of which
 SiO_2 is remarkably insol.) diminishes the
power of H_2O to retain SiO_2 in solution. SiO_2
is always more sol. in dil. than conc. NH_4OH
+ Aq . (Liebig, A. 94. 373.)

Silicic acid from the coagulation of the col-
loidal form (see p. 802) is sol. in about 5000
pts. H_2O when formed from a 1% solution,
and 10,000 pts. when formed from a 5% solu-
tion, but is insol. after being dried. (Graham,
A. 121. 36.)

Silicic acid is more sol. in dil. acids than in
 H_2O , because, when acid is added in excess
to moderately dil. $K_2SiO_3 + Aq$, the solution
remains clear, but if only enough acid is
added to neutralize the base present, silicic
acid will gradually separate out. If acid is
added to conc. $K_2SiO_3 + Aq$, silicic acid sep-
arates out insol. in excess of acid, but if
20-30 pts. H_2O are present to 1 pt. K_2SiO_3 ,
and an excess of acid added at once, the silicic
acid will remain in solution. This result is
obtained with HCl , HNO_3 , H_2SO_4 , or

$\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ These solutions may dissolve a neutral salt until saturated and no silicic acid will separate out. Therefore it is the acid that holds the SiO_2 in solution, and not the H_2O . (C. J. B. Karsten, (1826) Pogg. 6. 353.)

Even CO_2 has the power of holding SiO_2 in solution. (Karsten, *l. c.*)

Solubility in acids of silicic acid of Struckmann (see above): 100 pts. dil. $\text{HCl} + \text{Aq}$ of 1.088 sp. gr. dissolve 0.0172 g. SiO_2 in 11 days; 100 pts. H_2O sat. with CO_2 dissolve 0.0136 g. SiO_2 in 7 days.

Silicic acid obtained by passing SiF_4 into H_2O is sol. while still moist in 11,000 pts. cold, and 5500 pts. boiling $\text{HCl} + \text{Aq}$ of 1.115 sp. gr. (Fuchs, A. 82. 119.)

Silicic acid at the moment of separation (as in dissolving cast-iron, steel, etc.) is abundantly sol. in aqua regia (3 pts. $\text{HCl} + \text{Aq}$ of sp. gr. 1.13 and 1 pt. $\text{HNO}_3 + \text{Aq}$ of sp. gr. 1.33). (Wittstein, Z. anal. 7. 433.)

The aq. solution obtained by the hydrolysis of ethyl silicate is more stable in acids + Aq or alkali than in pure H_2O . (Jordis, Z. anorg. 1903, 35. 16.)

$\text{NH}_4\text{OH} + \text{Aq}$ dissolves considerable freshly precipitated silicic acid, $(\text{NH}_4)_2\text{CO}_3$ only a very little. (Karsten, Pogg. 6. 357.)

Dry or ignited SiO_2 is sol. in $\text{NH}_4\text{OH} + \text{Aq}$. 100 pts. $\text{NH}_4\text{OH} + \text{Aq}$ containing 10% NH_3 dissolve: 0.714 pt. SiO_2 from gelatinous silicic acid; 0.303 pt. from artificially dried silicic acid; 0.377 pt. from amorphous SiO_2 ; 0.017 pt. from quartz. (Pribram, Z. anal. 6. 119.)

$\text{NH}_4\text{OH} + \text{Aq}$ dissolves 0.382 pt. SiO_2 from dry silicic acid; 0.357 pt. from ignited SiO_2 ; 0.00827 pt. from quartz. (Souchay, Z. anal. 11. 182.)

Silicic acid precipitated from alkali silicates + Aq with CO_2 is sol. as follows: 100 pts. pure H_2O dissolve 0.021 pt. SiO_2 ; 100 pts. $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ containing 5% $(\text{NH}_4)_2\text{CO}_3$, 0.020 pt.; 100 pts. containing 1% $(\text{NH}_4)_2\text{CO}_3$, 0.062 pt.; 100 pts. $\text{NH}_4\text{OH} + \text{Aq}$ containing 19.2% NH_3 , 0.071 pt.; 100 pts. containing 1.6%, 0.0986 pt. (Struckmann, A. 94. 341.)

100 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (10% NH_3) dissolve of: crystallised SiO_2 , 0.017 pt.; amorphous SiO_2 , ignited, 0.38 pt.; amorphous $3\text{SiO}_2 \cdot 4\text{H}_2\text{O}$, 0.21 pt.; amorphous silicic acid in form of jelly, 0.71 pt. Upon evaporation no ppt. is formed, even when 80 mols. SiO_2 are present to 1 mol. NH_3 . (Wittstein, J. B. 1866. 192.)

Sol. in KOH or $\text{NaOH} + \text{Aq}$, especially if warm. (Dumas.)

Sol. in K_2SiO_3 or $\text{Na}_2\text{SiO}_3 + \text{Aq}$. (Fuchs.)

Easily sol. in boiling $\text{Na}_2\text{CO}_3 + \text{Aq}$, separating as a jelly on cooling. (Pfaff.)

NH_4Cl or other NH_4 salts ppt. SiO_2 from solution in $\text{Na}_2\text{CO}_3 + \text{Aq}$.

100 pts. Ti_2O in H_2O dissolve 4.17 pts. amorphous SiO_2 in 24 hours' boiling. (Flemming, Jena. Zeit. 4. 36.)

Sol. in butyl amine. (Wurtz, A. d. 166.)

Not more sol. in H_2O containing CO_2 than in pure H_2O . (Petzholdt, J. pr. 60.)

Soluble silicic acid.

Colloidal form by dialysis. Solution containing 4.9% SiO_2 may be evaporated; they contain 14% SiO_2 . The SiO_2 is from its solution thus made in man.

(1) By standing. This happens easily the more conc. the solution hastened by heat. A 10–12% solution gels at ordinary temp. in a few h immediately upon heating. A 5–6% may be kept 5–6 days, a 2% solution months, and a 1% solution may or more years without gelatinizing.

(2) When the solution is evaporated in vacuo at 15° a transparent jelly which is insol. in H_2O .

(3) The coagulation of colloidal is accelerated by powdered graphite, indifferent bodies, and it is brought about in a few minutes by a solution of the borates, even when only $\frac{1}{10,000}$ carbonate is present. (Graham, A. 121. 36.)

(4) Coagulation is also brought about by passing CO_2 through the solution.

CO_2 does not cause coagulation.

Coagulation is not caused by HNO_3 , $\text{HC}_2\text{H}_3\text{O}_2$, $\text{H}_2\text{C}_2\text{H}_3\text{O}_2$, or Aq , or by neutral or acid salts + ham.)

NaCl and $\text{Na}_2\text{SO}_4 + \text{Aq}$ coagulation. (Maschke.)

Alcohol, sugar, glycerine, or not coagulate.

Soluble $\text{Al}_2\text{O}_3 \cdot \text{H}_2\text{O}$, $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$, alk. casein precipitate soluble SiO_2 . (Graham, A. 121. 36.)

The jelly from colloidal SiO_2 is slightly alkaline H_2O . 1 pt. NaOH dissolves in an hour at 100° of the jelly corresponding to 20 (Graham.)

Other colloidal forms.

Various solutions of silicic acid obtained as follows:

The jelly formed when SiF_4 through H_2O dissolves in a large excess of H_2O and SiO_2 separates out on evaporation; it is still sol. in H_2O , but is made insol. by evaporation with HCl or H_2SO_4 . (Berzelius, A. ch. 14. 366.)

When SiF_4 is absorbed by H_2BO_3 , and the HF and H_2BO_3 a large excess of $\text{NH}_4\text{OH} + \text{Aq}$ is obtained which is very sol. in H_2O ; the solution is not decomp. by boiling; on evaporation an insol. powder remains. (Berzelius, A. ch. 14. 366.)

When $\text{K}_2\text{SiO}_3 + \text{Aq}$ is precipitated the precipitate washed and dissolved in Aq , the solution treated with H_2SO_4 boiled, a solution of silicic acid

ses with KOH or $\text{NH}_4\text{OH} + \text{Aq.}$ (3) 21. 40.)

$\text{O}_2 + \text{Aq}$ containing at most 3% ted with $\text{HCl} + \text{Aq}$ of 1.10 sp. IO_2 added until the solution is cent and carefully warmed to us mass is obtained which will O_2 by 12-16 hours' boiling if being exposed to the air. The htly opalescent. The solution ated by heat until it contains a vacuum or over H_2SO_4 , solu- ing 10% may be obtained. The it, freezing, alcohol, or H_2SO_4 coagulate the solution. (Kühn,

O_2 gives off H_2S , and forms a O_2 which, after dilution, can be hs. But when boiled or evapo- n a sol. silicate is added, it tinous. It leaves an insol. vaporated to dryness. (Fremy, 314.)

ns of silicic acid have been de- finite compounds of SiO_2 with nts of H_2O , but it is doubtful if ite compounds exist, as the per-) varies with the moisture of the is exposed. (See Ebelmen, A. 9; Doveri, A. ch. (3) 21. 40; 19; Merz, J. pr. 99. 177; van 11. 2232, etc.)

s are insol. in H_2O with the ex- alkali salts, and these are sol. ratio of the base to the acid is n limit.

icate, $2\text{Al}_2\text{O}_3, \text{SiO}_2 + 10\text{H}_2\text{O}$.

ite. Sol. in acids, with forma- H_2O . Becomes transparent in comp.

O_2 . Min. *Dillnite*.

Min. *Andalusite*, *Chiastolite*, *Disthene* or *Cyanite*. Insol. in

Min. *Allophane*. Completely ls; decomp. by conc. acids with $\text{SiO}_2, x\text{H}_2\text{O}$.

$\text{O}_2 + 4\text{H}_2\text{O}$. Min. *Pholerite*. In- $+ \text{Aq}$.

Min. *Glazerite*.

$\text{O}_2 + 2\text{H}_2\text{O}$. Min. *Kaolin*, *Clay*. HCl or $\text{HNO}_3 + \text{Aq}$; moderately q, when heated to evaporation, and some SiO_2 , and leaves the O_2 , sol. in boiling $\text{Na}_2\text{CO}_3 + \text{Aq}$. is dissolved by heating with 5-6 pt. H_2O until H_2SO_4 evaporates, ting with H_2O .

acked by $\text{H}_2\text{SiF}_6 + \text{Aq}$.

y boiling $\text{KOH} + \text{Aq}$, with resi- (Rammelsberg.)

extracts $\frac{1}{4}$ of the SiO_2 (Mala-

guti); is converted thereby into double sili- cates of K and Al, which are sol. in $\text{HCl} + \text{Aq}$. (Lemberg.)

Solubility in KOH and HCl increased if first heated to a low glow. (Glinka, C. C. 1899, II. 1063.)

Colloidal clay. (Schlösing, C. R. 79. 473.) $+ 4\text{H}_2\text{O}$. *Halloysite*. Decomp. by acids.

$4\text{Al}_2\text{O}_3, 9\text{SiO}_2 + 12\text{H}_2\text{O}$. Min. *Porcelain clay* from Passau.

$\text{Al}_2\text{O}_3, 3\text{SiO}_2 + 3\text{H}_2\text{O}$. Min. *Razoumoffskine*.

$\text{Al}_2\text{O}_3, 4\text{SiO}_2 + 7\text{H}_2\text{O}$. Min. *Montmorillonite*. Not decomp. by $\text{HCl} + \text{Aq}$, but by hot H_2SO_4 .

$+ \text{H}_2\text{O}$. Min. *Pyrophyllite*. Not decomp. by H_2SO_4 .

$+ 3\text{H}_2\text{O}$. Min. *Anauxite*.

$2\text{Al}_2\text{O}_3, 9\text{SiO}_2 + 6\text{H}_2\text{O}$. Min. *Cimolite*.

"Aluminum silicate" is insol. in acetone. (Naumann, B. 1904, 37. 4328); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Aluminum barium silicate, $\text{Al}_2\text{O}_3, \text{BaO}, 2\text{SiO}_2 + \text{H}_2\text{O}$ (?).

Min. *Edingtonite*. Decomp. by $\text{HCl} + \text{Aq}$ with separation of $\text{SiO}_2, x\text{H}_2\text{O}$.

$5\text{Al}_2\text{O}_3, 4\text{BaO}, 10\text{SiO}_2$. (Fremy and Feil, C. R. 85. 1033.)

$2\text{Al}_2\text{O}_3, 4\text{BaO}, 7\text{SiO}_2$. Min. *Barylite*. Very sl. decomp. by alkali carbonates $+ \text{Aq}$. (Blomstrand.)

Aluminum barium potassium silicate,

$\text{Al}_2\text{O}_3, (\text{Ba}, \text{K}_2)\text{O}, 5\text{SiO}_2 + 2\text{H}_2\text{O}$.

Min. *Harmotome*. When finely powdered, difficultly decomp. by $\text{HCl} + \text{Aq}$ with separation of pulverulent $\text{SiO}_2, x\text{H}_2\text{O}$.

$\text{Al}_2\text{O}_3, (\text{Ba}, \text{K}_2)\text{O}, 4\text{SiO}_2$. Min. *Hagalophane*. Scarcely attacked by acids.

Aluminum caesium silicate, $\text{H}_2\text{Cs}_2\text{Al}_2\text{Si}_2\text{O}_{11}$ (?)

Min. *Pollucite*. Very sl. decomp. by $\text{HCl} + \text{Aq}$.

Aluminum calcium silicate, $\text{Al}_2\text{O}_3, \text{CaO}, 2\text{SiO}_2$.

Min. *Anorthite*. Completely decomp. by $\text{HCl} + \text{Aq}$ with separation of pulverulent $\text{SiO}_2, x\text{H}_2\text{O}$.

Min. *Barsowite*. Instantaneously decomp. by $\text{HCl} + \text{Aq}$, with separation of gelatinous $\text{SiO}_2, x\text{H}_2\text{O}$.

$+ 4\text{H}_2\text{O}$. Min. *Gismondite*. Gelatinizes with $\text{HCl} + \text{Aq}$.

$\text{Al}_2\text{O}_3, \text{CaO}, 3\text{SiO}_2 + 3\text{H}_2\text{O}$. Min. *Scolezite*. Easily sol. in $\text{HCl} + \text{Aq}$, without formation of gelatinous SiO_2 . Sol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$ with pptn. of CaC_2O_4 .

Decomp. by, and sol. to a certain extent in $\text{H}_2\text{CO}_3 + \text{Aq}$, and decomp. also even by pure H_2O . (Rogers, Am. J. Sci. (2) 5. 408.)

$+ 5\text{H}_2\text{O}$. Min. *Leryn*. Decomp. by acids without gelatinizing.

$\text{Al}_2\text{O}_3, \text{CaO}, 4\text{SiO}_2 + 3\text{H}_2\text{O}$. Min. *Capor- cianite*. *Leonhardite*. Efflorescent. Easily

sol. in acids, with pptn. of gelatinous SiO_2 , $x\text{H}_2\text{O}$.

Al_2O_3 , CaO , $4\text{SiO}_2 + 4\text{H}_2\text{O}$. Min. *Laumontite*. Easily gelatinizes with HCl or HNO_3 + Aq, but is not affected by H_2SO_4 unless hot.

Al_2O_3 , CaO , $6\text{SiO}_2 + 5\text{H}_2\text{O}$. Min. *Epistilbite*. Gelatinizes with conc. HCl + Aq. (Goldschmidt, Z. anal. 17. 267.)

Scarcely decomp. by boiling conc. HCl + Aq. (Jannasch and Tenne, Miner. Jahrb. 1880, 1. 43.)

+ $6\text{H}_2\text{O}$. *Stilbite*. *Heulandite*. Slowly but completely gelatinized by HCl + Aq.

Al_2O_3 , 2CaO , $3\text{SiO}_2 + \text{H}_2\text{O}$. Min. *Prehnite*. Imperfectly decomp. by acids before ignition, but easily afterwards.

Al_2O_3 , 3CaO , 3SiO_2 . *Lime alumina garnet*. *Grossularite*. Partially decomp. by acids before ignition, but easily afterwards.

$2\text{Al}_2\text{O}_3$, CaO , $2\text{SiO}_2 + \text{H}_2\text{O}$. *Margarite*. Not attacked by acids.

$3\text{Al}_2\text{O}_3$, 4CaO , $6\text{SiO}_2 + \text{H}_2\text{O}$. *Zoisite*. Partially decomp. by HCl + Aq.

$4\text{Al}_2\text{O}_3$, 6CaO , 9SiO_2 . Min. *Meionite*. Completely sol. in HCl + Aq.

Aluminum calcium ferric silicate, $2\text{Al}_2\text{O}_3$, 4CaO , Fe_2O_3 , $6\text{SiO}_2 + \text{H}_2\text{O}$.

Min. *Epidote*. Only sl. attacked by HCl + Aq before ignition.

Aluminum calcium ferric magnesium silicate, $\text{H}_{14}(\text{Ca}, \text{Mg})_{40}(\text{Al}_2, \text{Fe}_2)_{10}\text{Si}_{38}\text{O}_{147}$.

Min. *Vesuvianite*, *Idiocrase*. Only partially decomp. by HCl + Aq before ignition.

Aluminum calcium iron, etc., silicate borate, $\text{H}_2\text{R}_6^{\text{II}}(\text{Al}_2, \text{B}_2)_3\text{Si}_3\text{O}_{32}$.

Min. *Axinite*. Not attacked by HCl + Aq before ignition.

Aluminum calcium magnesium silicate, $4\text{H}_4\text{Ca}_2\text{Mg}_2\text{Si}_4\text{O}_{24}$, $5\text{H}_2\text{CaMgAl}_4\text{O}_{12} = 15\text{Al}_2\text{O}_3$, 13CaO , 37MgO , $24\text{SiO}_2 + 13\text{H}_2\text{O}$.

Min. *Clintonite*. Completely decomp. by HCl + Aq without gelatinization.

$3\text{H}_4\text{Ca}_2\text{Mg}_2\text{Si}_4\text{O}_{24}$, $4\text{H}_2\text{CaMgAl}_4\text{O}_{12}$. Min. *Brandisite*. Not attacked by HCl + Aq. Slowly decomp. by boiling conc. H_2SO_4 .

$5\text{H}_4\text{Ca}_2\text{Mg}_2\text{Si}_4\text{O}_{24}$, $8\text{H}_2\text{CaMgAl}_4\text{O}_{12}$. Min. *Xanthophyllite*. Very sl. decomp. by hot HCl + Aq.

$3(\text{Ca}, \text{Mg})\text{O}$, Al_2O_3 , 2SiO_2 . Min. *Gehlenite*. Easily decomp. by acids.

Aluminum calcium potassium silicate, $(\text{H}, \text{K})_2\text{CaAl}_2\text{Si}_3\text{O}_{18} + 6\text{H}_2\text{O}$.

Min. *Chabasite*. Decomp. by HCl + Aq. $(\text{K}_2, \text{Ca})\text{Al}_2\text{Si}_3\text{O}_{10} + 4\text{H}_2\text{O}$. Min. *Zeagonite*. Completely sol. in HCl + Aq.

Aluminum calcium sodium silicate, $3\text{Al}_2\text{O}_3$, 8CaO , Na_2O , 9SiO_2 .

Min. *Sarcolite*. Decomp. by acids.

$2\text{Al}_2\text{O}_3$, $12(\text{Ca}, \text{Na}_2)\text{O}$, 9SiO_2 (?). *Mellilite*. Gelatinized by acids.

$\text{Na}_2\text{CaAl}_4\text{Si}_7\text{O}_{12}$ (?). Min. *Margarite*. $\text{Na}_2\text{CaAl}_4\text{Si}_{10}\text{O}_{28}$. Min. *Faujasite*. comp. by HCl + Aq.

$(\text{Na}_2, \text{Ca})\text{Al}_2\text{Si}_4\text{O}_{12}$. Min. *Gmelinite*. comp. by HCl + Aq.

$(\text{Ca}, \text{Na}_2)\text{Al}_2\text{Si}_4\text{O}_{18} + 6\text{H}_2\text{O}$. Min. *Difficultly decomp. by HCl + Aq.*

$(\text{Ca}, \text{Na}_2)\text{Al}_2\text{Si}_2\text{O}_8 + 2\frac{1}{2}\text{H}_2\text{O}$. Min. *sonite*. Gelatinizes with HCl + Aq.

$x\text{Na}_2\text{Al}_2\text{Si}_4\text{O}_{16}$, $y\text{CaAl}_2\text{Si}_2\text{O}_8$. Min. *clase*, *Labradorite*. Sl. decomp. by more easily the larger the amt. of Ca

Aluminum calcium sodium silicate, $2(\text{Na}_2, \text{Ca})\text{Al}_2(\text{SiO}_4)_3$, (Na_2, Ca)

Min. *Hauyn*. Gelatinizes with HCl

Aluminum glucinum silicate, Al_2O_3 , 6SiO_2 .

Min. *Beryl*. *Emerald*. Not dec. acids, excepting partially by H_2SO_4 ing ignited.

Al_2O_3 , 2GfO , $2\text{SiO}_2 + \text{H}_2\text{O}$. Min. Not attacked by acids.

Aluminum ferrous silicate, $\text{Al}_2\text{Fe}(\text{SiO}_4)_2$

Min. *Garnet*. Sl. decomp. by HCl

$\text{H}_2\text{FeAl}_2\text{SiO}_7$. Min. *Chloritoid*. tacked by HCl + Aq. Completely by H_2SO_4 .

Al_2O_3 , 3FeO , $3\text{SiO}_2 + 3\text{H}_2\text{O}$. Min.

Aluminum iron lithium potassium, $\text{K}_3\text{Li}_2\text{Fe}_4\text{Al}_{17}\text{Si}_{20}\text{O}_{66}$.

Min. *Zinnwaldite*. Sl. decomp. b

Aluminum ferrous magnesium sil, $6\text{Al}_2\text{O}_3$, $3(\text{Mg}, \text{Fe})\text{O}$, $6\text{SiO}_2 + \text{H}$

Min. *Staurolite*. Not attacked by

Aluminum ferric magnesium silicate, $2(\text{Al}_2, \text{Fe}_2)\text{O}_3$, 2MgO , 5SiO_2 .

Min. *Cordierite*. Sl. attacked by + $x\text{H}_2\text{O}$. Min. *Esmarkite*, *Chlor*

Aluminum ferrous manganous silica, FeO , 2MnO , 3SiO_2 .

Min. *Partschinite*.

Aluminum ferrous sodium, etc., silica, $\text{R}_1^{\text{I}}(\text{Al}_2)(\text{B}_2)\text{Si}_4\text{O}_{20} + \text{R}_2^{\text{II}}(\text{Al}_2)_2(\text{F}$ etc.

Min. *Tourmaline*. Not decomp. l Aq; very sl. decomp. by H_2SO_4 .

Aluminum lithium silicate, Al_2O_3 , Li

Not attacked by acids. (Hau C. R. 90. 541.)

Al_2O_3 , Li_2O , 6SiO_2 .

Al_2O_3 , Li_2O , 4SiO_2 . [Min. *Spectum* attacked by acids.

- $\text{Li}_2\text{O}, 30\text{SiO}_2$. Min. *Petalite*. Not y acids.
- lithium potassium silicate**, $(\text{Li}, \text{K})_2\text{Al}_2\text{Si}_2\text{O}_{10}$.
- vidolite*. Sl. decomp. by acids.
- magnesium silicate**, $5\text{Al}_2\text{O}_3, 4\text{MgO}$, *ophirine*.
- magnesium potassium silicate**, $\text{Al}_2\text{Si}_2\text{O}_{10}, \text{yMg}_{1-2}\text{Si}_2\text{O}_{10}$.
- pidomelane*. Easily decomp. by $\text{O}_2 + \text{Aq}$, with residue of a skeleton $2\text{MgO}, 2\text{K}_2\text{O}, 12\text{SiO}_2 + \text{H}_2\text{O}$. Min. $35\text{MgO}, 7\text{K}_2\text{O}, 36\text{SiO}_2$. Min.
- manganous silicate**, $2\text{Al}_2\text{O}_3, 6\text{MnO}$, imp. by very dil. $\text{HCl} + \text{Aq}$. (Gor-97. 1303.)
- potassium silicate**, $\text{Al}_2\text{O}_3, \text{K}_2\text{O}$, wly decomp. by cold H_2O ; 12% is y hot H_2O . Sol. in alkali hydrox- out insol. in carbonates + Aq . $\text{O}_2, 2\text{SiO}_2$. Insol. in cold H_2O , but es on boiling. Sol. in dil. acids. cali hydroxides or carbonates + Aq . . ch. (6) 10. 45.) $\text{O}_2, 3\text{SiO}_2 + 3\text{H}_2\text{O}$. Easily sol. in l. (Deville, A. ch. (3) 61. 313.) $\text{O}_2, 4\text{SiO}_2$. Min. *Leucite*. De- $\text{HCl} + \text{Aq}$ with separation of pul- O_2 . Ppt. (Deville, C. R. 54. 324) Si_2O_{10} . Min. *Muscovite*, "Mica." ed by HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$. $\text{O}_{17} + 3\text{H}_2\text{O}$. Min. *Pinite*. Partly $\text{HCl} + \text{Aq}$. O_{16} . Min. *Orthoclase*, *Feldspar*. tacked by acids. Slowly sol. in $\text{HCl} + \text{Aq}$ when finely powdered.
- potassium sodium silicate**, $\text{SiO}_2)_4, 5\text{Na}_2\text{Al}_2(\text{SiO}_4)_2$ (?).
- pheline*. Decomp. by $\text{HCl} + \text{Aq}$.
- silver silicate**, $\text{Al}_2\text{Ag}_2\text{Si}_2\text{O}_8$. $\text{NH}_4\text{OH} + \text{Aq}$. (Silber, B. 14. 941.) Si_2O_4 . As above. (Silber.)
- sodium silicate**, $\text{Al}_2\text{O}_3, \text{Na}_2\text{O}, \text{SiO}_2$. cold H_2O , but 38-40% dissolves in (Gorgeu.) $\text{Si}_2\text{O}_3, 2\text{SiO}_2$. Insol. in cold H_2O ;) dissolves 1-2%. Sol. in HCl or ted with 10-20 vols. H_2O . Insol.
- in alkali hydroxides or carbonates + Aq . (Gorgeu, A. ch. (6) 10. 145.) Not attacked by H_2O . (Silber, B. 14. 941.) + $3\text{H}_2\text{O}$. Easily sol. in $\text{HCl} + \text{Aq}$. (v. Ammon.) $\text{Al}_2\text{O}_3, \text{Na}_2\text{O}, 3\text{SiO}_2 + 3\text{H}_2\text{O}$. Decomp. by acids. (Deville, A. ch. (3) 61. 326.) $\text{Al}_2\text{O}_3, \text{Na}_2\text{O}, 4\text{SiO}_2 + 3\text{H}_2\text{O}$. Easily sol. in $\text{HCl} + \text{Aq}$ (v. Ammon.) $2\text{Al}_2\text{O}_3, 3\text{Na}_2\text{O}, 3\text{SiO}_2$. Insol. in cold H_2O , but 27-30% dissolves on boiling. (Gorgeu.) $\text{H}_4\text{Na}_2\text{Al}_4\text{Si}_4\text{O}_{34}$. Min. *Paragonite*. De- comp. by conc. H_2SO_4 . $\text{Na}_2\text{Al}_2\text{Si}_2\text{O}_{12} + 2\text{H}_2\text{O}$. Min. *Analcite*. Readily decomp. by $\text{HCl} + \text{Aq}$. $\text{Na}_2\text{Al}_2\text{Si}_2\text{O}_{10} + 2\text{H}_2\text{O}$. Min. *Natrolite*. Sol. in H_2O with separation of SiO_2 . Also sol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. $\text{Na}_2\text{Al}_2\text{Si}_2\text{O}_{10}$. Min. *Albite*. Not attacked by acids.
- Aluminum sodium silicate chloride**, $3\text{Na}_2\text{Al}_2(\text{SiO}_4)_2, 2\text{NaCl}$
- Min. *Sodalite*. Easily decomp. by HCl , and $\text{HNO}_3 + \text{Aq}$.
- Aluminum sodium silicate sulphate**, $3\text{Na}_2\text{Al}_2(\text{SiO}_4)_2, \text{Na}_2\text{SO}_4$
- Min. *Noceus*. Easily decomp. by $\text{HCl} + \text{Aq}$.
- Aluminum sodium silicate sulphide**.
- See *Ultramarine*.
- Barium silicate**, BaSiO_3 .
- Somewhat sol. in boiling H_2O . Completely sol. in dil. $\text{HCl} + \text{Aq}$. (v. Ammon.) + $6\text{H}_2\text{O}$, or $7\text{H}_2\text{O}$. Boiling H_2O decom- poses, and dissolves about $\frac{1}{2}$ the weight of this substance. (le Chatelier, C. R. 92. 931.) $2\text{BaO}, \text{SiO}_2$. Decomp. by H_2O into $\text{BaSiO}_3 + 6\text{H}_2\text{O}$. (Laudrin.)
- Bismuth silicate**, $2\text{Bi}_2\text{O}_3, 3\text{SiO}_2$.
- Min. *Eulytite*. Decomp. by $\text{HCl} + \text{Aq}$.
- Bismuth ferric silicate**, $\text{Bi}_2\text{Fe}_2\text{Si}_2\text{O}_{17}$.
- Min. *Bismuthoferrite*.
- Boron calcium silicate**.
- See *Borate silicate, calcium, and Silicate borate, calcium*.
- Cadmium silicate**, $\text{CdSiO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$.
- Sol. in $\text{HCl} + \text{Aq}$ with deposition of pul- verulent $\text{SiO}_2, x\text{H}_2\text{O}$. (Rousseau and Tite, C. R. 114. 1262.)
- Cæsium silicate**, Cs_2SiO_3 .
- (Kahlenberg, J. phys. Chem. 1898, 2. 82.)

Calcium silicate, CaSiO_3 .

Slowly sol. in H_2O ; sol. in $\text{HCl} + \text{Aq}$.

Sol. in about 100,000 pts. H_2O . (Gorgeu, A. ch. 1885, (6) 4. 550.)

100 cc. sat. aq. solution of air dried calcium silicate contains 0.0046 g. $\text{CaO} = 0.0095$ g. CaSiO_3 at 17° . (Weisberg, Bull. Soc. 1896, (3) 15. 1097.)

100 cc. sat. solution of air dried calcium silicate in 10% sugar solution at 17° contains 0.0065 g. $\text{CaO} = 0.0135$ g. CaSiO_3 ; 20% sugar solution, 0.0076 g. $\text{CaO} = 0.0175$ g. CaSiO_3 .

After boiling and filtering hot, 10% sugar solution contains 0.0094 g. $\text{CaO} = 0.0195$ g. CaSiO_3 ; 20% sugar solution, 0.0120 g. $\text{CaO} = 0.0249$ g. CaSiO_3 . (Weisberg.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

$4\text{CaO}, 3\text{SiO}_2$. (Laudrin.)

$5\text{CaO}, 3\text{SiO}_2 + 5\text{H}_2\text{O}$. When freshly precipitated is somewhat sol. in H_2O and easily decomp. by $\text{HCl} + \text{Aq}$. (v. Ammon.)

$\text{CaO}, 3\text{SiO}_2 + 2\text{H}_2\text{O}$. (Hjeldt, J. pr. 94. 129.)

$2\text{CaO}, 9\text{SiO}_2 + 3\text{H}_2\text{O}$. Ppt.

CaSiO_3 . Min. *Wollastonite*. Gelatinizes with $\text{HCl} + \text{Aq}$.

$\text{CaSi}_2\text{O}_6 + 2\text{H}_2\text{O}$. Min. *Okenite*. Easily decomp. by cold $\text{HCl} + \text{Aq}$ when powdered.

Calcium glucinum silicate sodium fluoride, $(\text{Ca}, \text{Gl})_{11}\text{Si}_{14}\text{O}_{43}, 6\text{NaF}$.

Min. *Leucophane*.

$7(\text{Ca}, \text{Gl})_2\text{Si}_2\text{O}_7, 6\text{NaF}$. Min. *Melinophane*.

Calcium ferrous silicate, $\text{CaSiO}_3, \text{FeSiO}_3$.

Min. *Hedenbergite*, *Pyroxene*. Sl. decomp. by acids.

Calcium ferric silicate, $\text{Ca}_2\text{Fe}_2(\text{SiO}_4)_2$.

Min. *Garnet*. Sl. decomp. by $\text{HCl} + \text{Aq}$.

$2\text{CaSiO}_3, 11 \text{Fe}_2(\text{SiO}_3)_2$. Min. *Szaboite*. Sl. attacked by $\text{HCl} + \text{Aq}$, and still less by $\text{H}_2\text{SO}_4 + \text{Aq}$.

Calcium ferroferric silicate, $2\text{CaO}, 4\text{FeO}, \text{Fe}_2\text{O}_3, 4\text{SiO}_2 + \text{H}_2\text{O} = \text{H}_2\text{Ca}_2\text{Fe}_4\text{Fe}_2\text{Si}_4\text{O}_{18}$.

Min. *Lievrite*, *Ilvaite*. Easily gelatinizes with $\text{HCl} + \text{Aq}$.

Calcium ferrous magnesium silicate, $(\text{Ca}, \text{Fe}, \text{Mg})\text{SiO}_3$.

Min. *Amphibole*, *Hornblende*, *Asbestos*, *Actinolite*, *Tremolite*. Only sl. attacked by acids.

Calcium ferroferric sodium silicate, $\text{CaSiO}_3, \text{FeSiO}_3, \text{Fe}_2(\text{SiO}_3)_2, \text{Na}_2\text{SiO}_3$.

Min. *Aegirite*.

Calcium magnesium silicate, $\text{CaO}, \text{MgO}, 4\text{SiO}_2$.

(Mutschler, A. 176. 86.)

$\text{Ca}_2\text{SiO}_4, \text{Mg}_2\text{SiO}_4$. Min. *Monticellite*. Completely sol. in dil. $\text{HCl} + \text{Aq}$.

$(\text{Ca}, \text{Mg})\text{SiO}_3$. Min. *Diopside*, *Pyroxene*. Very sl. attacked by acids.

Calcium manganous silicate, $\text{CaSiO}_3, 2\text{MnSiO}_3$.

Min. *Bustamite*.

Calcium potassium silicate.

See under Glass.

Calcium sodium silicate, $(\text{Ca}, \text{Na}, \text{H})$

Min. *Pectolite*. Decomp. by HCl . See under glass.

Calcium sodium silicate zirconate, $\text{Na}_4\text{Ca}(\text{Si}, \text{Zr})_2\text{O}_{21} + 9\text{H}_2\text{O}$.

Min. *Wöhlerite*. Decomp. by $\text{HCl} + \text{Aq}$.

Calcium uranyl silicate, $3\text{CaO}, 5\text{UO}_2, 18\text{H}_2\text{O}$.

Min. *Uranophane*. Gelatinizes with $\text{CaO}, 3\text{UO}_2, 3\text{SiO}_2 + 9\text{H}_2\text{O}$. Min. *Uranophane*.

Calcium silicate chloride, $2\text{CaO}, \text{SiO}_2$.

Insol. in H_2O or alcohol. Sol. in HCl . (le Chatelier, C. R. 97. 1510.)

Calcium silicate fluoride, $2\text{CaO}, 3\text{SiO}_2$.

(Deville, C. R. 52. 110.)

Calcium silicate potassium fluoride, $4\text{H}_2\text{CaSi}_2\text{O}_6, \text{KF} + 4\text{H}_2\text{O}$.

Min. *Apophyllite*. Decomp. by $\text{HCl} + \text{Aq}$.

Calcium silicate stannate.

See Silicostannate, calcium.

Calcium silicate titanate, CaO, SiO_2 .

(Hautefeuille, A. ch. (4) 4. 154.)

Min. *Titanite*. Incompletely decomp. by $\text{HCl} + \text{Aq}$, wholly by $\text{H}_2\text{SO}_4 + \text{Aq}$.

Cerous silicate, $\text{Ce}_2(\text{SiO}_3)_2$.

More or less attacked by HCl , $\text{H}_2\text{SO}_4 + \text{Aq}$, according to the concentration. (Didier, C. R. 101. 882.)

Cerium didymium lanthanum silicate, $2(\text{Ce}, \text{La}, \text{Di})_2\text{O}_3, 3\text{SiO}_2$.

Min. *Cerite*. Gelatinizes with $\text{HCl} + \text{Aq}$.

Cerium glucinum yttrium silicate, $(\text{Y}, \text{Ce}, \text{Gl})_2\text{SiO}_4$.

Min. *Gadolinite*. Easily gelatinizes with $\text{HCl} + \text{Aq}$.

Cerous silicate chloride, $2\text{Ce}_2\text{O}_3, 3\text{SiO}_2, 4\text{CeCl}_2 = \text{Ce}_4(\text{SiO}_4)_2, 4\text{CeCl}_2$.

Insol. in H_2O , but slowly decomp. (Didier, C. R. 101. 882.)

silicate, Co_2SiO_4 .
 es with $\text{HCl} + \text{Aq.}$ (Bourgeois, C. 77.)

ate, CuH_2SiO_4 .
ioptase. Sol. in HCl , HNO_3 , or Aq with separation of SiO_2 . Not y $\text{KOH} + \text{Aq.}$
 $-2\text{H}_2\text{O}$. Min. *Chrysocolla*. De-
 $\text{HCl} + \text{Aq.}$
 Min. *Asperolite*. Easily de-
 $\text{HCl} + \text{Aq.}$
 silicate" is insol. in methyl acetate.
 , B. 1909, 42. 3790.)

ate ammonia, $\text{CuSi}_2\text{O}_5, 2\text{NH}_3$.
 chiff, A. 123. 38.)

silicate, Gl_2SiO_4 .
henacite. Not attacked by acids.

ferrous manganous silicate ferrous
 mous sulphide, $3(\text{Gl}, \text{Fe}, \text{Mn})_2\text{SiO}_4$,
 $(\text{Fe})\text{S}$.

elvine. Decomp. by $\text{HCl} + \text{Aq.}$

us) silicate, Fe_2SiO_4 .
ayalite. Gelatinizes with $\text{HCl} +$

Min. *Grunerite*.
 Min. *Chlorophite*.
 iO_2 . (Zobel, Dingl. 154. 111.)

;) silicate, $\text{Fe}_2\text{Si}_2\text{O}_7 + 5\text{H}_2\text{O}$.
ontronite. Gelatinizes with hot

$9\text{SiO}_2 + 18\text{H}_2\text{O}$. Min. *Hisingerite*.
 $9\text{SiO}_2 + 2\text{H}_2\text{O}$. Min. *Anthosider-*

ferric) magnesium silicate,
 $(\text{Mg})_2\text{Fe}_2\text{Si}_2\text{O}_{10} + 4\text{H}_2\text{O}$.
Cronstadtite. Gelatinizes with

ferric) sodium silicate, $5\text{Na}_2\text{SiO}_3$,
 $\text{O}_3, 4\text{Fe}_2(\text{SiO}_3)_3$.

lknite. Sl. decomp. by acids.

ous) magnesium silicate, Fe_2SiO_4 ,
 iO_4 .

livene, *Chrysolite*, *Peridote*. Gelat-
 HCl or $\text{H}_2\text{SO}_4 + \text{Aq.}$

$\text{SiO}_2 + \frac{1}{2}\text{H}_2\text{O}$. Min. *Picrophyllite*.
). Min. *Monradite*.

SiO_2 . Min. *Bronzite*, *Hypersthene*.
 ed by acids.

$\text{Si}_2, \text{yFeSiO}_3$. Min. *Anthophyllite*.
 ed by acids.

ous) manganous silicate, Fe_2SiO_4 ,
 iO_4 .

nebelite. Gelatinizes with $\text{HCl} +$

Iron (ferrous) manganous silicate chloride,
 $7(\text{Fe}, \text{Mn})\text{SiO}_3, (\text{Fe}, \text{Mn})\text{Cl}_2 + 5\text{H}_2\text{O}$.

Min. *Pyrosmalite*. Completely decomp.
 by conc. $\text{HNO}_3 + \text{Aq.}$

Iron (ferric) potassium silicate, $\text{Fe}(\text{SiO}_3)_2$,
 K_2SiO_3 .

(Hautefeuille and Perrey, C. R. 107. 1150.)

Iron (ferric) sodium silicate, $\text{Na}_2\text{Fe}_2\text{Si}_4\text{O}_{11}$.

Min. *Crokydolite*. Not attacked by
 acids.

Lead silicate.

Insol. in acetone. (Naumann, B. 1904, 37.
 4329); methyl acetate. (Naumann, B. 1909,
 42. 3790.)

See under Glass.

Lithium silicate, $\text{Li}_2\text{Si}_2\text{O}_7$.

Li_4SiO_4 .

Li_2SiO_3 . More stable towards H_2O than
 the other alkali metasilicates. (Rieke and
 Endell, C. C. 1911, I. 7.)

Decomp. by boiling H_2O and acids. (Friedel,
 C. C. 1901, II. 89.)

Scarcely attacked by cold H_2O . (Friedel,
 Bull. Soc. Min. 1901, 24. 141.)

Insol. in ethyl acetate. (Naumann, B.
 1904, 37. 3601); methyl acetate. (Naumann,
 B. 1909, 42. 3790.)

Magnesium silicate, $\text{Mg}_2\text{Si}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Min. *Serpentine*. Decomp. by $\text{HCl} + \text{Aq}$,
 more easily by H_2SO_4 .

Min. *Chrysotile*.

$\text{Mg}_4\text{Si}_2\text{O}_{10} + 6\text{H}_2\text{O}$. Min. *Gymnite*, *Soap-*
stone. Decomp. by H_2SO_4 .

MgSiO_3 . Not completely decomp. by
 $\text{HCl} + \text{Aq.}$

$+ \frac{1}{4}\text{H}_2\text{O}$. Min. *Aphrodite*. Decomp. by
 hot acids.

$+ \frac{1}{2}\text{H}_2\text{O}$. Min. *Picrosmine*.

$+ 1\frac{1}{2}\text{H}_2\text{O}$. Sol. in dil. acids. (v. Ammon.)

Min. *Forsterite*.

$3\text{MgO}, 4\text{SiO}_2 + \text{H}_2\text{O}$ or $4\text{MgO}, 5\text{SiO}_2 +$
 $\frac{3}{4}\text{H}_2\text{O}$. Min. *Talc* or *Steatite*. Not at-
 tacked by HCl or $\text{H}_2\text{SO}_4 + \text{Aq.}$

$\text{Mg}_3\text{Si}_2\text{O}_{11} + 4\text{H}_2\text{O}$. Min. *Spadaite*. De-
 comp. by conc. $\text{HCl} + \text{Aq.}$

$\text{Mg}_2\text{Si}_2\text{O}_7 + 4\text{H}_2\text{O}$. Min. *Meerschaum*.
 Decomp. by $\text{HCl} + \text{Aq.}$

"Magnesium silicate," is insol. in methyl
 acetate. (Naumann, B. 1909, 42. 3790.)

Magnesium potassium silicate,

$\text{MgO}, \text{K}_2\text{O}, 3\text{SiO}_2$.

Easily sol. in acids with decomp. (Du-
 boin, C. R. 1895, 120. 681.)

Magnesium silicate fluosilicate, $\text{Mg}_2\text{Si}_2\text{O}_7$,
 $\text{Mg}_2\text{Si}_2\text{F}_{10}$.

Min. *Humite*, *Chondrodite*. Gelatinizes
 with HCl or $\text{H}_2\text{SO}_4 + \text{Aq.}$

Manganous silicate, Mn_2SiO_4 .

Min. *Tephroite*. Decomp. by $HCl + Aq$ with formation of a stiff jelly.

$MnSiO_3$. Min. *Rhodonite*, *Hermannite*. Not attacked by $HCl + Aq$.

$Mn_4Si_3O_{10} + 2H_2O$. Min. *Friedelite*. Easily gelatinized by $HCl + Aq$.

"Manganous silicate" is insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Manganous zinc silicate, $(Mn,Zn)_2SiO_4$.

Min. *Troostite*.

Manganous silicate chloride, $MnSiO_3$, MnO , $MnCl_2$.

Decomp. by H_2O . (Gorgeu.)

Nickel silicate, Ni_2SiO_4 .

Easily decomp. by acids. (Bourgeois, C. R. 108. 1077.)

Potassium silicate, K_2SiO_3 .

Completely sol. in H_2O . (Ordway, Sill. Am. J. (2) 33. 34.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

$K_2Si_2O_5$. Sol. in H_2O . Conc. $K_2Si_2O_5 + Aq$ contains 23% of the salt, and has sp. gr. 1.25. (Fuchs.)

Hydroscopic. Decomp. at once by H_2O . (Morey, J. Am. Chem. Soc. 1914, 36. 222.)

$K_2Si_8O_{17}$. Partially sol. in H_2O as K_2SiO_3 . $K_2Si_{14}O_{49} + 16H_2O$. Insol. in H_2O . (Forchhammer.)

The K silicates are pptd. from their aqueous solution by alcohol with partial decomp., but less readily than Na silicates.

More sol. in H_2O than the corresponding Na salts. (Ordway, Sill. Am. J. (2) 32. 155.)

Solution can be obtained which is perfectly clear when $4\frac{1}{2}SiO_2$ are present to $1K_2O$, if there are no impurities present. (Ordway.)

The K silicates resemble the Na salts, which see for further data.

Potassium hydrogen silicate, $KHSi_2O_5$.

Not readily affected by H_2O , even by treatment at 100° for several hours.

Decomp. by heating with dil. HCl . (Morey, J. Am. Chem. Soc. 1914, 36. 222.)

Potassium zinc silicate.

Sol. in $KOH + Aq$. (Schindler.)

$K_2O, 6ZnO, 4SiO_2$. Sol. in $HCl + Aq$. (Duboin, C. R. 1905, 141. 255.)

$8K_2O, 9ZnO, 17SiO_2$. Sol. in $HCl + Aq$. (Duboin.)

Potassium zirconium silicate, $K_2O, ZrO_2, 2SiO_2$.

Decomp. by $HCl + Aq$. (Melliss.)

Rubidium silicate, Rb_2SiO_3 .

(Kahlenberg, J. phys. Chem. 1898, 2. 82.)

Silver silicate, Ag_2SiO_3 .

Decomp. by all acids; sol. in $NH_4OH + Aq$. (Hawkins, Sill. Am. J. 139. 311.)

Sodium silicate, Na_2SiO_3 .

Rapidly decomp. by H_2O . (Morey, J. Am. Chem. Soc. 1914, 36. 224.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

+5, 6, and $8H_2O$. Easily sol. in H_2O . + $9H_2O$. Solubility in $\frac{1}{2}N NaOH + Aq$.

100 ccm. of the solution contain 25.5 g. $Na_2SiO_3 + 9H_2O$ at 17.5° .

Sp. gr. of the solution = 1.129. (Vesterberg, C. C. 1913. 777.)

100 ccm. of a sat. solution of sodium silicate in $\frac{1}{2}N NaCl + Aq$. contain 3.5 g. $Na_2SiO_3 + 9H_2O$ at 17.5° . Sp. gr. of solution = 1.15.

100 ccm. of a sat. solution of sodium silicate in sat. $NaCl + Aq$ contain 20.64 g. $Na_2SiO_3 + 9H_2O$ at 17.5° . (Vesterberg.)

$Na_2Si_2O_5$. Sol. in H_2O .

Slowly decomp. by cold H_2O . (Morey, J. Am. Chem. Soc. 1914, 36. 223.)

$Na_4Si_6O_{12}$.

$Na_2Si_2O_7$.

$Na_2Si_4O_{11}$. Slowly sol. in H_2O . + $12H_2O$.

Above compounds are all more or less indefinite.

Water glass. $xNa_2O, ySiO_2 + zH_2O$. Sol. in H_2O , but solution is decomposed by all weak acids, even CO_2 .

Fused water glass is but little acted on by cold H_2O , but when pure, easily dissolves in H_2O by long boiling. (Ordway, Am. J. Sci. (2) 32. 337.)

When the SiO_2 is present in greater proportion than in $Na_2O, 3SiO_2$, it is very difficult to dissolve in H_2O .

Na silicate is less easily sol. in H_2O than the corresponding K compound.

Solubility of water glass in H_2O is much impaired by earthy impurities, so that traces have great effect in preventing the solubility.

NH_4 salts decomp. water glass solutions. A solution containing $\frac{1}{2}\%$ Na_2SiO_3 is scarcely precipitated by NH_4Cl , but easily by NH_4NO_3 . (Fluckinger.)

Precipitated by $NH_4OH + Aq$ as Na_2SiO_3 .

Many sodium and potassium salts, especially the chlorides and acetates, form precipitates in solutions of water glass; these precipitates are larger the more concentrated the solution is, and the greater amount of SiO_2 it contains. Heating hastens the precipitation by chlorides, nitrates, and sulphates, but delays that by acetates. $KOH + Aq$ does not precipitate.

Cold sat. $Na_2SO_4 + Aq$ does not precipitate even on heating, but 1 pt. anhydrous Na_2SO_4 dissolved in 2 pts. H_2O precipitates a bit solution of Na_2SiO_3 .

lved in 1 pt. H_2O precipitate of 1.392 sp. gr.; $NaNO_3$ in 2 mixed with a solution of ove, if the two are present in uses no ppt. in the cold, but warmed to 54° , and redissolves rapidly, but if 2 vols. $NaNO_3$ ent to 1 vol. $Na_2SiO_3 + Aq$, does not disappear on cooling. $LiOH + Aq$ (0.921 sp. gr.) is . $Na_2SiO_3 + Aq$, no ppt. forms, ng the amt. of $NH_4OH + Aq$ greater pt. of the Na_2SiO_3 is solves on heating to 90° , separ- cooling. When 1 pt. NH_4OH to 6–8 pts. $Na_2SiO_3 + Aq$ 30° , a clear liquid is formed into two layers at ordinary

l. K, Na, Li, and NH_4 salts from conc. $Na_2SiO_3 + Aq$. salts lose this power by dilu- H_4 salts and KSON keep this solution is very dil. This is use with NH_4Cl and NH_4NO_3 . orine, propyl amine, crecote, red in glycerine, chloral hy- men solution, and glue solu- rom $Na_2SiO_3 + Aq$; but sugar, ine, urea, sl. alkaline solution, coniine, nicotine, saponine, jalappine, and colophonium $COH + Aq$ do not ppt. SiO_2 . ch. Pharm. (2) 144. 97.)

water glass as such from its n, even when this is very dil., ne decomposition, the alcohol d in solution a portion of a lkaine than that previously O , while the ppt. formed con- than the original silicate.

d K or Na salts ppt. water en added to aqueous solutions. ese solutions exert a decom- the ppt. being always more he original silicate. Na sili- rger deposit than K silicate; of one base is pptd. by a salt oth bases enter into the com- ppt, and the relative propor- K is very nearly the same as in he liquids mixed.

ts have very unequal pptg. ates and chlorides being parti-. Heat increases the pptg. chlorides, sulphates, and ni- nishes that of the acetates. etates are somewhat more he chlorides, but $NaC_2H_3O_2$ ght ppt. with Na_2O , $2\frac{1}{4}SiO_2$, time.

but little effect on the more s.

still less power than $NaNO_3$. no pptg. power, and Na_3AsO_4 e very little effect.

$MHSO_4$, $MHCO_3$, M_2HPO_4 , M_2HASO_4 , ppt. SiO_2 . NH_4 salts also have that effect.

Pptd. water glass, as mentioned above, is much more sol. in H_2O than ordinary water glass, and dissolves in H_2O without decomp. For numerous further details, see articles by Ordway in Sill. Am. J. Sci. vols. 32 and 33; also Storer's Dict.

Sp. gr. of water glass solution containing 14–15% SiO_2 , 13–14% Na_2O , and 70–72% H_2O is 1.30–1.35. (Hager, Comm. 1883.)

Sp. gr. of sat. $Na_2SiO_3 + Aq$ freshly prepared at 18° is 1.2600, and 1 litre contains 4.5 gramme-equivalents $\frac{1}{2}Na_2SiO_3$.

Sp. gr. of sat. solution of Na_2O , $3.4SiO_2$ is 1.366, and 1 litre contains 3.7 gramme-equivalents $\frac{1}{2}(Na_2O, 3.4SiO_2)$. (Kohl- rausch, Z. phys. Ch. 12. 773.)

Sodium zirconium silicate, Na_2O, ZrO_2, SiO_2 .

Decomp. by hot H_2O or $HCl + Aq$. (Gibbs, Pogg. 71. 559.)

$Na_2O, 8ZrO_2, SiO_2 + 11H_2O$. Decomp. by H_2SO_4 . (Melliss.)

Strontium silicate, $SrSiO_3$.

(Stein, Z. anorg. 1907, 55. 164.)
+ H_2O . Sol. in H_2O . (Jordis and Kanter, Z. anorg. 1903, 35. 90.)

Sr_2SiO_4 . (Stein, Z. anorg. 1907, 55. 167.)
 $3SrO, SiO_2$. Sl. sol. in H_2O . Sol. in acids. (Vauquelin.)

Thallous silicate, $3Tl_2O, 10SiO_2$.

100 pts. of a solution of Tl_2O dissolve 4.17 pts. SiO_2 by 24 hours' boiling. Sol. in H_2O . (Flemming, J. B. 1868. 251.)

Thorium silicate, ThO_2, SiO_2 .

Insol. in acids. Attacked by $KHSO_4$. (Troost and Ouyard, C. R. 105. 255.)

+ $1\frac{1}{2}H_2O$. Min. *Thorite*. Decomp. by $HCl + Aq$.

$ThO_2, 2SiO_2$. Insol. in acids or $KHSO_4$. (T. and O.)

Yttrium silicate, Y_2O_3, SiO_2 .

Attacked by HCl , HNO_3 , or $H_2SO_4 + Aq$. (Duboin, C. R. 107. 99.)

Zinc silicate, $ZnSiO_3$.

(Stein, Z. anorg. 1907, 55. 165.)
 Zn_2SiO_4 . Min. *Willemite*. Gelatinizes with $HCl + Aq$; sol. in $KOH + Aq$.

Decomp. by cold sat. citric acid + Aq . (Bolton, C. N. 1881, 43. 34.)

+ H_2O . Min. *Calamine*. Sol. in $HCl + Aq$ with separation of gelatinous SiO_2, xH_2O . Sol. in $HC_2H_3O_2 + Aq$, and $KOH + Aq$.

Insol. in $NH_4OH + (NH_4)_2CO_3 + Aq$. (Brandhorst, Zeit. angew. Ch. 1904, 17. 513.)

$ZnO, 3SiO_2$. (Bornträger, Ch. Z. 1893, 8. 186.)

Zirconium silicate, SiO_2 , ZrO_2 .

Min. *Zircon*. Insol. in acids, except H_2SO_4 , in which it is very slowly and sl. sol.

3SiO_2 , 2ZrO_2 Min. *Auerbachite*.

"Silicium oxide," $\text{Si}_2\text{H}_2\text{O}_4$.

(Geuther, J. pr. 95. 430.) This substance is identical with silicoformic anhydride according to Otto-Graham's Handb. anorgan. Chem. 7te Aufl. 2. 953.

Siliciuretted hydrogen.

See Silicon hydride.

Silicobromoform, HSiBr_3 .

Fumes on air; decomp. by H_2O .

Silicochloroform, HSiCl_3 .

Decomp. by H_2O and alcohol.

Completely miscible with CS_2 , CCl_4 , CHCl_3 , C_6H_6 , SiCl_4 , SnCl_4 , TiCl_4 , and AsCl_3 . (Ruff, B. 1905, 38. 2230.)

Silicoethane.

See Silicon hydride.

Silicofluoroform, SiHF_3 .

Decomp. by H_2O . Decomp. by NaOH and abs. alcohol with evolution of hydrogen. Decomp. by abs. ether. Sol. in toluene. (Ruff, B. 1905, 38. 63.)

Silicoformic anhydride,

$\text{H}_2\text{Si}_2\text{O}_3 = (\text{HSiO})_2\text{O}$.

Somewhat sol. in H_2O . Acids, even conc. HNO_3 + Aq, have no action, except HF , which dissolves it easily with evolution of hydrogen. Solutions of alkali hydrates, ammonium hydrate, and alkali carbonates + Aq also dissolve with evolution of hydrogen. (Ruff and Wöhler, A. 104. 101.)

Silicoiodoform, HSiI_3 .

Decomp. by H_2O . Sol. in CS_2 . (Friedel, A. 149. 96.)

Miscible with C_6H_6 and CS_2 . (Ruff, B. 1908, 41. 3739.)

Silicomethane, SiH_4 .

See Silicon hydride.

Silicomethyl chloride, SiH_3Cl .

Decomp. by H_2O and by alkalies. (Besson and Fournier, C. R. 1909, 148. 556.)

Silicomethylene chloride, SiH_2Cl_2 .

Decomp. by H_2O and by alkalies. (Besson and Fournier, C. R. 1909, 148. 556.)

Silicomolybdic acid, SiO_2 , 12MoO_3 + $26\text{H}_2\text{O}$.

Very easily sol. in H_2O and dil. acids. (Parmentier, C. R. 94. 213.)

Forms a solution with a little ether, separates into two layers by addition of or more ether. (Parmentier, C. R. 104. (Copaux, Bull. Soc. Min. 1906, 29. 79.) + $32\text{H}_2\text{O}$. Decomp. by alkali. (Asch, Z. anorg. 1901, 28. 293.)

+ $33\text{H}_2\text{O}$. (Copaux, Bull. Soc. Min. 29. 79.)

Aluminum silicomolybdate, $2\text{Al}_2\text{O}_3$, 3 12MoO_3 + $93\text{H}_2\text{O}$.

(Copaux, A. ch. 1906, (8) 7. 118.)

Aluminum sodium silicomolybdate, 4 Al_2O_3 , 2SiO_2 , Na_2MoO_4 + $7\text{H}_2\text{O}$.

Sol. in HCl + Aq. (Thugutt, Z. 1892, 2. 87.)

Ammonium silicomolybdate.

Sol. in H_2O . (Parmentier, C. R. 94. 213.)

Barium silicomolybdate, 2BaO , SiO_2 , 12 12MoO_3 + $16\text{H}_2\text{O}$.

(Copaux, A. ch. 1906, (8) 7. 118.)

+ $22\text{H}_2\text{O}$. Sol. in 4 pts. H_2O . (Copaux, Bull. Soc. Min. 1906, 29. 80.)

+ $24\text{H}_2\text{O}$. Efflorescent. Very sol. in H_2O . (Asch, Z. anorg. 1901, 28. 282.)

+ $29\text{H}_2\text{O}$. Efflorescent. (Copaux, A. 1906, (8) 7. 118.)

Cadmium silicomolybdate, 2CdO , SiO_2 , 12 12MoO_3 + $22\text{H}_2\text{O}$.

Very unusually sol. in H_2O . (Copaux, A. ch. 1906, (8) 7. 140.)

Cæsium silicomolybdate.

Sl. sol. in H_2O ; insol. in silicomolybdic acid + Aq.

Calcium silicomolybdate, 2CaO , SiO_2 , 12 12MoO_3 + $24\text{H}_2\text{O}$.

Efflorescent. Very sol. in H_2O . (Asch, Z. anorg. 1901, 28. 282.)

+ $26\text{H}_2\text{O}$. (Copaux, A. ch. 1906, (8) 7. 118.)

+ $31\text{H}_2\text{O}$. (Copaux.)

Chromium silicomolybdate, $2\text{Cr}_2\text{O}_3$, SiO_2 , 12 12MoO_3 + $93\text{H}_2\text{O}$.

(Copaux.)

Cupric silicomolybdate, 2CuO , SiO_2 , 12 12MoO_3 + $31\text{H}_2\text{O}$.

Very sol. in H_2O . (Copaux.)

Lithium silicomolybdate, $2\text{Li}_2\text{O}$, SiO_2 , 12 12MoO_3 + $29\text{H}_2\text{O}$.

Very sol. in H_2O . (Copaux.)

Magnesium silicomolybdate, 2MgO , SiO_2 , 12 12MoO_3 + $30\text{H}_2\text{O}$.

(Asch, Z. anorg. 1901, 28. 282.)

+ $31\text{H}_2\text{O}$. Very efflorescent and sol. in H_2O . (Copaux.)

um silicomolybdate, $2K_2O, SiO_2, MoO_3 + 16H_2O$.

rescent. Very sol. in H_2O . (Asch, Z. 1901, 28. 282.)

$2O, SiO_2, 12MoO_3 + 14H_2O$. (Asch.)
in H_2O with decomp. (Copaux.)

um silver silicomolybdate, $K_2O, Ag_2O, 2(SiO_2, 12MoO_3) + 14H_2O; 22H_2O; + 30H_2O$.

in H_2O with decomp. Sol. unchanged mother liquor. (Copaux, Bull. Soc. 907, 30. 293.)

um silicomolybdate.

sol. in H_2O .

silicomolybdate, $1.5Ag_2O, SiO_2, MoO_3 + 11H_2O$.

in cold H_2O . (Asch.)

$O, SiO_2, 12MoO_3 + 12H_2O$. Decomp. in H_2O . Sol. in $NH_4OH + Aq$.

$O, SiO_2, 12MoO_3 + 15H_2O$. (Asch.)

silicomolybdate, $2Na_2O, SiO_2, MoO_3 + 14H_2O$.

aux.)

$O, SiO_2, 12MoO_3 + 21H_2O$. Very sol. Efflorescent. (Asch.)

H_2O . (Copaux.)

$a_2O, SiO_2, 12MoO_3 + 17H_2O$. (Asch.)

$O, 2(SiO_2, 12MoO_3) + 17H_2O$. (Co-

um silicomolybdate, $2SrO, SiO_2, MoO_3 + 26H_2O$.

aux.)

icomolybdate, $2ZnO, SiO_2, 12MoO_3 + 12H_2O$.

very sol. in H_2O . (Copaux.)

, Si.

phous. Insol. in H_2O . Sol. before in cold HF. Insol. in other mineral and aqua regia. Sol. in conc. KOH + Aq. amorphous Si is ignited, it becomes HF and KOH + Aq.

phous Si is sol. in aqua regia and in fume of HNO_3 and HF. (Vigouroux, C. R. 1895, 120. 367.)

in liquid CO_2 . (Buchner, Z. phys. 6, 54. 674.)

in liquid NH_3 . (Gore, Am. Ch. J. 1, 830.)

itic. Sol. in $HNO_3 + HF$. (Berthel, 49. 247.)

illine. Insol. in all acids, except a trace of HF and HNO_3 . Sol. in moderately conc. $OH + Aq$ even when cold. (Deville.)

though it has been generally understood that crystallized Si is not attacked by HF, it has been found that this applies only to HF +

Aq. Gaseous HF readily attacks cryst. Si. (Newth, C. N. 1896, 72. 287.)

Si cryst. from Ag is incompletely sol. in HF. According to the temp. to which the Ag Si mixture has been heated, the following percentages of Si are dissolved in HF: 970°, 58.02%; 1150°, 27.66%; 1250°, 19%; 1470°, 16%. (Moissan and Siemens, C. R. 1904, 138. 657, 1300.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.)

Silicon amide, $Si(NH_2)_4$.

Unstable; decomp. by H_2O and partially decomp. by HNO_3 ; sol. in most organic solvents. (Lengfeld, Am. Ch. J. 1899, 21. 531.)

Decomp. by H_2O ; insol. in liquid NH_3 . (Vigouroux, C. R. 1903, 136. 1670.)

Silicon triboride, SiB_3 .

Slowly attacked by HNO_3 . Decomp. by hot conc. H_2SO_4 or fused KOH. (Moissan, C. R. 1900, 131. 142.)

Silicon hexaboride, SiB_6 .

Readily attacked by HNO_3 . Slowly decomp. by hot conc. H_2SO_4 . Not attacked by fused KOH. (Moissan, C. R. 1900, 131. 142.)

Silicon tribromide, Si_3Br_8 .

Decomp. by KOH + Aq. (Friedel and Ladenburg, A. 203. 253.)

$HSiBr_3$. See Silicobromoform.

Silicon tetrabromide, $SiBr_4$.

Rapidly decomp. by H_2O ; decomp. in several days by H_2SO_4 . (Friedel and Ladenburg, A. 147. 362.)

Silicon bromide, Si_3Br_8 .

(Besson, C. R. 1910, 151. 1056.)

Si_4Br_{10} . (Besson.)

Disilicon hydrogen pentabromide, HSi_2Br_5 or Si_2Br_5 (?).

Decomp. by H_2O . (Mahn, Zeit. Chem. (2) 5. 279.)

Silicon tetrabromide ammonia, $SiBr_4, 6NH_3$.

Decomp. by H_2O . (Lay, Dissert. 1910.)

$SiBr_4, 7NH_3$. Decomp. by H_2O . (Besson, C. R. 110. 240.)

Silicon bromoiodide, SiI_2Br_2 .

Decomp. by H_2O . Sol. in CS_2 . (Friedel, B. 2. 60.)

$SiBr_2I_2$. As above. (F.)

$SiBrI_3$. As above. (F.)

Silicon bromosulphide, $SiSBr_2$.

Decomp. in moist air. Violently decomp. by H_2O . Sol. in CS_2 and other organic solvents. (Blix, B. 1903, 38. 4218.)

Silicon carbide, SiC.

Very stable; insol. in H_2SO_4 and HNO_3 ; sol. in fused KOH at red heat. (Moissan, Bull. Soc. 1894, (3) 11. 997.)

Cryst. modification. Insol. in acids; sol. in fused alkalis. (Moissan, C. R. 1893, 117. 427.)

Insol. in all acids; sol. in molten alkalis. (Muhlhaeuser, Z. anorg. 1894, 5. 116.)

See Silundum.

Silicon subchloride, SiCl_2 (?).

Decomp. by H_2O . (Troost and Hautefeuille, A. ch. (5) 7. 463.)

Silicon trichloride, Si_2Cl_6 .

Decomp. by H_2O and alkalis. (Troost and Hautefeuille, A. ch. (5) 7. 459.)

SiHCl_3 . See Silicochloroform.

Silicon tetrachloride, SiCl_4 .

Decomp. by H_2O and alcohol.

Silicon octochloride, Si_3Cl_8 .

"Perchlorsilicopropane." Decomp. by H_2O . (Gattermann, B. 1894, 27. 1947.)

Silicon chloride, $\text{Si}_4\text{Cl}_{10}$.

(Besson, C. R. 1909, 149. 36.)

$\text{Si}_5\text{Cl}_{12}$. "Perchlorsilicobutane." Decomp. by H_2O . Fumes in the air. (Besson.)

$\text{Si}_6\text{Cl}_{14}$. "Perchlorsilicohexane." Decomp. by H_2O . (Besson, C. R. 1909, 148. 841.)

Silicon trichloride ammonia, $\text{Si}_2\text{Cl}_6, 5\text{NH}_3$.

Slowly decomp. by H_2O . (Besson, C. R. 110. 516.)

Silicon tetrachloride ammonia, $\text{SiCl}_4, 6\text{NH}_3$.

Decomp. by H_2O . (Persoz, A. ch. 44. 319.)

Silicon tetrachloride hydrazine, $\text{SiCl}_4, 4\text{N}_2\text{H}_4$.

Extremely hygroscopic and quickly decomp. by H_2O . (Lay, Dissert. 1910.)

Silicon chlorobromide, SiCl_3Br .

Decomp. by H_2O . (Friedel and Ladenburg, A. 145. 187.)

SiCl_2Br_2 . As above. (Friedel and Ladenburg.)

SiBr_3Cl . Decomp. by H_2O . (Reynolds, Chem. Soc. 51. 590.)

Silicon chlorobromide ammonia, $2\text{SiCl}_3\text{Br}, 11\text{NH}_3$.

Decomp. by H_2O . (Besson, C. R. 112. 788.)

$\text{SiCl}_2\text{Br}_2, 5\text{NH}_3$. As above. (B.)

$2\text{SiClBr}_3, 11\text{NH}_3$. As above. (B.)

Silicon chlorohydrosulphide, SiCl_3SH .

Decomp. by H_2O or alcohol. (Pierre, A. ch. (3) 24. 286.)

Silicon chloroiodide, SiCl_2I .

Decomp. by H_2O . (Besson, C. R. 1

SiCl_2I_2 . As above. (B.)

SiClI_3 . As above. (B.)

Silver chloroiodide ammonia, $2\text{SiCl}_2\text{I}$ (Besson.)

$\text{SiCl}_2\text{I}_2, 5\text{NH}_3$.

Silicon chloronitride, $\text{Si}_3\text{N}_2\text{Cl}_2$.

(Schützenberger, C. R. 92. 1508.)

Silicon chlorosulphide, $\text{Si}_2\text{Cl}_2\text{S}_2$.

Decomp. violently by H_2O . Sol (Besson, C. R. 113. 1040.)

SiSCl_2 . Violently decomp. by HCS_2 . (Blix, B. 1903, 36. 4223.)

Silicon difluoride, SiF_2 (?).

Decomp. by H_2O or $\text{NH}_4\text{OH} + \text{Ac}$ and Hautefeuille, A. ch. (5) 7. 464.

Silicon tetrafluoride, SiF_4 .

Abundantly absorbed by H_2O with 100 pts. H_2O absorb 140.6 pts. hours (Berzelius); 124.1 pts. SiF_4 (Davy).

Absorbed abundantly by E (Kuhlmann, A. 39. 319.)

Absorbed abundantly by alcohol; separation of silicic acid, if the a. contains less than 8% of water.

Sol. in conc. $\text{HF} + \text{Aq}$. Absorbed. Sl. sol. in naphtha, and oil of turp.

Silicon hydrogen fluoride, H_2SiF_6 .

See Fluosilicic acid.

Silicon fluoride with MF.

See Fluosilicate, M.

Silicon fluoride ammonia, SiF_4 .

Decomp. by H_2O . (Davy.)

Silicon hydride, SiH_4 .

Insol. in H_2O . Decomp. by I . Not changed by $\text{NH}_4\text{OH} + \text{Aq}$, H or $\text{HCl} + \text{Aq}$.

Si_2H_2 . "Silicoacetylene."

Sol. in 20% $\text{NaOH} + \text{Aq}$. with of H . (Bradley, C. N. 1900, 82. 1

Si_2H_6 . "Silicoethane." (Lebe 1909, 148. 44.)

Sl. sol. in H_2O . Best solvent orthosilicate. (Moissan, Bull. (2) 29. 443.)

Silicon nitrogen hydride, SiHN .

Decomp. by H_2O and NaOH . (1905, 38. 2241.)

oxide, $\text{SiO}_2, x\text{H}_2\text{O}$.

: acid.

See Silicooxalic acid.

See Silicoformic anhydride.

See Silicone.

e, $(\text{Si}(\text{NH})_2)$.

by H_2O . (Vigouroux, C. R. 671.)

1e, $\text{Si}(\text{NH})_2$.

by H_2O with evolution of much, B. 1903, 36. 4224.)

e hydrochloride, $\text{Si}(\text{NH}_2)_2, 2\text{HCl}$.
able in air. (Blix, B. 1903, 36.

lide, SiI_2 .

CS_2 , CHCl_3 , C_6H_6 , and SiCl_4 .
1 Ladenburg, A. 203. 247.)

dide, Si_2I_6 .

with H_2O even at 0° .

CS_2 dissolve 19 pts. Si_2I_6 at 19° ;
at 27° . (Friedel and Ladenburg,
2) 12. 92.)

ies Silicoiodoform.

iodide, SiI_4 .

by H_2O . Acts on alcohol and
 $\frac{1}{2}$ dissolves 2.2 pts. SiI_4 at 27° .
149. 96.)

ide, SiN .

decomp. by boiling with conc.

cked by dil. acids with the excep-
Decomp. by HF.

decomp. by boiling with alkalies +
s, Z. anorg. 1910, 65. 89.)

artially decomp. by boiling with
 $\frac{1}{4}$.

cked by dil. acids with the ex-
HF. Decomp. by HF.

decomp. by boiling with alkalies +
s, Z. anorg. 1910, 65. 89.)

Not attacked by H_2O .

decomp. by boiling with conc.

cked by dil. acids with the excep-
Decomp. by HF.

decomp. by boiling with alkalies +
s, Z. anorg. 1910, 65. 89.)

imide, $\text{Si}_2\text{N}_2\text{H}$.

."

HF, and rapidly in $\text{KOH} + \text{Aq}$.
berger, C. R. 92. 1508.)

. by cold, more rapidly by hot H_2O
more rapidly by alkalies. Sol. in

Not attacked by HNO_3 . Decomp.
 H_2SO_4 . (Lay, Dissert. 1910.)

Not decomposed by H_2O .

Sol. in hot alkalies + Aq with decomp.
(Blix, B. 1903, 36. 4227.)

Silicon suboxide, Si_2O_3 .

(Honigschmid, M. 1909, 30. 509.)

Silicon monoxide, SiO .

Much less easily sol. in $\text{HF} + \text{Aq}$ but more
easily sol. in alkalies + Aq than SiO_2 . (Potter,
C. C. 1907, II. 1952.)

Silicon dioxide, SiO_2 .

See also Silicic acid.

(a) *Crystalline*. Min. Quartz, Tridymite.

Insol. in H_2O , and acids, except HF.

Sl. sol. in boiling $\text{K}_2\text{CO}_3 + \text{Aq}$, and $\text{KOH} +$
Aq; see below.

Insol. in cold $\text{KOH} + \text{Aq}$; extremely slowly
sol. in boiling $\text{KOH} + \text{Aq}$. (Fuchs.)

Sol. in HF with formation of SiF_4 and H_2O .

Insol. in sugar + Aq, contrary to assertion
of Verdeil and Rissler. (Petzholdt, J. pr.
60. 368.)

(b) *Amorphous*. Min. Opal, etc.

Insol. in H_2O , and acids except HF.

100 pts. H_2O containing CO_2 dissolve
0.078 pt. amorphous SiO_2 (Maschke); 0.0136
pt. (Struckmann).

100 pts. cold $\text{HCl} + \text{Aq}$ of 1.088 sp. gr. dis-
solve 0.017 pt. SiO_2 . (Struckmann.) 100
pts. $\text{HCl} + \text{Aq}$ of 1.115 sp. gr. dissolve in the
cold 0.009 pt. SiO_2 , and 0.018 pt. on boiling.
100 pts. $\text{NH}_4\text{OH} + \text{Aq}$ (containing 10% NH_3)
dissolve 0.017 pt. quartz and 0.38 pt. ignited
 SiO_2 . (Pribram, Z. anal. 6. 119.)

Sol. in boiling K_2CO_3 or $\text{Na}_2\text{CO}_3 + \text{Aq}$,
separating out on cooling as a gelatinous
mass. (Pfaff, Schw. J. 29. 383.) The differ-
ent forms of SiO_2 have different degrees of
solubility in $\text{K}_2\text{CO}_3 + \text{Aq}$. Unignited amor-
phous SiO_2 from SiF_4 dissolves most readily,
then come opal, ignited amorphous SiO_2 ,
fused SiO_2 , and tridymite; quartz powder is
the most difficultly soluble. (Rose.) A
similar behaviour is shown to $\text{KOH} + \text{Aq}$.

Opal is much more sol. in $\text{KOH} + \text{Aq}$ than
quartz, and hyalite is the least sol. of the
varieties of opal. (Fuchs.)

Opal is easily sol. in $\text{KOH} + \text{Aq}$, even after
ignition. (Schaffgotsch, Pogg. 68. 147.)

Rammelsberg (Pogg. 112. 177) made the
following experiments on the solubility of
 SiO_2 in $\text{KOH} + \text{Aq}$. The $\text{KOH} + \text{Aq}$ used con-
tained 1 pt. KOH to 3 pts. H_2O . 1 pt. of
the powdered mineral was boiled half an hour
in a silver dish with such an amount of the
 $\text{KOH} + \text{Aq}$ that 20 pts. KOH were present.

7.75% of milky white quartz was dissolved
by repeating the above process three times.

12.8–15% of gray hornstone was dissolved
by twice boiling; 2.43% of moderately finely
powdered agate of 2.661 sp. gr. was dissolved
by once boiling; 9.7% of unignited hyalite
remained undissolved after thrice boiling;

21% of ignited hyalite remained undissolved after thrice boiling; 7.21% of semi-opal of 2.101 sp. gr. remained undissolved after thrice boiling; 18.5-19.2% of impure semi-opal of 2.101 sp. gr. remained undissolved after thrice boiling; 79.9% of chalcedony of 2.624 sp. gr. remained undissolved after thrice boiling; 6.12% of chalcedony of 2.567 sp. gr. remained undissolved after fourth boiling; 14.4% chrysoprase of 2.623 sp. gr. remained undissolved after once boiling; 49.41% of chrysoprase of 2.635 sp. gr. remained undissolved after thrice boiling; 6.62% of flint of 2.606 sp. gr. remained undissolved after twice boiling; 38.1% of fire-opal of 2.625 sp. gr. remained undissolved after fourth boiling; 26.6% of fire-opal of 2.625 sp. gr. remained undissolved after fifth boiling.

Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, **54**, 674.)

Insol. in acetone. (Naumann, B. 1904, **37**, 4329.)

The solubility of crystals of quartz on different faces in HF has been determined by Lebrun. (Belg. Acad. Bull. 1913, 953.)

Silicon thorium oxide.

See Silicate, thorium.

Silicon zirconium oxide.

See Silicate, zirconium.

Silicon oxychloride, Si_2OCl_4 .

Decomp. by H_2O and alcohol. Miscible with CS_2 , SiCl_4 , CCl_4 , CHCl_3 , or ether. (Friedel and Ladenburg, A. **147**, 355.)

$\text{Si}_4\text{O}_2\text{Cl}_{10}$; $\text{Si}_4\text{O}_4\text{Cl}_8$; $\text{Si}_8\text{O}_{10}\text{Cl}_{12}$; $(\text{Si}_2\text{O}_2\text{Cl}_2)_n$.
 $\text{Si}_4\text{O}_7\text{Cl}_2$. Sol. in above oxychlorides. (Troost and Hautefeuille, Bull. Soc. (2) **35**, 360.)

Silicon oxyfluorhydrin, $\text{Si}_2\text{O}_2\text{F}_2\text{OH}$.

(Landolt, A. Suppl. **4**, 27.)

Silicon selenide, SiSe_2 .

Decomp. by H_2O or $\text{KOH} + \text{Aq}$. (Sabatier, C. R. **113**, 132.)

Silicon sulphide, SiS_2 .

Sol. in H_2O with decomp. Acts on alcohol or ether in the cold. (Fremy, A. ch. (3) **38**, 314.)

SiS . Decomp. by H_2O ; easily sol. in dil. alkalis. (Schützenberger, Bull. Soc. (2) **38**, 56.)

Silicon sulphodiamide, $\text{SiS}(\text{NH}_2)_2$.

Slowly decomp. in the air. Insol. in cold liquid NH_3 . (Blix, B. 1903, **36**, 4219.)

Silicon sulphobromide.

See Silicon bromosulphide.

Silicon sulphochloride.

See Silicon chlorosulphide.

Silicon sulphourea, $\text{SiS}(\text{NH}_2)_2$.

Slowly decomp. in air.

Decomp. by H_2O .

Insol. in cold liquid NH_3 . (Blix, B. 1903, **36**, 4219.)

Silicone, $\text{Si}_4\text{H}_4\text{O}_8$.

Insol. in H_2O , but gives off hydrogen when warmed therewith. Not attacked by chlorine or nitric or sulphuric acids even on heating, but is gradually sol. in HF. Decomp. by alkalis, even by the most dil. $\text{NH}_4\text{OH} + \text{Aq}$, with greatest violence and evolution of heat and hydrogen gas. Insol. in alcohol, SiCl_4 , PCl_3 , or CS_2 . (Wöhler, A. **127**, 257.)

$\text{H}_2\text{Si}_2\text{O}_5$. Decomp. by H_2O and by dil acids. Violently decomp. by fuming HNO_3 . Not attacked by conc. H_2SO_4 . Very slowly decomp. by conc. HCl , rapidly by alkalis + Aq and by pyridine. (Honigschmid, M. 1909, **30**, 509.)

Insol. in H_2O , alcohol. SiCl_4 , PCl_3 , and CS_2 . Not attacked by acids except HF (Donath and Liesner, C. C. **1900**, II, 176.)

$\text{H}_{12}\text{Si}_{10}\text{O}_8$. Scarcely attacked by acids, but easily decomp. by hot H_2O , $\text{NaOH} + \text{Aq}$ etc. but not by $\text{NH}_4\text{OH} + \text{Aq}$. (Kolb, Z. anorg. **1909**, **64**, 353.)

$\text{H}_{10}\text{Si}_{10}\text{O}_8$. (Kolb.)

$\text{H}_8\text{Si}_8\text{O}_{12}$. As $\text{H}_{12}\text{Si}_{10}\text{O}_8$. (Kolb.)

Silicomesoaxalic acid, $\text{Si}(\text{OH})_2(\text{SiO.OH})_2$.

Insol. in cold H_2O , decomp. by hot H^+ (Gattermann, B. 1899, **32**, 1116.)

Silicooxalic acid, $\text{Si}_2\text{H}_2\text{O}_4 = \text{Si}_2\text{O}_2(\text{OH})_2$.

Decomp. by bases with evolution of hydrogen. Takes up HNO_3 to form compound, but not HCl or H_2SO_4 . (Troost and Hautefeuille, A. ch. (5) **7**, 463.)

Silicophosphoric acid, $\text{SiO}_2, \text{P}_2\text{O}_5$.

Slowly decomp. by H_2O . Unchanged by alcohol. Exists also in two modifications which are not attacked by H_2O . (Hautefeuille and Margottet, C. R. **99**, 789.)

$\text{SiO}_2, 2\text{P}_2\text{O}_5 + 4\text{H}_2\text{O}$. Decomp. by moist air. Sol. in H_2O at 0° , but decomp. by warming to ordinary temp. (Hautefeuille and Margottet, C. R. **104**, 56.)

Calcium silicophosphate.

See Phosphate silicate, calcium.

Silicostannic acid.

Calcium silicostannate, $\text{Ca}(\text{Si,Sn})\text{O}_3$.

Not attacked by acids, KHSO_4 , or alkalis + Aq . (Bourgeois, Bull. Soc. (2) **47**, 297.)

Silicodectungstic acid, $H_2W_{10}SiO_{38} + 3H_2O = 4H_2O, SiO_2, 10WO_3 + 3H_2O$.

Sometimes sol. in H_2O , but usually separates out gelatinous silica. (Marignac, A. ch. (4) 3. 55.)

See also **Silicoduodectungstic acid**.

Ammonium silicodectungstate, $(NH_4)_8W_{10}SiO_{38} + 8H_2O$.

Sol. in 4.5 pts. H_2O at 18° . Very sol. in hot H_2O . (Marignac, A. ch. (4) 3. 59.)

$(NH_4)_8H_2W_{10}SiO_{38} + 9H_2O$. (Marignac.)

Ammonium potassium —,

$(NH_4)_8K_4HSiW_{10}O_{38} + 15H_2O$. (Marignac.)

Barium —, $Ba_4SiW_{10}O_{38} + 22H_2O$.

Precipitate. Insol. in H_2O . (Marignac.)

Potassium —, $K_8SiW_{10}O_{38} + 17H_2O$.

Sol. in H_2O . (Marignac.)

$K_4H_4SiW_{10}O_{38} + 8H_2O$. Sol. in H_2O . (Marignac.)

Potassium — silicotungstate (?),

$K_8SiW_{11}O_{39} + 14H_2O$.

$K_4H_4SiW_{11}O_{39} + 10H_2O$. (Marignac.)

Silver —, $Ag_8W_{10}SiO_{38} + 3H_2O$.

Not appreciably sol. in cold H_2O . (Marignac, A. ch. (4) 3. 65.)

Silicotungstic acid or Silicoduodectungstic acid, $H_2SiW_{12}O_{42}$.

($H_4SiW_{10}O_{40}$, according to Copaux. (Bull. Soc. 1908, (4) 3. 101.)

+20 H_2O . Sol. in H_2O ; very sol. in alcohol; behaves with ether as the acid with 22 H_2O (Marignac, A. ch. (4) 3. 10.)

+22 H_2O . Solubility as acid with 29 H_2O . 100 pts. deliquesce with 13 pts. ether. To this mixture 20–25 pts. of ether can be added, but a further quantity no longer mixes with, but floats above the mixture. Ethereal solution is miscible with H_2O . Ether is taken up by a saturated aqueous solution with evolution of heat, until the volume has become doubled; more ether floats on the mixture. By warming the latter a liquid separates out which forms a layer between the two original layers. Alcoholic solution of the acid mixes with an equal vol. of ether, but on adding more ether a conc. ethereal solution separates as a syrupy layer. (Marignac, A. ch. (4) 3. 3. 10.)

+29 H_2O . Efflorescent. Sol. in H_2O . Saturated solution at 18° contains 1 pt. crystallized acid to 0.104 pt. H_2O , and has 2.843 sp. gr. Melts in crystal H_2O . Easily sol. in absolute alcohol and anhydrous ether.

+x H_2O . (Drechsel, B. 1887, 20. 1452.)

Aluminum silicotungstate, $Al_4H_{12}(SiW_{12}O_{42})_3 + 75H_2O$.

Not deliquescent; very sol. in H_2O . (Marignac.)

$Al_4(SiW_{12}O_{40})_3 + 60H_2O$. (Wyrouboff. Chem. Soc. 1897, 72, (2) 174.)

+87 H_2O . (Wyrouboff.)

+93 H_2O . Very efflorescent. (Wyrouboff.)

Aluminum ammonium —,

$Al_4(NH_4)_{12}(SiW_{12}O_{42})_3 + 75H_2O$.

Sol. in H_2O . (Marignac.)

Ammonium —, basic, $(NH_4)_4SiW_{12}O_{40} + 4NH_4OH + 14H_2O$.

(Wyrouboff, Chem. Soc. 1897, 72. (2) 174.)

Ammonium —, $(NH_4)_8SiW_{12}O_{42} + 16H_2O$.

Very sol. in hot H_2O . (Marignac, A. ch. (4) 3. 17.)

$(NH_4)_4H_4SiW_{12}O_{42} + 6H_2O$. Less soluble in H_2O than the preceding salt. (Marignac.)

$(NH_4)_4SiW_{12}O_{40} + 8H_2O$. (Wyrouboff, Chem. Soc. 1897, 72. (2) 174.)

Barium —, $Ba_2H_4SiW_{12}O_{42} + 14H_2O$.

Sol. in H_2O .

+22 H_2O . Gradually efflorescent. (Marignac.)

Sol. in cold H_2O , 1 : 0.7 pts. (Copaux, Bull. Soc. Min. 1906, 29. 80.)

Sol. in 4 pts. cold H_2O . (Wyrouboff, Bull. Soc. Min. 1896, 19. 278.)

$Ba_4SiW_{12}O_{42} + 27H_2O$. Nearly insol. in cold, sl. sol. in hot H_2O . (Marignac.)

$Ba_2SiW_{12}O_{40} + 16H_2O$. (Wyrouboff.)

Barium potassium —, $Ba_2K_2SiW_{12}O_{40} + 17H_2O$.

(Wyrouboff, Chem. Soc. 1897, 72. (2) 176.)

Barium sodium —, $Na_2Ba_2SiW_{12}O_{42} + 28H_2O$.

H_2O gradually dissolves out sodium silicotungstate.

Cadmium —, $Cd_2SiW_{12}O_{40} + 23H_2O$, and +27 H_2O .

(Wyrouboff.)

$4CdO, 3(SiO_2, 12WO_3) + 4H_2O$. (Wyrouboff.)

Cæsium —, $Cs_8SiW_{12}O_{42}$.

100 pts. H_2O dissolve only 0.005 pt. at 20° ; 0.52 pt. at 100° .

Completely insol. in alcohol, and $HCl + Aq$. Somewhat sol. in dil. $NH_4OH + Aq$. (Godefroy, B. 9. 1363.)

Cadmium hydrogen —, $2Cd_2SiW_{12}O_{40}, H_4SiW_{12}O_{40} + 42H_2O$.

(Wyrouboff.)

- Calcium silicotungstate**, $\text{Ca}_2\text{H}_4\text{SiW}_{12}\text{O}_{42} + 20\text{H}_2\text{O}$.
Not deliquescent. Extremely easily sol. in H_2O . (Marignac.)
 $\text{Ca}_2\text{SiW}_{12}\text{O}_{40} + 18, 24$ and $27\text{H}_2\text{O}$. (Wyruboff.)
- Calcium silicotungstate nitrate**, $\text{Ca}_2\text{SiW}_{12}\text{O}_{40}, \text{Ca}(\text{NO}_3)_2 + 15\text{H}_2\text{O}$.
(Wyruboff.)
- Cerous** —, $\text{Ce}_2\text{SiW}_{12}\text{O}_{40} + 27\text{H}_2\text{O}$.
(Wyruboff.)
 $\text{Ce}_2\text{SiW}_{12}\text{O}_{40}, \text{CeH}_2\text{SiW}_{12}\text{O}_{40} + 34\text{H}_2\text{O}$.
(Wyruboff.)
- Chromium** —, $\text{Cr}_4(\text{SiW}_{12}\text{O}_{40})_3 + 60, 87$ and $93\text{H}_2\text{O}$.
(Wyruboff.)
- Cupric** —, $\text{Cu}_2\text{SiW}_{12}\text{O}_{40} + 18, 27$ and $29\text{H}_2\text{O}$.
Very efflorescent. (Wyruboff.)
- Didymium** —, $\text{Di}_2\text{SiW}_{12}\text{O}_{40} + 26$ and $27\text{H}_2\text{O}$.
(Wyruboff.)
 $\text{Di}_2\text{SiW}_{12}\text{O}_{40}, \text{DiH}_2\text{SiW}_{12}\text{O}_{40} + 34\text{H}_2\text{O}$.
(Wyruboff.)
- Gallium** —, $\text{Ga}_4(\text{SiW}_{12}\text{O}_{40})_3 + 60, 87$, and $93\text{H}_2\text{O}$.
(Wyruboff.)
- Glucinum** —, $\text{Gl}_4(\text{SiW}_{12}\text{O}_{40})_3 + 45, 87$ and $93\text{H}_2\text{O}$.
(Wyruboff.)
- Indium** —, $\text{In}_2\text{O}_3, \text{H}_2\text{O}, 2(\text{SiO}_2, 12\text{WO}_3) + 40\text{H}_2\text{O}$.
(Wyruboff.)
 $2\text{In}_2\text{O}_3, 3(\text{SiO}_2, 12\text{WO}_3) + 63$ and $93\text{H}_2\text{O}$.
(Wyruboff.)
- Iron (ferric)** —, $\text{Fe}_4(\text{SiW}_{12}\text{O}_{40})_3 + 60$ and $93\text{H}_2\text{O}$.
(Wyruboff.)
- Lanthanum** —, $\text{La}_2\text{SiW}_{12}\text{O}_{40} + 27\text{H}_2\text{O}$.
Efflorescent. (Wyruboff.)
 $\text{La}_2\text{SiW}_{12}\text{O}_{40}, \text{LaH}_2\text{SiW}_{12}\text{O}_{40} + 34\text{H}_2\text{O}$.
(Wyruboff.)
- Lead** —, basic, $\text{Pb}_2\text{SiW}_{12}\text{O}_{40}, 2\text{PbO} + 20\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Wyruboff.)
- Lead** —, $\text{Pb}_2\text{SiW}_{12}\text{O}_{40} + 21 \text{H}_2\text{O}$.
Sol. in H_2O . (Wyruboff.)
- Lithium** —, $\text{Li}_2\text{O}, \text{SiO}_2, 12\text{WO}_3 + 14$ and $24\text{H}_2\text{O}$.
(Wyruboff.)
- Magnesium** —, $\text{Mg}_2\text{H}_4\text{SiW}_{12}\text{O}_{42} + 16\text{H}_2\text{O}$.
Stable on the air. (Marignac.)
- Mercurous silicotungstate, basic**, $\text{Hg}_2\text{SiW}_{12}\text{O}_{40}, 2\text{Hg}_2\text{O} + 5\text{H}_2\text{O}$.
Insol. in H_2O . Insol. in dil. HNO_3 . sol. in conc. warm. HNO_3 . (Wyruboff.)
- Mercurous** —, $\text{Hg}_2\text{SiW}_{12}\text{O}_{42}$.
Insol. in H_2O . Scarcely sol. in dil. l. Aq. (Marignac, A. ch. (4) 3. 43.)
- Mercuric** —, $\text{Hg}_2\text{SiW}_{12}\text{O}_{40} + 15\text{H}_2\text{O}$.
Very sol. in H_2O . Solution dec. boiling. (Wyruboff.)
- Potassium** —, basic, $\text{K}_2\text{SiW}_{12}\text{O}_{40}, 4 \text{H}_2\text{O}$.
1 pt. is sol. in 10 pts. H_2O at 18° .
boff.)
- Potassium** —, $\text{K}_2\text{SiW}_{12}\text{O}_{42} + 14\text{H}_2\text{O}$.
Sol. in 10 pts. H_2O at 18° , and 1 3 pts. at 100° . (Marignac.)
 $+ 20\text{H}_2\text{O}$. Much less sol. in cold H_2O . Extremely sol. in hot H_2O . 1 than above comp. (Marignac.)
 $\text{K}_4\text{H}_4\text{SiW}_{12}\text{O}_{42} + 7\text{H}_2\text{O}$. Solubility ceding salt.
 $\text{K}_4\text{H}_4\text{SiW}_{12}\text{O}_{42} + 16\text{H}_2\text{O}$. Sol. in 3 at 20° .
 $\text{K}_6\text{H}_{10}(\text{SiW}_{12}\text{O}_{42})_2 + 25\text{H}_2\text{O}$. Dec. dissolving in H_2O . (Marignac.)
 $\text{K}_4\text{SiW}_{12}\text{O}_{40} + 6$ and $15\text{H}_2\text{O}$. (Wyruboff.)
 $\text{K}_4\text{SiW}_{12}\text{O}_{40}, \text{K}_2\text{H}_2\text{SiW}_{12}\text{O}_{40} + 29\text{H}_2\text{O}$.
(Wyruboff.)
- Rubidium** —, $\text{Rb}_2\text{SiW}_{12}\text{O}_{42}$.
Sol. in 145–150 pts. H_2O at 20° and pts. at 100° . Insol. in alcohol; diff. in acidified, but extremely easily in al cal H_2O . (Godeffroy, B. 9. 1363.)
 $\text{Rb}_2\text{H}_2\text{SiW}_{12}\text{O}_{40} + 5\text{H}_2\text{O}$. (Wyruboff.)
 $\text{Rb}_4\text{SiW}_{12}\text{O}_{40}, \text{Rb}_2\text{H}_2\text{SiW}_{12}\text{O}_{40} + 22\text{H}_2\text{O}$.
(Wyruboff.)
- Silver** —, $\text{Ag}_4\text{H}_4\text{SiW}_{12}\text{O}_{42} + 7\text{H}_2\text{O}$.
Very sl. sol. in H_2O ; sol. in dil. l. Aq. (Marignac.)
Sl. sol. in H_2O . (Wyruboff.)
- Sodium** —, basic, $\text{Na}_2\text{SiW}_{12}\text{O}_{40}, 4 \text{H}_2\text{O}$.
(Wyruboff.)
- Sodium** —, $\text{Na}_2\text{SiW}_{12}\text{O}_{42} + 7\text{H}_2\text{O}$.
The saturated solution at 19° conta pt. H_2O to 1 pt. of the salt dried at 10 has sp. gr. = 3.05. (Marignac.)
 $\text{Na}_4\text{H}_4\text{SiW}_{12}\text{O}_{42} + 11\text{H}_2\text{O}$. Stable at $+ 18\text{H}_2\text{O}$. Efflorescent. (Marignac.)
 $\text{Na}_2\text{H}_2\text{SiW}_{12}\text{O}_{42} + 14\text{H}_2\text{O}$. Decomp. solving in H_2O . (Marignac.)
 $\text{Na}_4\text{SiW}_{12}\text{O}_{40} + 14, 16$ and $20\text{H}_2\text{O}$. (boff.)

silicotungstate nitrate, \cdot
 \cdot $\text{H}_4\text{SiW}_{12}\text{O}_{42}$, $4\text{NaNO}_3 + 39\text{H}_2\text{O}$.
 y deliquescent. (Marignac.)

n —, $\text{Sr}_2\text{SiW}_{12}\text{O}_{40} + 16, 17, 23$ and
 \cdot O .
 uboff.)

—, $\text{Ti}_2\text{H}_2\text{SiW}_{12}\text{O}_{40} + 9\text{H}_2\text{O}$.
 uboff.)

—, basic.
 in H_2O . (Wyrouboff.)

—, $\text{Th}_2\text{SiW}_{12}\text{O}_{40} + 27\text{H}_2\text{O}$.
 ol. in H_2O . (Wyrouboff.)
 $\text{V}_{12}\text{O}_{40}$, $2\text{H}_4\text{SiW}_{10}\text{O}_{40} + 45\text{H}_2\text{O}$.
 off.)

—.
 an salts are described by Wyrouboff.

—.
 salts are described by Wyrouboff.

—, $\text{Zn}_2\text{SiW}_{12}\text{O}_{40} + 18, 27$ and $29\text{H}_2\text{O}$.
 uboff.)

silicovanadiomolybdic acid.

am silicovanadiomolybdate,
 H_4O , SiO_2 , V_2O_5 , $9\text{MoO}_3 + 20\text{H}_2\text{O}$.
 of sat. solution containing 0.32016 g.
 ccm. at $18^\circ = 1.21322$. (Friedheim,
33. 1624.)

\cdot O , SiO_2 , V_2O_5 , $10\text{MoO}_3 + 21\text{H}_2\text{O}$.
 f sat. solution containing 0.35026 g.
 1 ccm. at $18^\circ = 1.25275$. (Fried-

\cdot O , V_2O_5 , SiO_2 , $11\text{MoO}_3 + 27\text{H}_2\text{O}$.
 f sat. solution containing 0.38086 g.
 ccm. at $18^\circ = 1.29266$. (Friedheim.)

\cdot O , V_2O_5 , SiO_2 , $15\text{MoO}_3 + 24\text{H}_2\text{O}$.
 f sat. solution containing 0.48997 g.
 ccm. at $18^\circ = 1.43761$. (Friedheim.)

um potassium —, $(\text{NH}_4)_2\text{O}$, $2\text{K}_2\text{O}$,
 \cdot V_2O_5 , $9\text{MoO}_3 + 20\text{H}_2\text{O}$.

of sat. solution containing 0.24021
 1 ccm. at $18^\circ = 1.17031$. (Fried-

\cdot O , $2\text{K}_2\text{O}$, SiO_2 , V_2O_5 , $10\text{MoO}_3 +$
 Sp. gr. of sat. solution contain-
 14 g. salt in 1 ccm. at $18^\circ = 1.19184$.
 m.)

\cdot O , $2\text{K}_2\text{O}$, SiO_2 , V_2O_5 , $11\text{MoO}_3 +$
 Sp. gr. of sat. solution contain-
 14 g. salt in 1 ccm. at $18^\circ = 1.21378$.
 m.)

um zinc —, $4(\text{NH}_4)_2\text{O}$, 2ZnO ,
 \cdot $3\text{V}_2\text{O}_5$, $18\text{MoO}_3 + 15\text{H}_2\text{O}$.
 in H_2O . (Blum, Dissert. 1904.)

Silicovanadiotungstic acid.

Ammonium silicovanadiotungstate, $3(\text{NH}_4)_2\text{O}$,
 SiO_2 , V_2O_5 , $9\text{WO}_3 + 24\text{H}_2\text{O}$.

Can be cryst. from H_2O . (Friedheim, B.
 1902, **35. 3244.**)

$(\text{NH}_4)_6\text{SiV}_2\text{W}_{10}\text{O}_{40} + 21\text{H}_2\text{O}$. 1 cc. of sat.
 solution in H_2O at 17.5° contains 0.6652 g.
 of the hydrated salt. Sp. gr. of this solution =
 1.4505. Decomp. by conc. acid and alkali.
 (Friedheim.)

Ammonium barium potassium —,
 $(\text{NH}_4)_2\text{K}_2\text{BaSiV}_2\text{W}_{10}\text{O}_{40} + 25\text{H}_2\text{O}$.

Sl. sol. in H_2O . Decomp. by conc. acids
 and alkalies. (Friedheim.)

Ammonium potassium —,
 $(\text{NH}_4)\text{K}_2\text{SiV}_2\text{W}_{10}\text{O}_{40} + 23\text{H}_2\text{O}$.

1 ccm. of sat. solution at 17.5° contains
 0.5072 g. of the salt. Sp. gr. of the solution
 at $20^\circ = 1.3462$. Can be cryst from H_2O .
 Decomp. by conc. acids and alkalies. (Fried-
 heim.)

Barium —, $\text{Ba}_2\text{SiV}_2\text{W}_{10}\text{O}_{40} + 28\text{H}_2\text{O}$.

1 ccm. of the sat. solution in H_2O at 17.5°
 contains 0.0384 g. of the salt. Sp. gr. of the
 solution = 1.0307. Decomp. by conc. acids
 and alkalies. (Friedheim, B. 1902, **35. 3245.**)

6BaO , 2SiO_2 , $3\text{V}_2\text{O}_5$, $18\text{WO}_3 + 50\text{H}_2\text{O}$.
 Sl. sol. in H_2O . (Friedheim.)

7BaO , 2SiO_2 , $3\text{V}_2\text{O}_5$, $18\text{WO}_3 + 83\text{H}_2\text{O}$.
 Sl. sol. in H_2O . (Friedheim.)

Potassium —, $\text{K}_6\text{SiV}_2\text{W}_{10}\text{O}_{40} + 22\text{H}_2\text{O}$.

Sol. in H_2O . Can be cryst. from H_2O
 without decomp. Decomp. by conc. acids
 and alkalies. (Friedheim.)

$6\text{K}_2\text{O}$, 2SiO_2 , $3\text{V}_2\text{O}_5$, $18\text{WO}_3 + 31\text{H}_2\text{O}$. Sol.
 in H_2O . (Friedheim.)

$7\text{K}_2\text{O}$, 2SiO_2 , $3\text{V}_2\text{O}_5$, $18\text{WO}_3 + 42\text{H}_2\text{O}$. Sol.
 in H_2O . (Friedheim.)

Sodium —, $\text{Na}_6\text{SiV}_2\text{W}_{10}\text{O}_{40} + 29\text{H}_2\text{O}$.

Very sol. in H_2O . Decomp. by conc acids
 and alkalies. (Friedheim.)

Silundum, Si_xC_y .

Not attacked by hot Cl or conc. acids.
 (Amberg, Z. Elektrochem. 1909, **15. 725.**)

Silver, Ag.

Not attacked by H_2O . Absolutely insol. in
 HCl or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Lea, Sill. Am. J.
144. 444.) Easily sol. in $\text{HNO}_3 + \text{Aq}$ on warm-
 ing, if not too conc. Only a minute trace is
 dissolved in an hour by cold dil. $\text{HNO}_3 + \text{Aq}$ (1
 pt. $\text{HNO}_3 + \text{Aq}$ of sp. gr. 1.40 : 10 pts. H_2O).
 (Lea.) Sol. in hot conc. H_2SO_4 with evolution
 of SO_2 . Sl. sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ (1 : 4), but
 with more dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ the different forms
 of Ag behave differently. (Lea.)

Sol. in $\text{HI} + \text{Aq}$ at ordinary temperature.

Sol. in KI + Aq with access of air. Sol. in hot KCN + Aq. (Christomanos, Z. anal. 7. 301.)

Sol. in chromic, iodic, chloric and bromic acids. Dil. H_2SO_4 alone is incapable of dissolving finely divided Ag, and the seeming solvent action is due to the oxygen of the air, oxygen dissolved in the acid, or derived from some external source. (Hendrixson, J. Am. Chem. Soc. 1903, 25. 637.)

Boiling H_2SO_4 dissolves pure Ag only when concentration equals 60°B . More dil. acid dissolves only the impure metal. (Pannani, Gazz. ch. it. 1909, 39. (2) 234.)

Slowly decomp. into AgCl by alkali chlorides + Aq; also by CuCl_2 , etc. + Aq.

Somewhat sol. in NH_4OH + Aq in presence of O. (Lea, Sill. Am. J. 144. 444.)

Sol. in KMnO_4 + dil. H_2SO_4 + Aq. (Friedheim, B. 20. 2554.)

Sol. in $\text{Fe}_2(\text{SO}_4)_3$ + Aq, especially on heating, but completely insol. in FeSO_4 + Aq. (Vogel.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Allotropic forms—(a). Very sol. in H_2O . Solution is pptd. by saline solutions or almost any neutral substance. Alkali sulphates, nitrates, and citrates ppt. it in a sol. form, while MgSO_4 , CuSO_4 , FeSO_4 , NiSO_4 , $\text{K}_2\text{Cr}_2\text{O}_7$, $\text{K}_4\text{Fe}(\text{CN})_6$, $\text{Ba}(\text{NO}_3)_2$, and even AgNO_3 + Aq ppt. it in an insol. form, which, however, may be made sol. again by treatment with many substances, as $\text{Na}_2\text{B}_4\text{O}_7$, K_2SO_4 , or Na_2SO_4 + Aq. NaNO_3 + Aq ppts. the Ag from its solution in a perfectly insol. form.

(β). The ppt. from aqueous solution by salts is sol. in NH_4OH + Aq. (Lea, Sill. Am. J. 137. 476.)

Many other allotropic forms exist. (Lea.)

Pure colloidal silver is also sol. in alcohol. (Schneider, B. 25. 1164.)

Entirely sol. in H_2O , even when dry. (Schneider, Z. anorg. 1894, 7. 339.)

Silver acetylide, Ag_2C_2 .

Sol. in KCN + Aq with evolution of C_2H_2 . Decomp. by HCl + Aq. (Arth, C. R. 1897, 124. 1535.)

Silver amide, AgNH_2 .

Ppt.; sol. in ammonium salts + Aq and in excess of potassium amide. Sol. in liquid NH_3 . Insol. in Ag salts + Aq. (Franklin, J. Am. Chem. Soc. 1905, 27. 833.)

Sol. in excess of KNH_2 . (Franklin, Z. anorg. 1905, 46. 16.)

Silver antimonide, Ag_3Sb or Ag_4Sb .

Min. *Discrasite*. Sol. in HNO_3 + Aq. Ag_3Sb . Insol. in HCl + Aq; decomp. by HNO_3 + Aq. (Christofle.)

Silver azoimide, AgN_3 .

Insol. in hot or cold H_2O or dil. acids; sol. in conc. mineral acids. Sol. in NH_4OH + Aq. (Curtius, B. 23. 3023.)

Silver bromide, AgBr .

Insol. in H_2O , or H_2O acidulated HNO_3 , H_2SO_4 , or $\text{HC}_2\text{H}_3\text{O}_2$ between 33° . If flocculent or pulverulent, it is sol. therein above 33° , but if granular above 50° , and then very slightly. (Sch. (5) 3. 289.) Ag can be detected in 10,000,000 pts. H_2O . (Stas.)

Calculated from the electrical conductivity of AgBr + Aq, AgBr is sol. in 1,971.6 H_2O at 20.2° , and 775,400 pts. at 38° . (Pannani, Z. phys. Ch. 12. 133.)

By same method Kohlrausch and R. calculate that 1 l. H_2O dissolves 0.4 mg at 18° . (Z. phys. Ch. 12. 240.)

Solubility in H_2O = 0.109 mg. per l. homme, J. chim. Phys. 9. 519.)

Solubility in H_2O = 6.6×10^{-7} at 25° . (Win, Z. phys. Ch. 1894, 13. 645.)

Solubility of AgBr in H_2O at 25° 8.1×10^{-7} g. mols. per l. (Thiel, Z. 1900. 24. 57.)

Aq. solution sat. at 21.1° contains 10^{-6} gr. equiv. per litre. (Kohlrausch 1901, II. 1299.)

1 l. H_2O dissolves 0.000137 g. AgBr (Abegg and Cox, Z. phys. Ch. 1903, 29. 108.)

0.84×10^{-4} g. are dissolved per litre solution at 20° . (Böttger, Z. phys. Ch. 46. 603.)

1 l. H_2O dissolves 0.107 mg. AgBr (Kohlrausch, Z. phys. Ch. 1904, 30. 108.)

3.7 mg. AgBr are contained in 1 lit solution at 100° . (Böttger, Z. phys. Ch. 1906, 56. 93.)

1 mg. in 1 l. of sat. solution at 21° (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

Solubility in H_2O = 8.8×10^{-7} per litre at 25° . (A. E. Hill, J. Am. Soc. 1908, 30. 74.)

Boiling H_2O dissolves 0.0000035 weight of AgBr . HNO_3 + Aq (1°) dissolves 0.00000543 of its weight of 100° with sl. decomposition. The is pptd. by AgNO_3 + Aq or HBr (or Aq, but not completely. 1 pt. of solution requires 3 pts. of Br as HBr), or of Ag as AgNO_3 in order to precipitated. (Stas.)

Not attacked by boiling HNO_3 , sol. in conc. HBr or HCl + Aq. Boiling conc. H_2SO_4 decomposes it hardly acts on it (Dumas), dissolve quantity, which is repptd. by H_2S (Stas.)

Very sl. sol. in dil., easily in conc. + Aq. 100 pts. NH_4OH + Aq (0.98) dissolve 0.51 pt. AgBr (dried at 100° and about double that amount of pptd. AgBr . (Pohl, W. A. B. 41. 108.)

1 g. freshly pptd. AgBr is sol. in 10% NH_4OH + Aq, but insol. in an aq. solution of AgCl . (Seiner, F. Trans. (3) 14. 1.)

1 g. AgBr dissolves in 8779.4 g. H_2O + Aq (sp. gr. = 0.998) at 12° , and in

$\text{NH}_4\text{OH} + \text{Aq}$ (sp. gr. = 0.96) at 12° .
agi, Gass. ch. it. 13. 87.)

Solubility of AgBr in $\text{NH}_4\text{OH} + \text{Aq}$ at 15° .
G. mols. per l.

NH_3	AgBr	Sp. gr. 15.5°
0.085	0.0011	0.9932
0.365	0.0031	0.9853
0.410	0.0050	0.9793
0.590	0.0074	0.9720
0.725	0.0101	0.9655

Bodländer, Z. phys. Ch. 1892, 9. 734.)

Solubility in $\text{NH}_4\text{OH} + \text{Aq}$ at 0° .

100 ccm. of the solution contain g NH_3 and
Ag Br.

NH_3	Mg. AgBr	g. NH_3	Mg. AgBr
0.307	8.0	2.627	106.7
0.488	9.6	3.126	156.8
0.669	17.2	3.389	198.7
0.829	21.2	3.652	266.9
0.151	34.9	3.722	288.8
0.532	55.7	3.770	293.0
0.809	72.2	3.926	289.2
0.953	74.1	3.995	285.0

(Jarry, A. ch. 1899, (7) 17. 364.)

Solubility of AgBr in $\text{NH}_4\text{OH} + \text{Aq}$ at 25° .
G. mols. per l.

NH_3	AgBr	NH_3	AgBr
1932	0.00060	1.965	0.00692
3849	0.00120	3.024	0.01163
7573	0.00223	5.244	0.02443

Bodländer and Fittig, 1902, Z. phys. Ch. 39.
597.)

Solubility in $\text{NH}_4\text{OH} + \text{Aq}$ at 25° .

g. at. Ag per l.	Mols. NH_3 per l.
0.00170	0.450
0.00159	0.497
0.000941	0.268
0.00107	0.273
0.000391	0.115
0.000386	0.118
0.000278	0.0764
0.000264	0.0777

Atney and Melcher, J. Am. Chem. Soc.
1903, 25. 79.)

sl. in hot $\text{NH}_4\text{Cl} + \text{Aq}$. Very sl. sol. in
carbonate, sulphate, or succinate + Aq,
still less in nitrate. (Wittstein.) Not
easily sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ when sus-
pended in much H_2O , and is separated out
by $\text{KBr} + \text{Aq}$. (Field, C. N. 3. 17.)
sl. in $\text{KCN} + \text{Aq}$. Sl. sol. in conc. KCl ,

KBr , NaCl , NaBr , NH_4Cl , or $\text{NH}_4\text{Br} + \text{Aq}$;
but insol. when dilute.

Traces only dissolve in alkali nitrates + Aq.
(Fresenius, Quant. Anal.)

Abundantly sol. in $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$. 100
ccm. H_2O containing 10 ccm. normal
 $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$ dissolve 0.0383 g. AgBr.
(Stas.)

Solubility of AgBr in $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$ at 25° .
G. mols. per l.

$\text{Hg}(\text{NO}_3)_2$ HNO_3	AgBr	$\text{Hg}(\text{NO}_3)_2$ HNO_3	AgBr
1	0.03660	0.025	0.00459
0.10	0.00873	0.0125	0.00329
0.05	0.00639	0.0100	0.00306

HNO_3 was present in all cases, and it was
found that there was no difference in solubili-
ty of AgBr with concentrations between
0.1N and 2N HNO_3 . Cryst. and amorphous
AgBr showed the same solubility. (Morse,
Z. phys. Ch. 1902, 45. 708.)

Difficultly sol. in hot conc. $\text{AgNO}_3 + \text{Aq}$.
(Riese, A. 111. 39.)

100 ccm. of a 3-N solution of AgNO_3 dis-
solve 0.04 g. AgBr at 25° . Much less sol.
in $\text{AgNO}_3 + \text{Aq}$ than AgI. (Hellwig, Z.
anorg. 1900, 25. 176.)

Solubility in $\text{AgNO}_3 + \text{Aq}$.

Volumetric measurements

AgNO_3	N/10 KBr ccm.	Opal- escent at	G. AgBr re- tained per 100 g. AgNO_3
10 g. made up to 32 g. per 100 g. of solution	0.65	22°	0.129
	0.72	35	0.144
	0.8	44	0.159
	0.9	62	0.178
	1.0	67	0.188
	1.1	77	0.207
	1.2	79	0.226
	6.0	37	1.13
10 g. made up to 70 g. per 100 g. of solution	8.0	53	1.50
	10.0	67	1.88
	11.25	72	2.12
	12.0	74	2.26
	12.75	79	2.40
	13.5	82	2.54
	15.5	85.5	2.92
	17.5	90	3.29

Gravimetric measurements at 14.5°

G. AgNO_3	ccm H_2O	Strength of AgNO_3 %	G. AgBr re- tained per 100 g. AgNO_3
7.326	9.32	44	0.144
8.290	7.65	52	0.185
7.255	4.84	60	0.283
7.35	3.95	65	0.365

(Lowry, Roy. Soc. Proc. 1914, 91. A. 65.)

100 g. KBr in conc. KBr+Aq dissolve 3019 mg. AgBr at 15°; 95 g. NaCl+10 g. KBr dissolve only 75 mg. AgBr at 15°. (Schierholz, W. A. B. 101, 2b. 4.)

Solubility in KBr+Aq at 25°.

Mol. KBr in 1 litre	G. AgBr in 1 litre
4.864	26.44
4.44	17.95
4.18	13.50
3.68	7.50
2.81	2.34
2.76	2.20

(Hellwig, Z. anorg. 1900, 25. 183.)

Sol. in conc. KBr or NaBr+Aq (Lowig), but less than AgI in KI+Aq (Field).

100 g. NaCl in conc. NaCl+Aq dissolve 474 mg. AgBr at 15°; 100 g. NaCl in 21% NaCl+Aq dissolve 188 mg. AgBr at 15°. (Schierholz, W. A. B. 101, 2b. 4.)

Solubility of AgBr in Na₂SO₃+Aq at 25°. G. formula weights per l.

So ₃	Ag	So ₃	Ag
0.232	0.0025	0.466	0.0053
0.406	0.0023	0.474	0.0055
0.448	0.0023	0.675	0.0084

(Luther and Leubner, Z. anorg. 1912, 74. 393.)

Solubility of AgBr in Na₂SO₃ at (?)°. (g. salts per l. of solution.)

Na ₂ SO ₃	AgBr	Na ₂ SO ₃	AgBr
83.75	0.790	2.08	0.0159
70.75	0.570	1.13	0.0086
38.2	0.265	0.59	0.0045
17.65	0.116	0.3	0.0039
9.47	0.0526	0.17	0.0022
4.85	0.0329	0.08	0.00075

(Mees and Piper, Photog. J. 1912, 36. 234.)

Solubility in Na₂S₂O₃+Aq at 35°.

g. Na ₂ S ₂ O ₃ in 1 liter	g. AgBr corresponding to each g. Na ₂ S ₂ O ₃ .
100	0.376
200	0.390
300	0.397
500	0.427

(Richards and Faber, Am. Ch. J. 1899, 21. 169.)

(NH₄)₂S₂O₃+Aq. dissolves AgBr more rapidly than does Na₂S₂O₃+Aq. (Lumière and Seyewitz, C. C. 1906, II. 1138.)

Solubility in salts+Aq.

Solvent	% Conc.	AgBr sol. in 1 l. of 1% solution
Sodium thiosulphate	1	0.3
" " "	5	1.9
" " "	10	3.5
" " "	15	4.2
" " "	20	5.8
Sodium sulphite	10	0.0
" "	20	0.0
Ammonium sulphite	10	Tr
Potassium cyanide	5	6.5
Ammonium sulphocyanide	5	0.5
" " "	10	2.0
" " "	15	5.5
Potassium " "	10	0.
Calcium " "	10	0.
Barium " "	10	0.
Aluminum " "	10	4.
Thiocarbamide	10	1
Thiosinamine	1	0
"	5	0
"	10	0

(Valenta, M. 1894, 15. 250)

Solubility of AgBr in salts+A (G. AgBr sol. in 1 l. of 1% solution)

NaSCN	2.06
NH ₄ SCN	0.03
(NH ₄) ₂ CO ₃	0.004
Na ₂ SO ₄	0.055

(Mees and Piper, Photog. J., 1912)

In a solution of NaC₂H₃O₂+Aq, 10 ccm. of sat. NaC₂H₃O₂+Aq, 20 ccm. normal HC₂H₃O₂+Aq, 970 ccm. H₂O, about double the amount of AgBr is dissolved in the solution. This solution required 3 pts. of Aq. ppt. the AgBr in solution. Pure granular AgBr are wholly insol. in conc. acetates+Aq. (Stas.)

Sol. in Hg(C₂H₃O₂)₂+Aq. 100 ccm. H₂O containing 10% Hg(C₂H₃O₂)₂+Aq dissolves 0.012 g. AgBr at 20°. (Schierholz.)

Very sol. in liquid NH₃. (Fra Ch. J. 1898, 20. 829.)

Solubility in 10 cc, methylamine at different concentrations at 11.5°.

g. NH ₂ CH ₃	4.844	4.311	3.5
mg. AgBr	289	127	75

g. NH ₂ CH ₃	1.797	1.513	1.3
mg. AgBr	28	16	15

(Jarry, A. ch. 1890, (7) 17. 3)

methylamine + Aq at 25°.

G. mol. per l.	
	AgBr
	0.00026
	0.00034
	0.000395
	0.00041
	0.00045

h, B. 1902, **35**. 2416.)

gBr in methylamine + Aq at 25°.

G. mol. per l.		
AgBr	CH ₃ NH ₂	AgBr
0025	0.102	0.00026
0013	0.051	0.00012
00049		

l Eberlein, B. 1903, **36**. 3948.)

in ethylamine + Aq at 25°.

G. mol. per l.	
	AgBr
	0.0000867
	0.000137
	0.000193
	0.000258
	0.000711

h, B. 1902, **35**. 2416.)

l. ethylamine + Aq containing C₂H₅NH₂ dissolves 0.00231 g. 00 g. mol. C₂H₅NH₂, 0.0097 g. (Bodländer and Eberlein, B.)

benzonitrile. (Naumann, B.)

stone. (Eidmann, C. C. 1899, Naumann, B. 1904, **37**. 4329.)

thoholic thiourea. (Reynolds, 92, **61**. 251.)

thyl acetate. (Bezold, Dissert. ann, B. 1909, **42**. 3790.) ethyl

iers, Dissert. 1906); (Naumann 14.)

is dissolved in 1 l. of 1% thio- q. (Mees and Piper, Photog. 9.)

rm pyridine.

terminated in piperidine. (Wer- 1897, **15**. 16.)

rite, Bromite.

ammonia, AgBr, NH₃.

l Crozier, C. R. 894, **118**. 1150.)

AgBr, 1½NH₃. (Jarry, A. ch. 1899, (7) 17. 356.)

2AgBr, 3NH₃. (Joannis and Crozier.)

AgBr, 3NH₃. Decomp. by H₂O. Sl. sol. in liquid NH₃. (Jarry.)

AgBr, 5NH₃. (Jarry, C. R. 1898, **126**. 1141.)

Silver carbide, Ag₂C.

(Gay-Lussac.)

Ag₂C(?). Sol. in HNO₃ + Aq with residue of C. (Liebig, A. **38**. 129.)

Ag₂C₂. Sol. in HNO₃ + Aq with residue of C. (Regnault, A. **19**. 153.)

Silver subchloride, Ag₂Cl₂.

NH₄OH + Aq dissolves the greater part, the residue (20%) being sol. in HNO₃ + Aq. KCN dissolves the greater part; H₂SO₄ dissolves about 2%; acetic acid and KOH are without action. (Bibra, J. pr. 1875, (2) **12**. 52.)

Argentous chloride, Ag₂Cl.

Obtained in a pure state by Guntz (C. R. **112**. 861). Dil. HNO₃ + Aq does not attack but warm conc. HNO₃ + Aq decomp. Easily sol. in KCN + Aq. (Guntz, C. R. **112**. 1212.)

The following data are for a more or less impure Ag₂Cl.

Boiling conc. HCl + Aq, NaCl + Aq, or NH₄OH + Aq dissolve out AgCl, and leave Ag. (Scheele, Wetzlar. Dulk, Wöhler.)

According to Berthollet, wholly sol. in NH₄OH + Aq. Sol. for the most part in NH₄OH + Aq, and the residue is sol. in HNO₃ + Aq (= Ag + AgCl). (v. Bibra, B. 7. 741.)

Silver chloride, AgCl.

Nearly insol. in H₂O.

When AgCl is left in contact for some hours with pure H₂O at 20-22°, and especially at 75°, traces go into solution; more Cl is dissolved than Ag. When 1 pt. Ag is pptd. as AgCl in presence of 1 million pts. H₂O a slight bluish milkyiness is observed; but in order to have a distinct ppt. 4 pts. Ag should be present.

Dil. HNO₃ + Aq does not increase the solubility of AgCl, but AgCl is not absolutely insol. in stronger HNO₃ + Aq. (Mulder.)

1 pt. AgNO₃, when mixed with HCl + Aq in presence of 120,000 (Pfaff), 240,000 (Harting), pts. H₂O, causes an opalescence.

1 pt. Ag gives a slight turbidity with HCl + Aq in presence of 200,000 pts. H₂O, a scarcely opalescent cloudiness with 400,000 pts. H₂O, and the same after the lapse of 15 minutes in presence of 800,000 pts. H₂O. (Lassaigne.)

1 pt. Ag can be detected as AgCl in 1 million parts H₂O at ordinary temp., but not in 2 million parts. In NaNO₃ + Aq containing 0.79 pt. NaNO₃ in 200,000 pts. H₂O, 1 pt. Ag can be detected as AgCl. This dissolves at 75°, and is visible again on cooling.

If the same liquid contains 1574 pts. NaNO₃, the AgCl remains in solution after cooling.

In 100 ccm. H_2O containing 0.787 g. $NaNO_3$, 13 drops of $NaCl$ and silver solution, each drop of which contains 0.05 mg. Ag , cause a precipitate at 5° , 20 drops at $15-17^\circ$, 60 drops at $45-55^\circ$.

$AgCl$ is somewhat less sol. in $HNO_3 + Aq$ than in $NaNO_3 + Aq$ when the amount of H_2O remains the same.

Therefore, if HCl is used instead of $NaCl$, about $\frac{1}{7}$ less $AgCl$ remains in solution.

In 100,000 pts. of H_2O , which contain HNO_3 , and an amount of HCl corresponding to the amount of Ag salt, 1.596 pts. $AgCl$ dissolve at 25° . The solution is precipitated by either $AgNO_3$ or HCl . (Mulder, Silber Probir-methode, Leipzig, 1859. 62.)

(For further older data, see Storer's Dictionary.)

White flaky $AgCl$ is appreciably sol. in hot H_2O , 1000 ccm. boiling H_2O dissolving about 2 mg. $AgCl$. Far less sol. in H_2O containing $AgNO_3$, being practically insol. in H_2O containing 0.1 g. $AgNO_3$ in a litre. Solubility is also diminished one-half by addition of HCl . (Cooke, Sill. Am. J. (3) 21. 220.)

Solubility in H_2O rapidly diminishes as the temp. falls. (Cooke, *l. c.*)

Not completely insol. in H_2O . According to Stas (C. R. 73. 998) there are four modifications: (1) gelatinous; (2) cheesy-flocculent; (3) pulverulent; (4) granular, crystalline, or fused. (4) is almost absolutely insol. in H_2O at the ordinary temp., but the solubility increases with the temp., and is considerable at 100° ; (2), which is formed by the precipitation of a cold dilute Ag solution, has the greatest solubility in pure H_2O , and it changes its solubility by standing, or if made pulverulent by shaking with H_2O ; (3) is also sol. in H_2O ; the solution of (2) or (3) in pure H_2O , or H_2O acidified with HNO_3 , is precipitated by $AgNO_3$ or $NaCl + Aq$.

In order to ppt. 1 pt. $AgCl$ in above solution 3 pts. of Cl as chloride or Ag as nitrate are necessary; the pptn. is then complete.

Solubility of granular variety in boiling H_2O is proportionately large, and pptn. is brought about by 3 pts. Cl or Ag as above, but the pptn. in this case is not complete.

The salts formed simultaneously with the $AgCl$ have no influence on the solubility of the $AgCl$. Presence of HNO_3 does not increase the solubility of (2), but has that effect on (3) in proportion to the amt. of HNO_3 present. (Stas, C. R. 73. 998.)

Further determination by Stas are as follows:—

Between 0° and 30° granular $AgCl$ is insol. in pure H_2O , or H_2O acidulated with HNO_3 .

Between 0° and 30° the flocculent and pulverulent forms of $AgCl$ dissolve without alteration in pure H_2O , in acidulated H_2O , in alkali acetates + Aq , and in $Hg(C_2H_3O_2)_2 + Aq$ containing an alkali acetate. Their degree of solubility is a function of the state of the chloride, of the temp., and of the nature

and quantity of the solvent within limits of temp. ($0^\circ-30^\circ$). These soln. they contain either Ag in the state of salt, or Cl as chloride or HCl in an amount three times that which they can dissolve. $AgCl$, exercise no solvent action on any modifications of $AgCl$. And reciprocal $AgCl + Aq$ is pptd. instantly by a dec. solution of $AgNO_3$ or MCl (or HCl). $AgCl$ is wholly pptd. when the quantity of the Ag or Cl thus added is equal to times the quantity of the Ag or Cl dissolved as $AgCl$.

Between 50° and 100° , however, dec. solutions of Ag or chlorides, which give instant ppts. in solutions sat. with any modifications of $AgCl$, do not eliminate the dissolved $AgCl$. At 100° , they contain 60% of the amt. dissolved. (Stas, A. 3. 323.)

Calculated from electrical conductance $AgCl + Aq$, $AgCl$ is sol. in 715,800 pts. at 13.8° , and 384,100 pts. at 26.5° . (van't Hoff, Z. phys. Ch. 12. 132.)

Calculated in the same way, $l.$ dissolves 0.76 mg. at 2° ; 0.97 mg. at 18° ; 2.24 mg. at 26° ; 3.03 mg. at 42° . (Kohlrausch and Fieser, Z. phys. Ch. 12. 242.)

Solubility in $H_2O = 1.25 \times 10^{-5}$ mol. (Goodwin, Z. phys. Ch. 1894, 13. 64)

Solubility of $AgCl$ in H_2O at $25^\circ = 10^{-5}$ (in normality). (Thiel, Z. anorg. Ch. 24. 57.)

2.16×10^{-5} moles are sol. in 1 litre at 25° . (Noyes and Kohr, Z. phys. Ch. 42. 341.)

1.53×10^{-3} g. per liter are dissolved in aq. solution at 20° . (Böttger, Z. anorg. Ch. 1903, 46. 603.)

1 l. H_2O dissolves 1.6 mg. $AgCl$ (Kohlrausch, Z. phys. Ch. 1904, 50. 168.)

21.8 milligrams are dissolved in sat. solution at 100° . (Böttger, Z. anorg. Ch. 1906, 56. 93.)

1.34 mg. are contained in 1 l. of sat. solution at 18° . (Kohlrausch, Z. phys. Ch. 64. 168.)

1 l. sat. solution at t° contains mg

t°	mg. $AgCl$	t°
1.55	0.56	17.51
4.68	0.66	25.86
9.97	0.89	34.12

(Kohlrausch, Z. phys. Ch. 1908, 6)

Solubility in $H_2O = 1.6 \times 10^{-5}$ g. per litre at 25° . (A. E. Hill, J. Am. Chem. Soc. 1908, 30. 74.)

1 l. H_2O dissolves 0.00154 g. $AgCl$ at 100° . (Whitby, Z. anorg. Ch. 67. 108.)

lated from electrical conductivity of Aq, 1 l. H₂O dissolves:

05 milli-equivalents AgCl at 18°.
65 " " " " 50°.
7 " " " " 100°.

her, J. Am. Chem. Soc. 1910, 32. 55.)

I₂O dissolves 1.02×10⁻⁵ g. equiv. at 19×10⁻⁵ g. equiv. at 25°. (van Rossen, 112, I. 1539.)

most probable average value for solu-
AgCl in H₂O is 1.04×10⁻⁵ g. equiv.
18°, and 1.43×10⁻⁵ g. equiv. per l.
(van Rossen, C. C. 1912, II. 1807.)

10⁻⁵ g. AgCl are sol. in 1 l. H₂O at
Howczynski, C. A. 1915, 741.)

sol. in H₂O than AgSCN. (Normand,
Soc. 1912, 101. 1853.)

a conc. HCl+Aq, and also when not
ic.; thus the solution of 1 pt. AgNO₃+
5,000 pts. H₂O is clouded by a little
q, but clears up by the addition of
Reinsch, J. pr. 13. 133.)

AgCl dissolves in 200 pts. conc. HCl+
in 600 pts. HCl+Aq diluted with 2
(Pierre, J. Pharm. (3) 12. 237.)

what sol. in hot alcohol, to which HCl
added, but is precipitated on cooling.
an, J. pr. 19. 341.)

ts. sat. HCl+Aq (sp. gr. 1.165) dis-
2980 pt. AgCl, or AgCl is sol. in 336
l+Aq at ord. temp.; 100 pts. HCl+
gr. 1.165) at b.-pt. dissolve 0.56 g.
AgCl is sol. in 178 pts. HCl+Aq.

ility of AgCl in dil. HCl+Aq. 100
Cl+Aq (sp. gr. 1.165), to which the
nt. H₂O has been added, dissolve g.

	ccm. H ₂ O	g. AgCl	Pts. HCl which dis- solve 1 pt. AgCl
	10	0.056	1,785
	20	0.018	5,555
	30	0.0089	11,235
	50	0.0035	18,571

ogel, N. Rep. Pharm. 23. 335.)

Cl is added to a solution in which
pt. Ag is suspended, the milkiness dis-

Solubility in HCl+Aq increases
temp., the AgCl separating out on
(Mulder.)

amounts of AgCl which dissolve in
q are directly proportional to the
of acid (of fixed concentration) used.

, J. Am. Chem. Soc. 1906, 28. 1448.)

HCl dissolves 0.0002 g. AgCl at 21°.
HCl dissolves 0.0033 g. AgCl at 21°.
HCl dissolves 0.0555 g. AgCl at 21°.
Whitby, Z. anorg. 1910, 67. 108.)

Solubility in HCl+Aq at 25°.	
HCl g.-equivalents per l.	Ag × 10 ⁻³ g.-equivalents per l.
0.649	0.032
1.300	0.126
1.911	0.266
2.149	0.374
2.569	0.610
2.975	0.814
3.576	1.358
4.182	2.147
4.735	3.168
5.508	4.126

(Forbes, J. Am. Chem. Soc. 1911, 33. 1941.)

Solubility in 20% HCl +Aq.			
HCl	N/10 AgNO ₃ ccm.	Opalescent at t°	G. AgCl to 100 g. anhy- drous HCl
20 g. of 20% acid	1.0
	1.1	0.0	0.39
	2.0	29.5	0.72
	3.0	51.5	1.076
	3.75	70.0	1.346
	4.25	82.0	1.525
	4.75	90.0	1.74
	5.80	107.0	2.08

(Lowry, Roy, Soc. Proc. 1914, 91. A. 62.)

Sl. sol. in conc. HBr+Aq. (Löwig.)
Insol. in HNO₃+Aq. (Wackenroder.)
Entirely unacted upon by HNO₃ of 1.43
sp. gr. (Wurtz, Am. J. Sci. (2) 25. 382.)

Solubility in dil. HNO₃+Aq is the same as
solubility in H₂O, i. e. 1/2,000,000 pt. of Ag can-
not be detected in H₂O with or without HNO₃,
but 1/1,000,000 pt. can be detected in both cases.
(Mulder.)

1 pt. Ag in the form of AgCl dissolves at
25° in 83,000 pts. H₂O containing free HNO₃,
and 0.33 pt. of HCl. (Mulder.)

100,000 pts. conc. HNO₃+Aq dissolve
about 2 pts. AgCl, and solubility is not sen-
sibly affected by lower nitrogen oxides.
(Thorpe, Chem. Soc. (2) 10. 453.)

Solubility of AgCl in HNO ₃ +Aq at 25°.			
G. per liter			
HNO ₃	AgCl	HNO ₃	AgCl
0.0315	0.001647	18.9	0.00225
0.063	0.001705	94.5	0.0245
0.630	0.00176		

(Glowczynski, Kolloidchem. Beih. 1914, 6.
147.)

Insol. in cold conc. H_2SO_4 , but on boiling is in part decomp. and in part dissolved, and does not separate on cooling.

$AgCl$ is not more sol. in dil. $H_2SO_4 + Aq$ than in dil. $HNO_3 + Aq$.

Unacted upon by cold $H_2SO_4 + Aq$, and but slightly decomp. on heating. (Vogel.)

Abundantly sol. in $H_2PtCl_6 + Aq$ without decomp. (Birnbäum, Z. Ch. 1867. 520.)

Insol. in cold dil. caustic alkalis + Aq but decomp. by hot conc. solutions. (Gregory.)

Decomp. by $K_2CO_3 + Aq$.

Sl. sol. in cold $K_2CO_3 + Aq$.

Easily sol. even in dil. $NH_4OH + Aq$.

1 pt. $AgCl$ dissolves in 1288 pts. $NH_4OH + Aq$ of 0.89 sp. gr. (Wallace and Lamont, Chem. Gaz. 1893. 137.)

100 pts. $NH_4OH + Aq$ of 0.986 sp. gr. dissolve at 80° 1.492 pts. $AgCl$, dried at 100° . (Pohl, W. A. B. 41. 627.)

1 l. $NH_4OH + Aq$ of 0.949 sp. gr. dissolves 51.6 g. Ag as freshly precipitated $AgCl$, and 47.6 g. when diluted with 1 l. H_2O .

1 l. $NH_4OH + Aq$ of 0.924 sp. gr. dissolves 58 g. Ag as freshly precipitated $AgCl$; 1 l. $NH_4OH + Aq$ of 0.899 sp. gr. dissolves 49.6 g.; 0.5 l. $NH_4OH + Aq$ (of 0.049 sp. gr.) + 0.5 l. saturated $NaCl + Aq$ dissolves 20.8 g.; 0.5 l. $NH_4OH + Aq$ (of 0.949 sp. gr.) + 0.5 l. saturated $KCl + Aq$ dissolves 20.4 g.; 0.5 l. $NH_4OH + Aq$ (of 0.949 sp. gr.) + 0.5 l. saturated $NH_4Cl + Aq$ dissolves 22.4 g. Ag as freshly pptd. $AgCl$. (Millon and Commaile, C. R. 56. 309.)

1 g. $AgCl$ dissolves in 428.64 g. 5% $NH_4OH + Aq$ (sp. gr. 0.998) at 12° ; 1 g. $AgCl$ dissolves in 12.76 g. 10% $NH_4OH + Aq$ (sp. gr. 0.96) at 18° . (Longi, Gazz. ch. it. 13. 87.)

1 g. freshly pptd. $AgCl$ is sol. in 17 cc. 10% $NH_4OH + Aq$. Solubility is diminished by presence of $AgBr$. (Senier, Pharm. J. Trans. (3) 14. 1.)

Solubility in $NH_4OH + Aq$ at 0° .

G. per 100 g. solution.

NH_3	$AgCl$	NH_3	$AgCl$
1.45	0.49	28.16	5.69
1.94	1.36	29.80	7.09
5.00	3.44	30.19	7.25
6.24	4.00	32.43	5.87
11.77	4.68	34.56	4.77
16.36	5.18	37.48	3.90

(Jarry, A. ch. 1899, (7) 17. 342.)

Solubility in $NH_4OH + Aq$ increases with the temp. (Jarry.)

Solubility in $NH_4OH + Aq$ at 25°

g. at Ag per l.	Mols. NH_3 per l.	g. at Ag per l.	Mols. NH_3 per l.
0.151	2.042	0.0140	0.253
0.149	2.017	0.0140	0.253
0.149	2.013	0.0140	0.252
0.147	1.991	0.0139	0.252
0.0616	0.961	0.00621	0.115
0.0583	0.916	0.00621	0.115
0.0584	0.909	0.00619	0.115
0.0572	0.903	0.00625	0.115
0.0569	0.896	0.00304	0.059
0.0555	0.873	0.00297	0.058
0.0541	0.863	0.00300	0.058
0.0514	0.818	0.00149	0.025
0.0249	0.428	0.00143	0.025
0.0240	0.416	0.00142	0.025
0.0235	0.411	0.00141	0.025
0.0227	0.397		

(Whitney and Melcher, J. Am. Chem. 1903, 25. 78.)

Solubility of $AgCl$ in $NH_4OH + Aq$ at :

$Ag = g$ at Ag in 1000 g. H_2O .

$NH_3 = g$ mol NH_3 in 1000 g. H_2O .

Conc. = Molecular concentration of NH_3 .

Ag	NH_3	Conc.	Solid phase
0.023	0.437	0.391	$AgCl$
0.025	0.428	0.378	"
0.1197	1.700	1.461	"
0.1308	1.688	1.426	"
0.372	3.782	3.038	"
0.378	3.945	3.181	"
0.574	5.10	3.95	"
0.609	5.33	4.11	"
0.633	5.545	4.279	"
0.745	6.26	4.77	$AgCl + 2AgCl, 2N$
0.754	6.27	4.76	"
0.757	6.25	4.74	"
0.760	6.25	4.73	"
0.775	6.52	4.97	$2AgCl, 3NH_3$
0.848	8.28	6.58	"
0.968	11.19	9.25	"
0.980	11.78	9.82	"
0.978	12.23	10.27	"
0.965	12.26	10.33	"
1.03	12.68	10.62	"
1.09	12.96	10.78	"
1.049	14.34	12.24	"
1.039	14.47	12.39	"

(Straub, Z. phys. Ch. 1911, 77. 332.)

Easily (Brett), difficultly (Wittmann), in $NH_4Cl + Aq$, but not in other NH_4 salts.

Solubility in NH₄Cl + Aq at 15°.

% NH ₄ Cl	% AgCl
10.0	0.0050
14.29	0.0143
17.70	0.0354
19.23	0.0577
21.98	0.110
25.31	0.228
28.45	0.340*
Sat.	0.177

5°.
Schierholz, W. A. B., 1890, 101. 2b. 8.)
Solubility in NH₄Cl + Aq (26.31%) at t°.

t°	% AgCl
15	0.276
40	0.329
60	0.421
80	0.592
90	0.711
100	0.856
110	1.053

(Schierholz.)

25°, 1 l. NH₄Cl + Aq containing 0.00053 H₂Cl dissolves 0.001604 g. AgCl; 0.00530 H₂Cl, 0.002379 g. AgCl. (Glowczynski, Kolloidchem. Beih. 1914, 6. 147.)

also Forbes, page 826.

KClO₃ + Aq dissolves 1.8 mg. (Guye, Ann. Chim. Phys. 10. 145.)

sol. in conc. KCl + Aq, NaCl + Aq, and in other chlorides.

LiCl, KCl, NH₄Cl, CaCl₂, ZnCl₂ + Aq, etc., dissolve appreciable quantities of AgCl, especially if hot and concentrated, but it separates out for the most part on cooling.

sol. in solutions of all the metallic chlorides which are sol. in H₂O, thus NaCl, KCl, CaCl₂, LiCl, and BaCl₂ + Aq all dissolve AgCl, especially if hot. MgCl₂, NH₄Cl, and HgCl₂ (at) also dissolve AgCl. (Mulder.)

sol. in conc. CaCl₂ + Aq. (Wetzlar.)

sol. in roseocobaltic chloride + Aq. (Gibbs, Z. Chem. 1891, 1. 100.)

sol. in SnCl₄, HgCl₂, CuCl₂, ZnCl₂, CdCl₂, NiCl₂, or CoCl₂ + Aq. (Vogel.)

Solubility of AgCl in sat. solutions of chlorides at ordinary temperatures.

Salt	100 pts. sat. solution dissolve pts. AgCl	Pts. solution required to dissolve 1 pt. AgCl
Li ₂	0.0143	6,993
Na ₂	0.0884	1,185
K ₂	0.0930	1,075
Li	0.0950	1,050
Na	0.0475	2,122
KCl	0.1575	634
Li ₂	0.1710	584
Na ₂	0.2980	336

(Vogel, N. Rep. Pharm. 23. 335.)

Experiments by Hahn give different results from those of Vogel as follows:—

Solubility in various salts + Aq.

Salt	% salt	Sat. at t°	% AgCl
KCl	24.95	19.6	0.0776
NaCl	25.96	"	0.1053
NH ₄ Cl	28.45	24.5	0.3397
CaCl ₂	41.26	"	0.5713
MgCl ₂	36.35	"	0.5313
BaCl ₂	27.32	"	0.0570
FeCl ₃	0.1686
FeCl ₂	0.0058
MnCl ₂	24.5	0.1996
ZnCl ₂	0.0134
CuCl ₂	24.5	0.0532
PbCl ₂	"	0.0000

(Hahn, Wyandotte Silver Smelting Works, 1877.)

1 l. 4-N KCl + Aq dissolves 0.915 g. KCl at 25°. (Hellwig, Z. anorg. 1900, 25. 166.)

Solubility in KCl + Aq at t°.

t°	G. equiv. per l.	
	Ag × 10 ⁻³	KCl
1.0	1.734	3.325
25.0	2.415	3.083
35.0	2.786	2.955

(Forbes, J. Am. Chem. Soc. 1911, 33. 1937.)

Solubility in KCl + Aq at 25°.
G. per liter.

KCl	AgCl	KCl	AgCl
0.00236	0.00184	0.01491	0.00305
0.00471	0.00218	0.02984	0.00321

(Glowczynski, Kolloidchem. Beih. 1914, 6. 147.)

Solubility in CaCl₂ + Aq.

t°	G. equiv. per l.	
	Ag × 10 ³	$\frac{\text{CaCl}_2}{2}$
1.0	0.964	3.512
25.0	1.514	3.320
35.0	1.806	3.221

(Forbes, l. c.)

Sat. CuCl₂ + Aq at 0° dissolves 2.835 g. AgCl per l; at 100°, 8.147 g. Solubility in sat. MgCl₂ + Aq is still greater. (Hahn, Eng. Min. J. 65. 434.)

More sol. in $\text{HgCl}_2 + \text{Aq}$ than in H_2O . (Finzi, Gazz. ch. it. 1902, 32. (2) 324.)
At 15° , 100 g. NaCl in 280 ccm. H_2O dissolve 485 mg. AgCl ; 100 g. KCl in 300 ccm. H_2O dissolve 334 mg.; 100 g. NH_4Cl in 280 ccm. H_2O dissolve 1051 mg.
The solubility decreases with dilution rapidly at first until about an equal vol. of H_2O has been added, and then much more slowly to a minimum quantity, when the dilution is 1 : 10 for NaCl and KCl , and 1 : 20 for NH_4Cl .
100 g. NaCl in 280 ccm. H_2O dissolve 2170 mg. AgCl at 109° ; 100 g. NH_4Cl in 280 ccm. H_2O dissolve 4000 mg. AgCl at 110° ; 100 g. NaCl in 620 ccm. H_2O (14% solution) dissolve 15 mg. AgCl at 15° , and 774 mg. at 104° . (Schierholz, W. A. B. 101, 2b. 4.)
The solubility of AgCl in $\text{NaCl} + \text{Aq}$ decreases with diminishing concentration of $\text{NaCl} + \text{Aq}$. (Barlow, J. Am. Chem. Soc. 1906, 28. 1448.)

Solubility in $\text{NaCl} + \text{Aq}$.

Gravimetric measurements, 15°	
Strength of salt solution	G. AgCl retained per 100 g. NaCl
15% NaCl	0.063
20% NaCl	0.134
28% NaCl	0.279

Volumetric measurements			
NaCl	N/10 AgNO_3 ccm.	Opalescent at t°	G. AgCl retained per 100 g. NaCl
20 g. of 15% solution	0.25	28	0.119
	0.4	40	0.191
	0.7	64	0.335
	1.0	78	0.478
	1.25	89	0.598
	1.7	102.5	0.812
20 g. of 20% solution	0.43	17.0	0.156
	0.65	26.0	0.234
	0.82	37.0	0.295
	1.2	51.5	0.430
	1.6	67.0	0.524
	2.12	79.5	0.765
	2.52	88.5	0.910
	3.08	97.0	1.10
20 g. of 28% solution	3.52	105.0	1.27
	2.25	36.5	0.675
	2.75	45.0	0.704
	3.5	56.0	0.896
	4.5	69.0	1.153
	5.5	84.0	1.411
	6.5	94.0	1.664
	7.75	107.5	1.958

Solubility in salts + Aq at 25° .
 C = concentration of the salt in salt solution in g.-equivalents per litre.

Salt	C	$\text{Ag} \times 10^{-4}$ g.-equivalents per
NaCl	0.933	0.086
	1.190	0.130
	1.433	0.184
	1.617	0.245
	1.871	0.348
	2.094	0.446
	2.272	0.570
	2.449	0.684
	2.658	0.851
	2.841	1.040
	3.000	1.194
	3.270	1.583
	3.471	1.897
	3.747	4.462
	3.977	2.879
	4.170	3.335
CaCl_2 2	4.363	3.810
	4.535	4.298
	5.039	6.039
	1.748	0.289
	2.201	0.501
	2.741	0.900
	3.264	1.463
	3.737	2.182
NH_4Cl	4.033	2.802
	4.538	4.175
	5.005	5.823
	0.513	0.042
	0.926	0.113
	1.141	0.172
	1.574	0.365
	2.143	0.842
	2.566	1.425
	2.918	2.160
	3.162	2.795
	3.510	4.029
SrCl_2 2	4.363	9.353
	4.902	14.92
	5.503	24.04
	5.764	30.17
	0.550	0.033
	0.989	0.082
NH_4Cl	1.359	0.173
	1.572	0.236
	1.698	0.284
	1.818	0.348
	2.140	0.510
	2.476	0.747
	2.992	1.252
	3.494	2.018
	4.152	3.594
	5.216	8.174
	5.775	12.04

(Lowry, Roy, Soc. Proc. 1914, 91. A, 61.)

solubility in salts + Aq. at 25°—Continued.

lt	C	Ag×10 ⁻³ g.-equivalents per l.
	1.111	0.141
	1.425	0.235
	1.713	0.391
	2.022	0.616
	2.396	1.050
	2.628	1.390
	2.850	1.845
	3.081	2.435
	3.424	3.602
	3.843	5.725
l ₂	1.248	0.186
	1.610	0.339
	2.676	1.274
	3.260	2.366

Forbes, J. Am. Chem. Soc. 1911, 33, 1940).

sl. in NaNO₃, KNO₃, Ca(NO₃)₂, Mg(NO₃)₂, NH₄NO₃ + Aq; sl. sol. at ord. temp., but solubility is much increased by heat.

Solubility in NaNO₃ + Aq at 15–20°.

mm. H ₂ O	g. NaNO ₃	mg. AgCl dissolved
100	0.787	1.33
200	0.787	1.93
300	2.361	3.99
100	2.787	2.53

Solubility increases with ascending temp.

Temp.	ccm. H ₂ O	g. NaNO ₃	mg. AgCl dissolved
5°	100	0.787	0.86
5–17°	100	0.787	1.33
18°	100	0.787	1.46
30°	100	0.787	2.33
5–55°	100	0.787	3.99

(Mulder.)

At 25°, 100,000 pts. H₂O containing a little HNO₃ and 0.787 g. NaNO₃ dissolve 2.128 AgCl. By adding 2 g. more NaNO₃ to the same solution, 2.5269 mg. (1/3 more) AgCl are dissolved. (Mulder.)

Solubility in H₂O is not appreciably increased by 1/10 N to N-KNO₃ or NH₄NO₃ + (van Rossen, C. C. 1912, II. 1807.)

In presence of NaNO₃ and excess of HCl, H₂O dissolves 0.03 mg. AgCl. (Richards & Wells.)

Hg(NO₃)₂ + Aq dissolves considerable quantities of AgCl, but the other nitrates do not. (Mulder.)

Much more sol. in hot than in cold (NO₃)₂ + Aq, and much more sol. therein than in NH₄NO₃ + Aq. NaCl ppts. AgCl in this solution; much less sol. therein in

presence of NaC₂H₃O₂ or NH₄OH + Aq. AgCl is pptd. from above solution by NaC₂H₃O₂ + Aq. (Mulder.)

Sol. in Hg(NO₃)₂ + Aq (Wackenroder, A. 41. 317); in considerable amount (Liebig, A. 81. 128); and is precipitated by HCl, NH₄Cl, NaCl, KC₂H₃O₂ (Debray, C. R. 70. 849); incompletely precipitated by AgNO₃ and not by HNO₃ (Wackenroder).

Solubility of AgCl in Hg(NO₃)₂ + Aq at 25°.
(G. mols. per l.)

Hg(NO ₃) ₂ HNO ₃	AgCl	Hg(NO ₃) ₂ HNO ₃	AgCl
0.0100	0.00432	0.050	0.00914
0.0125	0.00499	0.100	0.01395
0.025	0.00690	1.000	0.04810

HNO₃ was present in all cases, and it was found that there was no difference in solubility of AgCl with concentrations between 0.1N and 2N HNO₃. (Morse, Z. phys. Ch. 1902, 45. 708.)

Not sol. to appreciable extent in Cu(NO₃)₂, Fe₂(NO₃)₆, Mn(NO₃)₂, Co(NO₃)₂, Zn(NO₃)₂, or Ni(NO₃)₂ + Aq; insol. or exceedingly sl. sol. in Pb(NO₃)₂ + Aq. (Mulder.)

Imperfectly sol. in AgNO₃ + Aq. (Wackenroder.)

Conc. AgNO₃ + Aq dissolves AgCl perceptibly.

Less sol. in AgNO₃ + Aq than AgBr. (Risse, A. 111. 39.)

Solubility in 0.02N AgNO₃ + Aq = 0.15 × 10⁻⁷ g. mols. per l. (Böttger.)

100 ccm. of 3-N solution of AgNO₃ dissolve 0.08 g. AgCl at 25°. More dil. solutions dissolve very slight amounts of AgCl. (Hellwig, Z. anorg. 1900, 25. 177.)

Solubility in 2-N AgNO₃ + Aq at ord. temp. = 0.03 × 10⁻³ g. equiv. AgCl. (Forbes, J. Am. Chem. Soc. 1912, 33. 1946.)

Solubility in AgNO₃ + Aq at t°.

(Det. by volumetric method.)

AgNO ₃ :H ₂ O = 2:1			
G. AgNO ₃	n/10 NaCl ccm.	t°	g. AgCl retained per 100 g. AgNO ₃
6	2	57	0.478
7	2	45	0.410
8	2	40	0.359
9	2	35	0.319
11	2	30	0.261
7	1	26	0.205
10	1	22	0.143
10	4	65	0.572
10	5	86	0.715

Solubility in AgNO₃ + Aq at t°.—Continued.

AgNO ₃ :H ₂ O = 1:1			
5	1	94	0.286
6	1	84	0.239
7	1	75	0.205
8	1	66	0.179
9	1	58	0.159
5.5	0.5	48	0.130
6.5	0.5	40	0.110
12	0.5	23	0.060

AgNO ₃ :H ₂ O = 1:2			
6	0.5	104	0.120
7	0.5	92	0.103
8	0.5	85	0.090
10	0.5	73	0.072
12	0.5	61	0.060
8	0.25	45	0.045
12	0.25	28	0.030

(Lowry, Roy, Soc. Proc. 1914, 91. A. 58.)

Solubility in AgNO₃ + Aq at 20°. (Det. by gravimetric method.)

g. AgNO ₃	g. H ₂ O	g. AgCl retained per 100 g. AgNO ₃
220	110	0.1372
220	165	0.1009
220	220	0.0722
220	330	0.0402
220	440	0.0294

(Lowry, Roy. Soc. Proc. 1914, 91. A. 56.)

Insol. in Na₂SO₄ + Aq.

Solubility of AgCl in Na₂SO₃ + Aq at 25°. G. formula weights per l.

SO ₃	Ag	SO ₃	Ag
0.080	0.011	0.483 *	0.059 *
0.106	0.017	0.470	0.070
0.220	0.033	0.652	0.103
0.234	0.036	0.890	0.140
0.478 *	0.057 *	0.937	0.142

* In presence of 0.05 Cl. (Luther and Leubner, Z. anorg. 1912, 74. 393.)

Easily sol. in Na₂S₂O₃ or KCN + Aq. When freshly pptd., very sol. in solutions of soluble thiosulphates, and especially in conc Na₂S₂O₃ + Aq, which dissolves AgCl almost as readily as H₂O dissolves sugar. K₂S₂O₃ + Aq, even when very dil., also dissolves AgCl; also SrS₂O₃ + Aq. (Herschel, 1819.) Sol. in KAsO₂ + Aq. (Reynoso.) Cold NaHSO₃ + Aq dissolves a considerable amount of AgCl. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 78.) Sol. in cold sat. (NH₄)₂S₂O₃ + Aq. (Rosen-

heim and Steinhäuser, Z. anorg. 1900, 25. 103.)

Solubility in Na thiosulphate + Aq at 16°.

g. Na ₂ S ₂ O ₃ ·5H ₂ O in 100 cc. water	g. dissolved AgCl	
	experimental	calculated
2.08	0.29	0.80
4.16	0.64	1.60
6.24	0.88	2.40
8.35	1.26	3.21
16.70	2.54	6.42
20.83	3.28	7.99

(Abney, Z. phys. Ch. 1895, 18. 65.)

A solution of Na₂S₂O₃ + Aq containing 200 g. Na₂S₂O₃ per liter, dissolves 0.454 g. AgCl per g. of Na₂S₂O₃ at 35°. (Richards and Faber, Am. Ch. J. 1899, 21. 170.)

Solubility in salts + Aq.

Solvent	% Conc.	Grams AgCl sol. in 100 grams solvent
Sodium thiosulphate	1	0.40
" " "	5	2.00
" " "	10	4.10
" " "	20	6.10
Ammonium thiosulphate	1	0.57
" " "	5	1.32
" " "	10	3.92
Sodium sulphite	10	0.44
" " "	20	0.95
Ammonium sulphite	10	Trace
" carbonate	10	0.06
Ammonia + Aq	3	1.40
"	15	7.58
Magnesium chloride	50	0.50
Potassium cyanide	5	2.75
Ammonium sulphocyanide	5	0.08
" " "	10	0.54
" " "	15	2.88
Potassium	10	0.11
Calcium	10	0.15
Barium	10	0.30
Aluminum	10	2.02
Thiocarbamide	10	0.83
Thiosinamine	1	0.40
"	5	1.90
"	10	3.90

(Valenta, M. 1894, 15. 250.)

Solubility in salts + Aq.

31.71 cc. of a solution of sodium thiosulphate containing 31.869 g. Na₂S₂O₃ per liter (i. e. 5 g. of the hydrate in 100 cc. of the solution) dissolve 0.6124 g. AgCl. 21.88 cc. of a solution of ammonium thiosulphate containing 50 g. (NH₄)₂S₂O₃ per liter dissolve 0.7024 g. AgCl. 27.34 cc. of a solution of potassium cyanide containing 49.511 g. KCN per liter dissolve

5 g. AgCl. (Cohn, Z. phys. Ch. 1895, 1.)

solubility of AgCl in sodium thiosulphate and potassium cyanide solutions may be determined without reference to experimental data. (Cohn.)

sol. in liquid NH_3 . (Franklin, Am. Ch. Soc. Trans. 18, 20. 829.)

sol. in moderately dil. $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$. 1 ccm. normal $\text{Hg}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$ contains 1 g. Hg dissolve 0.01892 g. AgCl at 15°.

10 ccm. of a solution of a mixture of Na and Hg acetates dissolve 0.00175 g. AgCl. (A. ch. (5) 3. 145.)

sl. sol. in liquid NH_3 .

solubility curve for AgCl, AgCl , 3NH_3 , 5NH_3 . (Jarry, A. ch. 1899, 17. 342.)

sol. in alcoholic ammonia. (Bodländer, Z. phys. Ch. 1892, 9. 731.)

early insol. in ether. (Mylius and Hüttig, Z. 1911, 44. 1316.)

slightly sol. on warming with solution of tartaric acid, but nearly the whole is dissolved on cooling.

sol. in acetone. (Naumann, B. 1904, 37. 1014.) ; insol. in acetone and in methylal. (Naumann, C. C. 1899, II. 1014.)

sol. in methyl acetate. (Bezold, Dissert. 1890, Naumann, B. 1909, 42. 3790.)

sol. in ethyl acetate. (Hamers, Dissert. 1910, Naumann, B. 1910, 43. 314.)

sol. in methylamine + Aq. (Wurtz, A. ch. 1850, 453.)

solubility of AgCl in methylamine at 11.5°.

CH_3NH_2	% AgCl	% CH_3NH_2	% AgCl
78	0.16	13.70	3.29
44	0.62	18.69	5.43
51	0.83	36.69	9.93
66	1.32		

(Jarry, A. ch. 1899, (7) 17. 342.)

solubility in methylamine + Aq at 25°.
G. mols. per l.

CH_3NH_2	AgCl
0.0200	0.000300
0.0400	0.000370
0.0740	0.000424
0.0947	0.000447
0.1950	0.000481

(Wuth, B. 1902, 35. 2416.)

solubility in methylamine + Aq at t° .
G. mols. per l.

CH_3NH_2	Ag
0.93	0.0315
0.93	0.0338
0.93	0.0335

(Euler, B. 1903, 36. 2880.)

At 25°, 1 l. methylamine + Aq, containing 1.017 g. mols. CH_3NH_2 , dissolves 0.0387 g. mol. AgCl; 0.508 g. mol. CH_3NH_2 , 0.0178 g. mol. AgCl. (Bodländer and Eberlein, B. 1903, 30. 3948.)

Solubility in ethylamine + Aq at 25°.
G. mols. per l.

$\text{C}_2\text{H}_5\text{NH}_2$	AgCl
0.01272	0.000114
0.03942	0.000156
0.05512	0.000235
0.06572	0.000312
0.10300	0.000824

(Wuth, B. 1902, 35. 2416.)

Solubility in ethylamine + Aq at t° .
G. mols. per l.

t°	$\text{C}_2\text{H}_5\text{NH}_2$	Ag
18	0.094	0.00458
25	0.093	0.00474
25	0.094	0.00478
18	0.236	0.0132
25	0.234	0.0136
18	0.462	0.0251

(Euler, B. 1903, 36. 2880.)

At 25°, 1 l. ethylamine + Aq, containing 0.483 g. mol. $\text{C}_2\text{H}_5\text{NH}_2$, dissolves 0.0314 g. mols. AgCl; 0.200 g. mol. $\text{C}_2\text{H}_5\text{NH}_2$, 0.0115 g. mol. AgCl; 0.100 g. mol. $\text{C}_2\text{H}_5\text{NH}_2$, 0.0062 g. mol. AgCl. (Bodländer and Eberlein.)

Sol. in amylamine + Aq, but less than in $\text{NH}_4\text{OH} + \text{Aq}$.

Sol. in caprylamine + Aq.

Easily sol. on warming in ethylene diamine + Aq. (Kurnakow, Z. anorg. 1898, 17. 220.)

Easily sol. in alcoholic solution of thiocetamide. (Kurnakow, J. pr. 1895, (2) 51. 251.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Solubility in pyridine at t° .

t°	g. AgCl sol. in 100 g. pyridine	Solid phase
-52	0.70	AgCl, $2\text{C}_5\text{H}_5\text{N}$
-49	0.77	
-35	0.99	
-30	1.36	
-25	1.80	
-22	2.20	
transition point	2.75	AgCl, $\text{C}_5\text{H}_5\text{N}$
-20	3.71	
-18	3.85	
-10	4.35	
-5	5.05	
-1	5.60	

Solubility in pyridine at t° .—*Continued.*

t°	g. AgCl sol. in 100 g. pyridine	Solid phase
transition point	...	
0	5.35	AgCl
10	3.17	
20	1.91	
30	1.20	
40	0.80	
50	0.53	
60	0.403	
70	0.32	
80	0.25	
90	0.22	
100	0.18	
110	0.12	

(Kahlenberg, J. phys. Chem. 1909, **13**. 423.)Easily sol. in warm piperidine. (Varet, C. R. 1892, **115**. 335.)Mol. wt. determined in piperidine. (Werner, Z. anorg. 1897, **15**. 16.)Quinoline dissolves traces of AgCl. (Varet, C. R. 1893, **116**. 60.)As sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$ as in $\text{NH}_4\text{OH} + \text{Aq}$. (Blyth, Chem. Soc. **1**. 350.)Sol. in sinamine, and thiosinamine + Aq.
Min. *Cerargyrite*.**Silver chloride ammonia**, $\text{AgCl} \cdot 2\text{NH}_3$.Decomp. by H_2O . (Terreil, A. Phys. Beibl. **7**. 149.) $2\text{AgCl} \cdot 3\text{NH}_3$. Decomp. on air and in H_2O to AgCl. Sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$, from which it can be crystallised. (Rose.)Insol. in alcohol. (Bodländer, Z. phys. Ch. **9**. 730.) $\text{AgCl} \cdot 3\text{NH}_3$. More easily decomp. than $2\text{AgCl} \cdot 3\text{NH}_3$.Sl. sol. in liquid NH_3 . (Jarry, A. ch. 1899, (7) **17**. 343.) $\text{AgCl} \cdot 5\text{NH}_3$. Sl. sol. in liquid NH_3 . (Jarry, A. ch. 1899, (7) **17**. 336.)**Silver chlorobromiodides.**(Rodwell, Proc. Roy. Soc. **25**. 292.)**Silver subfluoride (argentous fluoride)**, Ag_2F .Decomp. by H_2O into Ag and AgF. (Guntz, C. R. **110**. 1337.)Decomp. by H_2O .Insol. in abs. alcohol, ether, acetone and xylene. (Wöhler and Rodewald, Z. anorg. 1909, **61**. 63.)Decomp. by H_2O until the solution contains 64.5% AgF, independent of temp. (Guntz, C. R. 1913, **157**. 981.)**Silver fluoride**, AgF.

Extremely deliquescent. (Gore.)

Sol. in 0.55 pt. H_2O at 15.5° with evolution of heat. Sp. gr. of sat. solution at $15.5^\circ = 2.61$. (Gore.)Solubility of AgF in H_2O at t° .
G. per 100 g. H_2O .

t°	AgF	Solid phase
-14.2	60	Ice + AgF, 4H ₂ O
+18.5	165	AgF, 4H ₂ O
18.65	169.5	" + AgF, 2H ₂ O
20	172	AgF, 2H ₂ O
24	178	"
25	179.5	"
28.5	215	"
32	193	"
39.5	222	AgF, 2H ₂ O + AgF
108	205	

(Guntz, A. ch. 1914, (9) **2**. 101.)Sp. gr. AgF + Aq at 18° .

% AgF	Sp. gr.
7.20	1.07
29.60	1.38
49.20	1.82
56.40	2.00
66.20	2.62

(Guntz, A. ch. 1914, (9) **2**. 104.)

Data on solubility of AgF in HF + given by Guntz (l. c.).

Sl. sol. in liquid NH_3 . (Gore, Am. 1898, **20**. 829.)+ H_2O . Deliquescent. Sol. in (Guntz, A. ch. 1914, (9) **2**. 101.)+ $2\text{H}_2\text{O}$. Deliquescent. Sol. in (Guntz.)+ $4\text{H}_2\text{O}$. Not deliquescent. Sol. in (Guntz.)+ $\frac{1}{2}\text{H}_2\text{O}$. Unstable in the presence of crystals of $\text{AgF} + 2\text{H}_2\text{O}$. (Guntz, A. ch. (9) **2**. 101.)**Silver hydrogen fluoride**, $\text{AgF} \cdot \text{HF}$.

(Guntz.)

 $\text{AgF} \cdot 3\text{HF}$. Very unstable.Sol. in HF. (Guntz, Bull. Soc. **18**. 114.)**Silver stannic fluoride.**

See Fluostannate, silver.

Silver tungstyl fluoride.

See Fluoxtungstate, silver.

Silver, fulminating.

See Silver nitride.

Silver hydride, AgH .Not decomp. by H_2O . (Bartlett, Am. J. 1896, **19**. 52.)**Argentous hydroxide**, $\text{Ag}_2\text{O} \cdot \text{H}_2\text{O}$.Sol. in H_2O . Known only in sol. (Weltzein, A. **142**. 105.)**Silver hydroxide**, AgOH .Decomp. into Ag_2O and H_2O above 100° .
See Silver oxide.

s iodide, Ag_2I .

, C. R. 112. 861.)

midosulphamide, $\text{AgN}(\text{SO}_2\text{NH}_2)_2 + \text{I}_2\text{O}$.

p. slowly in the air. Somewhat sol. in boiling H_2O , from which unchanged on cooling. In aqueous is stable toward alkali. Decomp.

Difficultly sol. in dry pyridine; l. in pyridine + Aq. (Hantzsch, B. 1035)

$\text{V}_2\text{H}_2\text{Ag}_2 + 5\frac{1}{2}\text{H}_2\text{O}$. Nearly insol. O. (Ephraim and Michel, B. 1909,)

$\text{V}_2\text{H}_2\text{Ag}_2 + 4\text{H}_2\text{O}$. (Ephraim and

$\text{V}_2\text{H}_2\text{Ag}_2 + 1\frac{1}{2}$, 11, and 28 H_2O .

sol. in HNO_3 and $\text{NH}_4\text{OH} + \text{Aq}$. pyridine. Very sol. in pyridine con-
pyridine nitrate and can be recryst.
l. (Ephraim and Michel.)

$\text{V}_2\text{Ag}_2 + 8\text{H}_2\text{O}$. (Ephraim and

ide, AgI .

n H_2O .

ded from electrical conductivity of
 AgI is sol. in 1,074,040 pts. H_2O
and 420,260 pts. at 40° . (Holleman,
Ch. 12. 130.)

H_2O dissolves 0.1 mg. AgI at 18° .
sch and Rose, Z. phys. Ch. 12. 241.)
ty in $\text{H}_2\text{O} = 1 \times 10^{-6}\text{N}$. (Rolla.)

ty in $\text{H}_2\text{O} = 0.97 \times 10^{-6}\text{g. mola. per l.}$
Woodwin, Z. phys. Ch. 1894, 12. 645.)

ty of AgI in H_2O at 25° is 1.05×10^{-6}
lity). (Thiel, Z. anorg. 1900, 24. 57.)

aq. solution at 20.8° contains 0.0020
equiv. per l. (Kohlrausch, C. C.
1299.)

H_2O dissolves 0.0035 mg. AgI at 21° .
sch, Z. phys. Ch. 1904, 50. 356.)

H_2O dissolves 0.00253 mg. AgI at 60° .
Z. phys. Ch. 1905, 53. 644.)

ty in $\text{H}_2\text{O} = 1.23 \times 10^{-6}\text{g.-mol.}$
at 25° . (A. E. Hill, J. Am. Chem.
Soc. 30. 74.)

mg. are contained in 1 l. of sat. solu-
tion. (Kohlrausch, Z. phys. Ch. 1908,

n dil. $\text{HNO}_3 + \text{Aq}$ or $\text{H}_2\text{PO}_4 + \text{Aq}$.
by hot. conc. $\text{HNO}_3 + \text{Aq}$ or H_2SO_4 .
in conc. $\text{HI} + \text{Aq}$.

AgI dissolves in 2510 pts. $\text{NH}_4\text{OH} +$
sp. gr. (Martini, Schw. J. 56. 154);
sp. gr. of 0.89 sp. gr. (Wallace and La-
Gaz. 1859. 137).

AgI dissolves in 26,300 g. 10%
Aq (sp. gr. = 0.96) at 12° . Insol.
in $\text{OH} + \text{Aq}$. (Longi, Gazz. ch. it. 19.

ent of solubility in $\text{NH}_4\text{OH} + \text{Aq}$
0.926) is found lower than previ-
ously determined and of the order of $\frac{1}{100000}$ at
Nancy, Bull. Soc. 1908, (4) 3. 772.)

According to Field, insol. in cold conc. KCl
or $\text{NaCl} + \text{Aq}$, and only in traces on boiling,
and separates out on cooling.

100 g. NaCl in conc. $\text{NaCl} + \text{Aq}$ dissolve
0.95 mg. AgI at 15° ; 100 g. NH_4Cl in conc.
 $\text{NH}_4\text{Cl} + \text{Aq}$ dissolve 2.9 mg. AgI at 15° ; 95 g.
 $\text{NaCl} + 10$ g. KBr in conc. solution dissolve
1.2 mg. AgI at 15° ; 100 g. $\text{KBr} + 225$ g. H_2O
dissolve 430 mg. AgI at 15° ; 100 g. KBr in
conc. $\text{KBr} + \text{Aq}$ dissolve 525 mg. AgI at 15° ;
100 g. $\text{KI} + 69$ g. H_2O dissolve 89.8 g. AgI at
 15° ; 100 g. $\text{KI} + 92$ g. H_2O dissolve 54.0 g.
 AgI at 15° ; 100 g. $\text{KI} + 366$ g. H_2O dissolve
7.25 g. AgI at 15° . (Schierholz, W. A. B.
101, 2b. 4.)

Sol. in conc. $\text{KI} + \text{Aq}$, from which it is pre-
cipitated by H_2O . (Field, C. N. 3. 17.)

KI gives a ppt. with AgNO_3 in presence of
30,000 pts. H_2O . (Harting.)

Solubility in $\text{KI} + \text{Aq}$ at 15° .

% KI	% AgI	% KI	% AgI
59.16	53.13	33.3	7.33
57.16	51.13	25.0	2.75
50.0	25.0	21.74	1.576
40.0	13.0	20	0.80

(Schierholz, W. A. B. 1890, 101. 2b. 10.)

Solubility in $\text{KI} + \text{Aq}$ at 25° .

Mol. KI per l.	g. AgI per l.
1.937	46.42
1.6304	24.01
1.482	15.46
1.406	12.55
1.018	3.47
1.008	3.32
0.734	1.032
0.586	0.512
0.335	0.0853

Hellwig, Z. anorg. 1900, 25. 180.)

Solubility in $\text{KI} + \text{Aq}$.

t = 50°		
% AgI	% KI	Solid phase
2.5	24.8	AgI
16.0	33.8	"
28.0	36.7	"
39.0	38.1	"
51.8	36.2	"
53.5	36.5	"
53.5	36.6	$\text{AgI} + \text{AgI, KI}$
53.5	37.1	AgI, KI
53.4	37.6	$\text{KI} + \text{AgI, KI}$
50.4	40.2	KI
45.0	43.2	"
38.0	47.1	"
22.8	55.5	"
10.7	59.1	"

t = 30°		
% AgI	% KI	Solid phase
0.1	10.2	AgI
10.0	31.4	"
29.4	37.6	"
42.8	38.8	"
49.7	38.6	AgI + AgI, 2KI
49.6	39.5	AgI, 2KI
47.7	40.9	"
46.3	41.4	"
44.1	43.2	AgI, 2KI + KI
42.8	43.9	KI
35.8	46.9	"
16.0	55.5	"
0	60.35	"

t = 0°		
% AgI	% KI	Solid phase
0.2	9.8	AgI
1.5	20.5	"
6.5	26.1	"
26.6	34.6	"
28.1	36.4	"
38.0	41.3	AgI + AgI, KI
37.9	42.0	AgI, KI
37.6	42.7	"
37.9	44.0	AgI, KI + KI
31.3	46.6	KI
21.7	50.5	"
18.0	51.2	"
9.0	53.0	"
0	56.1	"
27.5	48.7	AgI, 2KI + KI
21.0	50.3	AgI, 2KI

(Van Dam and Donk, Chem. Weekbl. 1911, 8. 848.)

Very sol. in KI₂ + Aq. (Muth, Dissert. 1895.)
Very sol. in H₂O in presence of NaI. (Kurnakow, Ch. Z. 1900, 24. 60.)

Solubility in KI + Aq at 15°.

Composition of the sat. solution in mols. per 1000 mols. H ₂ O		Solid phase
Mols. NaI ₂	Mols. Ag ₂ I ₂	
35.63	8.14	AgI
40.54	10.94	"
61.55	25.15	"
80.55	38.19	"
94.25	47.79	"
107.52	57.52	AgI + AgI, NaI, 3½H ₂ O
117.96	51.70	AgI, NaI, 3½H ₂ O
134.40	46.82	"
135.83	46.36	AgI, NaI, 3½H ₂ O + NaI
133.81	43.03	NaI
129.02	34.85	"
122.56	22.82	"
117.11	11.93	"
111.52	...	"

(Krym, J. Russ. Phys. Chem. Soc. 1909, 41. 382.)

Traces are dissolved by alkali nitrate. Easily sol. in hot KOH + Aq, from which pptd. by H₂O or alcohol. Not decomposed by boiling KOH + Aq. (Vogel, N. Rep. 20. 129.)

100 pts. of AgNO₃ + Aq sat. at 11° dissolves 2.3 pts. AgI in the cold, and 12.3 pts. in boiling. (Schnauss.)

Solubility of AgI in AgNO₃ + Aq

Mol. AgNO ₃ in 1 l.	g. AgI in 1 l.	Solubility
0.20	0.0680	AgI
0.25	0.080	
0.30	0.090	
0.35	0.125	
0.40	0.167	
0.45	0.224	
0.50	0.299	
0.55	0.400	
0.60	0.528	
0.65	0.672	Ag ₂ I
0.70	0.850	
1.215	3.08	
1.63	6.26	Ag ₂ I
2.04	10.90	
2.54	16.1	Ag ₂ I
3.115	22.7	
3.75	33.2	
4.055	40.0	
4.69	53.2	
5.90	85.0	

(Hellwig, Z. anorg. 1900, 25.)

Solubility of AgI in 25% Aq reaches a maximum at about 60°; at this point of maximum solubility the dissolved amounts to about 5 g. AgI per g. AgNO₃. (Lowry, Roy. Soc. P. 91, A, 66.)

Sol. in hot Hg(NO₃)₂ + Aq, from which it crystallizes on cooling.

Solubility of AgI in Hg(NO₃)₂ + Aq

Mols. Hg(NO ₃) ₂ per l.	g. AgI per l.	Mols. Hg(NO ₃) ₂ per l.
0.010	0.800	0.050
0.0125	0.841	0.100
0.025	1.118	1.000

Solubility is not affected by 0.1 to 2N HNO₃.

(Morse, Z. phys. Ch. 1902, 41.)

Sol. in KCN + Aq.

Sl. sol. in Na₂S₂O₃ + Aq when saturated with H₂O, but separates again on addition of KI + Aq. (Field.)

Insol. in Na₂S₂O₃ + Aq. (Fogel, Z. anorg. 1890, 110. 711.)

Solubility in salts + Aq.

t	% Conc.	grams AgI sol. in 100 grams solvent	
bate	1	0.03	20
"	5	0.15	
"	10	0.30	
"	15	0.40	
"	20	0.60	
"	10	0.01	25
"	20	0.02	
"	10	Traces	
bite	5	8.23	20
de	5	0.02	
bocyanide	10	0.08	
"	15	0.13	
"	10	0.03	
"	10	0.02	25
"	10	0.02	
"	10	0.79	
"	1	0.008	
"	5	0.05	
"	10	0.09	

nta, M. 1894, 15. 250.)

liquid NH₃. (Franklin, Am. 10. 829; Jarry, A. ch. 1899, (7)

n liquid NH₃. (Ruff and Geisel, 662.)

etone. (Eidmann, C. C. 1899, umann, B. 1904, 37. 4329.)

ethyl acetate. (Bezold, Dis-Naumann, B. 1909, 42. 3790.)

z. (Arctowski, Z. anorg. 1894,

ol. in hot alcoholic thiourea than gBr. (Reynolds, Chem. Soc.

benzonitrile. (Naumann, B. .)

in piperidine at 100°. (Varet, 5. 336.)

sol. in 100 pts. pyridine at 10°.

sol. in 100 pts. pyridine at 121°. B. 1894, 27. 2288.)

etermined in piperidine. (Wer-1897, 15. 16.)

te.

en iodide, 3AgI, HI+7H₂O. C. R. 91. 1024.)

iodide, 2AgI, NaI.

acetone. (Marsh, Chem. Soc. 4.)

-3½H₂O. (Krym, J. Russ. Soc. 1909, 41. 382.)

al under AgI.

ummonia, AgI, NH₃.

id NH₃. (Jarry, A. ch. 1899,

2AgI, NH₃. (Rammelsberg, Pogg. 48. 170.)

Composition is AgI, NH₃. (Longi, Gazz. ch. it. 13. 86.)

Sol. in liquid NH₃. (Jarry, A. ch. 1899, (7) 17. 371.)

AgI, 2NH₃. (Terreil, C. R. 98. 1279.)

Silver nitride, Ag₃N.

Berthollet's "knallsilber." Very explosive. Insol. in H₂O. Sol. in KCN+Aq. Slowly

sol. in NH₄OH+Aq. (Raschig, A. 233. 93.) (Angeli, Chem. Soc. 1894, 66. (2) 93.)

Argentous oxide, Ag₂O.

Insol. in H₂O. Decomp. by acids into argentic oxide and silver. Insol. in NH₄OH+

Aq or HC₂H₃O₂. (v. der Pfordten, B. 20. 1458.)

Contains H, and is a hydroxide Ag₂H₂O. (v. der Pfordten, B. 21. 2288.)

The above substance is a mixture, according to Friedheim (B. 20. 2557.)

Silver oxide, Ag₂O.

Somewhat sol. in H₂O. (Bucholz.)

Sol. in 3000 pts. H₂O. (Bineau, C. R. 41. 509); sol. in 98 pts. H₂O. (Abl.)

Sol. in 15,360 pts. H₂O. (Levi, Gazz. ch. it. 1901, 31. (1) 1.)

Solubility in H₂O at 25° = 2.16 × 10⁻⁴ mols. AgOH per litre. (Noyes, J. Am. Chem. Soc. 1902, 24. 1147.)

1 liter sat. aqueous solution at 19.96° contains 2.14 × 10⁻² g.; at 24.94° contains 2.5 × 10⁻² g. Ag₂O. (Böttger, Z. phys. Ch. 1903, 46. 603.)

1 l. H₂O at 25° dissolves 1.8 × 10⁻⁴ gram—atoms of silver. Determined from its solubility in NH₃. (Abegg and Cox, Z. phys. Ch. 1903, 46. 11.)

1 l. H₂O dissolves 0.0215 g. Ag₂O at 20°. (Whitby, Z. anorg. 1910, 67. 108.)

The solubility of Ag₂O in H₂O varies with the method of preparation.

Solubility of Ag₂O (prepared by action of NaOH, freshly prepared by the solution of Na in H₂O, on a dil. solution of AgNO₃) =

2.16 × 10⁻⁴ g.-mol. in 1 l. H₂O at 25°; 2.97 × 10⁻⁴ g.-mol. at 50°.

Solubility of Ag₂O (prepared by action of aqueous barium hydroxide on AgNO₃) =

2.23 × 10⁻⁴ g.-mol. in 1 l. H₂O at 25°; 3.09 × 10⁻⁴ g.-mol. in 1 l. H₂O at 50°.

Solubility of Ag₂O (prepared by action of conc. NaOH+Aq on moist, freshly pptd. AgCl) = 2.32 × 10⁻⁴ g.-mol. in 1 l. H₂O at 25°; 3.55 × 10⁻⁴ g.-mol. at 50°.

Solubility of Ag₂O (prepared by action of conc. NaOH+Aq. on moist, freshly pptd. Ag₂CO₃) = 2.95 × 10⁻⁴ g.-mol. in 1 l. H₂O at 25°; 3.89 × 10⁻⁴ g.-mol. at 50°. (Rebière, Bull. Soc. 1915, (4) 7. 311.)

Sol. in acids, NH₄OH, and (NH₄)₂CO₃+Aq. Decomp. by alkali chlorides, bromides,

and iodides + Aq. Sol. in alkali cyanides, and thiosulphates + Aq. Sl. sol. in nitrates + Aq; insol. in sulphates + Aq. When freshly pptd., sol. in NH_4SCN + Aq. Sl. sol. in NH_4NO_3 + Aq. Abundantly sol. in $\text{Ba}(\text{NO}_3)_2$ + Aq without pptn. of BaO_2H_2 . Sol. in boiling $\text{Mn}(\text{NO}_3)_2$, $\text{Ni}(\text{NO}_3)_2$, $\text{Co}(\text{NO}_3)_2$, $\text{Cu}(\text{NO}_3)_2$, and $\text{Ce}_2(\text{NO}_3)_6$ + Aq with pptn. of oxides. (Persoz.)

Insol. in KOH , and NaOH + Aq. Sl. sol. in BaO_2H_2 + Aq. (Berzelius (?)).

Solubility in NH_4OH + Aq at 25° .

G. at. Ag per l.	* Mol. NH_3 per l.
0.0654	0.214
0.0658	0.220
0.134	0.458
0.140	0.469
0.205	0.671
0.205	0.684
0.225	0.720
0.224	0.733
0.251	0.811
0.248	0.827
0.242	0.830
0.257	0.876
0.278	0.899
0.276	0.915
0.299	0.999
0.343	1.147
0.454	1.498
0.470	1.522

(Whitney and Melcher, J. Am. Chem. Soc. 1903, 25, 78.)

Insol. in liquid NH_3 . (Franklin, Am. ch. J. 1898, 20, 829.)

Insol. in acetone. (Eidmann, C. C. 1899, II, 1014); (Naumann, B. 1904, 37, 4329.)

Insol. in ethyl acetate. (Hamers, Dissert. 1906; Naumann, B. 1910, 43, 314.)

Sl. sol. in amylamine + Aq, easily in methylamine + Aq (Wurtz, A. ch. 30, 453); also in ethylamine, and thioethylamine + Aq.

Solubility in methylamine + Aq. at 18° .
G. mols. per l.

CH_3NH_2	Ag
0.1	0.0221
0.5	0.118
1.0	0.228

(Euler, B. 1903, 36, 2879.)

Solubility in ethylamine + Aq at 18° .
G. mols. per l.

$\text{C}_2\text{H}_5\text{NH}_2$	Ag
0.1	0.0322
0.5 (interpolated)	0.160
1.0 "	0.314
0.581	0.180
0.927	0.291

(Euler.)

Silver peroxide, Ag_2O_2 .

Sol. in conc. H_2SO_4 (Rose), and HNO_3 + Aq without decomp. Sol. in + Aq. (Schönbein, J. pr. 41, 321.)

Sol. in HNO_3 and H_2SO_4 with (Mulder, R. t. c. 1898, 17, 151.)

Insol. in liquid NH_3 . (Gore, Am 1898, 20, 829.)

Silver oxide ammonia.

See Silver nitride.

Silver oxybromide, Ag_2OBr .

Insol. in H_2O . Insol. in HNO_3 , hot ammonia and in NaOCl + Aq. (C. R. 1912, 164, 357.)

Silver oxyfluoride, AgF , AgOH .

Decomp. by H_2O with separation (Pfaundler.)

Silver peroxyfluoride, $2\text{Ag}_2\text{O}_4$, AgF

(Tanatar, Z. anorg. 1901, 28, 335. $4\text{Ag}_2\text{O}_4$, 3AgF . (Tanatar, Z. an. 28, 335.)

Silver oxyiodide, Ag_2O , Ag_2I_2

(Seyewitz, Bull. Soc. 1894, (3) 11)

Silver phosphide, Ag_3P .

Sol. in HNO_3 . Attacked by aq (Granger, C. R. 1897, 124, 897.)

Ag_3P . Insol. in HCl + Aq; easily in HNO_3 + Aq. (Schrötter, J. B. 1899, 12, 1014.)

Ag_3P . (Hackspill, C. R. 1913, 1, 1014.) Ag_3P (?). (Fresenius and Neel anal. 1, 340.)

Silver phosphoselenide, Ag_3Se , P_2Se_3

Insol. in H_2O or HCl + Aq. Sol. in Ao. Insol. in cold, decomp. by hot. Aq. (Hahn, J. pr. 93, 436.)

$2\text{Ag}_3\text{Se}$, P_2Se_3 . Insol. in H_2O , HNO_3 + Aq; slowly sol. in red fums (Hahn, J. pr. 93, 440.)

$2\text{Ag}_3\text{Se}$, P_2Se_3 . Sol. only in fums (Hahn.)

Silver phosphosulphide, $2\text{Ag}_3\text{S}$, P_2S_5

Ag_3S , P_2S_5 . (Berzelius, A. 46, 261. $2\text{Ag}_3\text{S}$, P_2S_5 . Easily sol. in H without separation of P. (Berzelius.)

$\text{Ag}_3\text{P}_2\text{S}_5$. (Berzelius.)

Ag_3PS_2 . Easily attacked by HCl . Sl. decomp. Insol. in hot H_2O comp. by aqua regia. (Ferrand, A. (7) 17, 413.)

Silver selenide, Ag_2Se .

Sol. in boiling HNO_3 + Aq as which separates out by dilution (Berzelius.)

Insol. in $\text{Hg}_2(\text{NO}_3)_2$ + Aq. (Was A. 41, 321.)

fin. *Naumannite*. Insol. in dil., but sol. in conc. $\text{HNO}_3 + \text{Aq}$.

er sulphamide (silver thionyl amide), $\text{SO}_2(\text{NHAg})_2$.

sol. in pyridine. (Hantzsch and Holl, B. 1, 34. 3436.)

H_2O . (Ephraim and Gurevitch, B. 43. 146.)

entous sulphide, Ag_2S .

easily sol. in warm dil. $\text{HNO}_3 + \text{Aq}$, and in conc. H_2SO_4 without separation of S. Sol. in conc. $\text{KCN} + \text{Aq}$. (v. der Pfordten, B. 20. 3; Guntz, C. R. 112. 861.)

er sulphide, Ag_2S .

less sol. in H_2O than AgI . (Lucas, Z. anorg. Chem. 1904, 41. 210.)

1. H_2O dissolves about 4×10^{-11} g. at. Ag_2S at 18° . (Bernfeld, Z. phys. Ch. 1898, 72.)

1. H_2O dissolves 0.8×10^{-6} g. mols. at 18° . (Biltz, Z. phys. Ch. 1907, 58. 291.)

1. H_2O dissolves 0.552×10^{-6} g. mols. at 18° . (Weigel, Z. phys. Ch. 1907, 58.)

1. in conc. $\text{HNO}_3 + \text{Aq}$ with separation of S. Sol. in hot conc. $\text{HCl} + \text{Aq}$. Not decomp. by $\text{CuCl}_2 + \text{Aq}$, but by $\text{CuCl}_2 + \text{NaCl}$.

1. Insol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in $\text{H}_2\text{S} + \text{Aq}$, or in $\text{Hg}(\text{NO}_3)_2 + \text{Aq}$.

1. sol. in H_2O , dil. acids, alkalis, and alkali hydrides + Aq . (Fresenius.)

1. in $\text{HCN} + \text{Aq}$. (Hahn, C. C. 1870.)

1. **ptd.** Ag_2S is very sol. in HNO_3 containing more than 5% HNO_3 . (Gruener, J. Am. Chem. Soc. 1910, 32. 1032.)

1. **ly** very sl. sol. in $\text{AgNO}_3 + \text{Aq}$, even at 10° . (Lowry, Roy. Soc. Proc. 1914, 91, A.)

1. in $\text{KCN} + \text{Aq}$. (Hahn, C. C. 1870.)

1. **fficultly** sol. in $\text{KCN} + \text{Aq}$; less difficultly sol. Ag_2S is pptd. from a very dil. solution.

1. of KCN present also has influence on solubility. Ag_2S dissolved in conc. KCN separates out on dilution. (Béchamp, C. R. 60. 64.)

1. sol. in NH_4Cl or $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Brett.)

1. in *Argentite*. *Acanthite*. Sol. in conc. $\text{H}_2\text{S} + \text{Aq}$ with separation of S.

1. in citric acid + Aq with addition of H_2O . (Bolton, C. N. 37. 48.)

er disulphide, Ag_2S_2 .

1. in H_2O with decomp.; also sol. with decomp. in HCl , HNO_3 . CS_2 does not dissolve it. (Hantzsch, Z. anorg. 1898, 19. 105.)

er sodium sulphide, $3\text{Ag}_2\text{S}, \text{Na}_2\text{S} + 2\text{H}_2\text{O}$.

1. in conc. $\text{Na}_2\text{S} + \text{Aq}$ with decomp.; sol. in H_2O with decomp. (Ditte, C. R. 1895, 93.)

Silver zinc sulphide, $\text{Ag}_2\text{S}, 3\text{ZnS}$.

(Schneider, J. pr. (2) 8. 29.)

Silver sulphimide (silver thionyl imide), $\text{SO}_2\text{N}^+\text{Ag}$.

Very sl. sol. in cold, more sol. in hot H_2O . Very sol. in dil. HNO_3 . (Traube, B. 1892, 25. 2474.)

Silver sulphophosphide.

See Silver phosphosulphide.

Silver telluride, Ag_2Te .

Min. *Hessite*. Sol. in warm $\text{HNO}_3 + \text{Aq}$.

Sodammonium, $\text{Na}_2(\text{NH}_3)_2$.

100 g. liq. NH_3 dissolve 60.5 g. at -23° ; 56.4 g. at 0° ; 56 g. at $+5^\circ$; 55 g. at 9° . (Joannis, A. ch. 1906, (8) 7. 41.)

Sodium, Na_2 .

Violently decomposes H_2O , alcohol, etc. Insol. in hydrocarbons. Easily sol. in acids with violent action.

Solubility in fused NaOH .

G. sol. in 100 g. fused NaOH at temp.

t°	G. per 100 g. NaOH
480	25.3
600	10.1
610	9.9
670	9.5
760	7.9
800	6.9

(Hevesy, Z. Elektrochem. 1909, 15. 531.)

Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54. 674.)

Sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

1 gram atom dissolves:—

at $+22^\circ$ in 6.14 mol. liquid NH_3 .

“ 0° “ 5.87 “ “ “

“ -30° “ 5.52 “ “ “

“ -50° “ 5.39 “ “ “

“ -70° “ 5.20 “ “ “

“ -105° “ 4.98 “ “ “

(Ruff, B. 1906, 39. 839.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0449 g. Na in 6 days. (Gates, J. phys. Chem. 1911, 15. 143.)

Insol. in ethylamine and in secondary and tertiary amines. (Kraus, J. Am. Chem. Soc. 1907, 29. 1561.)

Sodium acetylde acetylene, $\text{Na}_2\text{C}_2, \text{C}_2\text{H}_2$.

Very deliquescent. Decomp. by H_2O and by absolute alcohol. Insol. in ether, ligroin, etc. (Moissan, C. R. 1898, 127. 915.)

Sodium amalgam.

NaHg₈. Stable in contact with the liquid amalgam from 0°–40.5°. Can be cryst. from Hg without decomp. at any temp. between these limits.

NaHg₁. Stable in contact with the liquid amalgam from 40.5°–150°. Can be cryst. from Hg without decomp. at any temp. between these limits. (Kerp, Z. anorg. 1900, 25. 68.)

Sodium amide, NaNH₂.

Decomp. by H₂O and alcohol.

Sodium amidochloride, Na₂NH₂Cl.

Sol. in H₂O with decomp. (Joannis, C. R. 112. 392.)

Sodium arsenide, Na₃As.

Decomp. H₂O. (Lebeau, C. R. 1900, 130. 504.)

Sodium arsenide ammonia, Na₃As, NH₃.

Easily sol. in liquid NH₃. (Lebeau, C. R. 1900, 130. 502.)

Sl. sol. in liquid NH₃. (Hugot, C. R. 1898, 127. 554.)

Sodium azoimide, NaN₃.

Not hygroscopic. Sol. in H₂O. Insol. in alcohol and ether. (Curtius, B. 24. 3344.)

40.16 pts. are sol. in 100 pts. H₂O at 10°.
40.7 " " " " 100 " H₂O " 15.2.
41.7 " " " " 100 " H₂O " 17.0°.
0.3153 pt. is sol. in 100 pts. abs. alcohol at 16°.

Insol. in pure ether. (Curtius, J. pr. 1898, (2) 58. 279.)

Sodium bromide, NaBr, and +2H₂O.

Not deliquescent. Solubility in H₂O differs according as NaBr or NaBr+2H₂O is used. The following data for anhydrous NaBr were found.

Pts. NaBr dissolved by 100 pts. H₂O at t°.

t°	Pts. NaBr	t°	Pts. NaBr	t°	Pts. NaBr
44.1	115.6	74.5	118.4	97.2	119.9
51.5	116.2	80.5	118.6	100.3	120.6
55.1	116.8	86.0	118.8	110.6	122.7
60.3	117.0	90.5	119.7	114.3	124.0
64.5	117.3

Solubility is represented by a straight line of the formula $S = 110.34 + 0.1075t$.

Below 50° the salt usually crystallizes with

2H₂O, of which the solubility in 100 pts was found to be as follows:

t°	Pts. NaBr	t°	Pts. NaBr	t°	Pts. NaBr
–21	71.1	+5	82.0	30	86.1
–20	71.4	10	84.5	35	87.1
–15	73.1	15	87.3	40	88.1
–10	75.1	20	90.3	45	89.1
–5	77.1	25	93.8	50	90.1
0	79.5

(Coppet, A. ch. (5) 30. 420.)

If solubility $S = \text{pts. NaBr in 100 pts. H}_2\text{O}$, $S = 40.0 + 0.1746t$ from -20° to 50° , $S = 52.3 + 0.0125t$ from 50° to 150° . (C. R. 98. 1432.)

100 pts. H₂O dissolve: at 0°, 77.5 pts.; at 20°, 88.4 pts.; at 40°, 104.2 pts.; at 60°, 111.1 pts.; at 80°, 112.4 pts.; at 100°, 113.1 pts. (Kremers.)

Sat. solution boils at 121°. (Kremers, 97. 14.)

Sat. NaBr + Aq contains at:

–22°	–10°	+140°	163°
40.1	42.5	56.5	57.5
180°	180°	210°	212°
59.5	59.0	60.9	61.0
			62.0

(Étard, A. ch. 1894, (7) 2. 53)

100 g. sat. NaBr + Aq at 16.4° contains 100 g. NaBr. (Greenish, Pharm. J. 1900, 1901, 1902, 1903, 1904, 1905, 1906, 1907, 1908, 1909, 1910, 1911.) Solubility of NaBr + 2H₂O in H₂O 65.5% anhydrous NaBr. (Cocheret, 1911.)

Sp. gr. of NaBr + Aq at 19.5° contains

5	10	15	20	25
1.040	1.080	1.125	1.174	1.226

30	35	40	45	50
1.281	1.334	1.410	1.483	1.565

(Gerlach, Z. anal. 8. 285.)

NaBr + Aq containing 17.15% NaBr sp. gr. 20°/20° = 1.1473.

NaBr + Aq containing 22.72% NaBr sp. gr. 20°/20° = 1.2060.

(Le Blanc and Rohland, Z. phys. 19. 278.)

Sp. gr. of NaBr + Aq at 20.5°

Normality of NaBr + Aq	g. NaBr in 100 g. of solution	Sp. gr.
4.33	33.57	1.1473
3.00	25.10	1.1473
1.99	17.77	1.1473
0.98	9.41	1.1473

(Oppenheimer, Z. phys. Ch. 1898, 2

gr. at 20° of NaBr+Aq containing
mols. NaBr per liter.

0.01	0.025	0.05	0.075
r. 1.000732	1.002177	1.004074	1.005972

0.10	0.25	0.50	0.75
r. 1.00788	1.01964	1.03908	1.05811

1.0	1.5	2.0
r. 1.07632	1.11963	1.15240

es and Pearce, Am. Ch. J. 1907, 38. 728.)

l. in H₂SO₄. (Walden, Z. anorg. 1902, 84.)

0 pts. NaBr+Aq sat. at 18-19° contain
i pts. NaBr; 100 pts. NaBr+NaCl+Aq
at 18-19° contain 46.59 pts. of the two
; 100 pts. NaBr+NaI+Aq sat. at 18-19°
in 63.15 pts. of the two salts; 100 pts.
r+NaCl+NaI+Aq sat. at 18-19° con-
63.20 pts. of the three salts. (v. Hauer,
98. 137.)

ubility of NaBr in NaOH+Aq at 17°.
(G. per 100 g. H₂O.)

OH	NaBr	NaOH	NaBr
0	91.38	22.35	59.60
.26	79.86	24.74	55.03
.24	68.85	28.43	48.00
.43	64.90	36.61	38.41
.17	63.06	46.96	29.37
.12	62.51	54.52	24.76

(Ditte, C. R. 1897, 124. 30.)

aily sol. in liquid HF. (Franklin, Z.
. 1905, 46. 2.)

y sl. sol. in alcohol.
Br+2H₂O is sol. in 1.10 pts. H₂O at
n 159 pts. absolute alcohol at 15°; in
pts. absolute ether at 15°. (Eder,
. 221. 89.)

Br+2H₂O is sol. in 2.25 pts. 60% alco-
nd 7 pts. 90% alcohol. NaBr is sol. in
60% alcohol, and 10 pts. 90% alcohol.
er.)

1 pts. absolute methyl alcohol dissolve
pts. at 19.5°. (de Bruyn, Z. phys. Ch.
13.)

1 g. NaBr+CH₃OH contain 0.9 g. NaBr
ie critical temp. (Centnerszwer, Z.
Ch. 1910, 72. 437.)

room temp., 1 pt. NaBr by weight is
1:

4.6	pts. methyl alcohol	D ₁₅ 0.7990.
4.0	“ ethyl	“ D ₁₅ 0.8100.
9.7	“ propyl	“ D ₁₅ 0.8160.

Rohland, Z. anorg. 1898, 18. 325.)

Solubility in ethyl alcohol at 30°.

Wt. %		Solid phase
Alcohol	NaBr	
0	59.4	NaBr, 2H ₂ O
11.79	42.90	“
31.78	32.12	“
43.22	26.79	“
54.59	20.83	“
65.51	16.08	“
72.36	13.41	“
76.92	12.03	NaBr, 2H ₂ O+NaBr
87.35	7.44	NaBr
97.08	3.01	“

(Cocheret, Dissert. 1911.)

Solubility in mixtures of methyl and ethyl
alcohol at 25°.

P = % methyl alcohol in the solvent.
G = g. NaBr in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0.00	0.293	0.8189
4.37	0.365	0.8265
10.40	0.404	0.8273
41.02	0.724	0.8593
80.69	1.251	0.9079
84.77	1.286	0.9104
91.25	1.432	0.9235
100.00	1.440	0.9238

(Herz and Kuhn, Z. anorg. 1908, 60. 155.)

Solubility in mixtures of methyl and propyl
alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. NaBr in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	1.440	0.9238
11.11	1.243	0.9048
23.8	1.053	0.8887
65.2	0.442	0.8390
91.8	0.147	0.8153
93.75	0.126	0.8144
100	0.074	0.8093

(Herz and Kuhn, Z. anorg. 1908, 60. 156.)

Solubility in mixtures of propyl and ethyl alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. NaBr in 10 ccm. of the solution.
S = Sp. gr. of the sat solution.

P	G	S 25°/4°
0	0.293	0.8189
8.1	0.249	0.8147
17.85	0.247	0.8145
56.6	0.190	0.8107
88.6	0.111	0.8116
91.2	0.083	0.8083
95.2	0.082	0.8090
100	0.074	0.8093

(Herz and Kuhn, Z. anorg. 1908, 60. 159.)

2.05 g. are sol. in 100 g. propyl alcohol. (Schlamp, Z. phys. Ch. 1894, 14. 276.)

Sl. sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

100 g. 95% formic acid dissolve 22.3 g. NaBr at 18.5°. (Aschan, Ch. Ztg. 1913, 37. 1117.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314); benzonitrile. (Naumann, B. 1914, 47. 1370.)

The composition of the hydrates formed by NaBr at different dilutions is calculated from determinations of the lowering of the fr-pt. produced by NaBr and of the conductivity and sp. gr. of NaBr + Aq. (Jones, Am. Ch. J. 1905, 34. 303.)

Sodium stannic bromide.

See Bromostannate, sodium.

Sodium uranous bromide, Na₂UBr₆.

As K salt. (Aloy, Bull. Soc. 1899, (3) 21. 264.)

Sodium zinc bromide, NaBr, ZnBr₂ + H₂O.

Hygroscopic. (Ephraim, Z. anorg. 1908, 59. 63.)

2NaBr, ZnBr₂ + 5H₂O. Hygroscopic. (Ephraim.)

Sodium carbide, Na₂C₂.

Insol. in all neutral solvents; decomp. on heating and by H₂O. (Matignon, C. R. 1897, 125. 1034.)

Sodium carbonyl, Na₂C₂O₃.

Decomp. by H₂O with explosion. (Joannis, C. R. 116. 1518.)

Sodium subchloride, Na₂Cl₂.

Decomp. by H₂O into NaCl and NaOH + Aq. (Kreutz, B. 1897, 30. 403.)

Sodium chloride, NaCl.

Sol. in H₂O.

100 pts. H₂O at t° dissolve pts. NaCl

t°	Pts. NaCl	Authority
0	More than at 13.89°	Gay-Lussac, A. ch. 310.
13.89	35.81	
16.90	35.88	
59.93	37.14	
109.73	40.38	
12	35.91	Fehling, A. 77. 30
100	39.92	
18.75	37.731	Bischof.
10-15	35.42	Bergmann.
106+	42.86	Griffiths, 1883.
20	35.9	Schiff, A. 100. 35
All temps.	37	Fuchs and Reeb 1886.
25	35.7	Kopp, A. 84. 263.
18.75	36.53	C. J. B. Karsten.
1	36.121	G. Karsten.
18.75	36.724	
100	41.076	
1.25	36.119	Unger, J. pr. 8. 26
Boiling	39.324	
18.75	35.40	Karsten (?), cit Unger, l.c.
100	36.95	
15.56	34.2-35.42	Ure's Dict.
100	36.16	
15	35.837	Michel and Kraft

1 pt. NaCl is sol. in 2.789 pts. H₂O at 15° (G in 3 pts. H₂O at 18.75° (Abl); in 2.8235 pts. H₂O temp. (Bergmann); in 2.7647 pts. boiling H₂O mann); in 2.857 pts. hot or cold H₂O (Fourcroy

Not deposited from boiling aqueous solution the vessel containing it is open to the air. (Tag

Solubility in 100 pts. H₂O at t°.

t°	Pts. NaCl	t°	Pts.
1 5	33.6	70	
13.75	35.8	108.5	

(Nordenskjöld, Pogg. 196. 315.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. NaCl	t°	Pts.
13.89	35.8	59.93	
16.90	35.9	109.73	

(Gay-Lussac, A. ch. 11. 296.)

Solubility of NaCl at various pressures. The represent pts. NaCl in 100 pts. sat. NaCl t° and A pressure in atmospheres.

A	0°	9°	12°	15°	20°	25°
1	26.25	26.32	26.35	26.30	26.35	26.37
20	26.35	26.38	..	26.39	26.37	26.47
40	26.44	26.40

(Müller, Pogg. 117. 386.)

100 pts. H ₂ O dissolve at t°.			
t°	Pts. NaCl	t°	Pts. NaCl
-15	32.73	40	36.64
-10	33.49	50	36.98
-5	34.22	60	37.25
0	35.52	70	37.88
5	35.63	80	38.22
9	35.74	90	38.87
14	35.87	100	39.61
25	36.13	109.7	40.35

(Poggiale, A. ch. (3) 8. 649.)

100 pts. H₂O dissolve at:
0° 9° 12° 15°
.59 35.72 35.77 35.68 pts. NaCl,

20° 25° 30°
35.77 35.81 36.00 pts. NaCl.
(Müller, Pogg. 122. 337.)

100 pts. H₂O dissolve 35.76–36.26 pts. NaCl at 15.6°, and the sp. gr. of sat. solution 1.204. (Page and Keightley, Chem. Soc. 10. 566.)
100 pts. NaCl+Aq sat. at 18–19° contain 47 pts. NaCl. (v. Hauer, J. pr. 98. 137.)

Solubility of NaCl in 100 pts. H₂O at t°.

t°	Pts. NaCl	t°	Pts. NaCl	t°	Pts. NaCl
0	35.7	30	36.3	60	37.3
1	35.7	31	36.3	61	37.3
2	35.7	32	36.3	62	37.4
3	35.7	33	36.4	63	37.4
4	35.7	34	36.4	64	37.5
5	35.7	35	36.4	65	37.5
6	35.7	36	36.5	66	37.6
7	35.7	37	36.5	67	37.7
8	35.7	38	36.5	68	37.7
9	35.7	39	36.6	69	37.8
0	35.8	40	36.6	70	37.9
1	35.8	41	36.6	71	37.9
2	35.8	42	36.7	72	38.0
3	35.8	43	36.7	73	38.0
4	35.8	44	36.8	74	38.1
5	35.9	45	36.8	75	38.2
6	35.9	46	36.8	76	38.2
7	35.9	47	36.9	77	38.2
8	35.9	48	36.9	78	38.2
9	36.0	49	36.9	79	38.3
0	36.0	50	37.0	80	38.4
1	36.0	51	37.0	81	38.4
2	36.0	52	37.0	82	38.5
3	36.1	53	37.1	83	38.6
4	36.1	54	37.1	84	38.6
5	36.1	55	37.1	85	38.7
6	36.1	56	37.2	86	38.7
7	36.2	57	37.2	87	38.8
8	36.2	58	37.2	88	38.9
9	36.2	59	37.3	89	39.0

Solubility of NaCl in 100 pts. H₂O at t°.—
Continued.

t°	Pts. NaCl	t°	Pts. NaCl	t°	Pts. NaCl
90	39.1	97	39.5	104	40.0
91	39.1	98	39.6	105	40.1
92	39.2	99	39.7	106	40.1
93	39.3	100	39.8	107	40.2
94	39.3	101	39.8	108	40.3
95	39.4	102	39.9	109	40.3
96	39.4	103	40.0	109.7	40.4

(Calculated by Mulder from his own and other observations, Scheik. Verhandel. 1864. 37.)

Solubility in 100 pts. H₂O at:
0–4° 20° 40° 60° 80°
35.630 35.825 36.32 37.06 38.00
(Andreae, J. pr. (2) 29. 456.)

Solubility in 100 pts. H₂O from most careful experiments.
0° 20° 60° 80°
35.571 35.853 37.091 38.046
(Raupenstrauch, M. Ch. 6. 563.)

Solubility of NaCl in 100 pts. H₂O at t°.

t°	Pts. NaCl	t°	Pts. NaCl
-14.0	32.5	44.75	36.64
-13.8	32.15	52.5	37.04
-6.25	34.22	55.0	36.99
-5.95	34.15	59.75	37.31
0	35.7	71.3	37.96
3.6	35.79	74.45	37.96
5.3	35.8	82.05	38.41
14.45	35.94	86.7	38.47
20.85	35.63	93.65	38.90
25.45	35.90	101.7	40.76
38.55	36.52

Solubility above 20° is represented by the formula $S=34.359+0.0527t$. (Coppet, A. ch. (5) 30. 426.)

Solubility of NaCl in 100 pts. H₂O at high temp.

t°	Pts. NaCl	t°	Pts. NaCl
118	39.8	160	43.6
140	42.1	180	44.9

(Tilden and Shenstone, Phil. Trans. 1884. 23.)

Sat. NaCl+Aq contains % NaCl at t°.			
t°	% NaCl	t°	% NaCl
-21	23.7	77	28.0
-21	23.4	90	28.2
-18	23.5	115	29.1
-17	23.3	135	28.9
- 7	25.5	140	28.8
0	25.8	150	29.6
+15	26.7	180	30.2
55	26.8	215	31.6

(Étard, A. ch. 1894, (7) 2. 532.)

100 g. H₂O dissolve 0.616 gram-equivalent NaCl at 25°. (Van't Hoff and Meyerhoffer, Z. phys. Ch. 1904, 49. 315.)

Solubility of NaCl in H₂O at t°.

Most careful experiments.

t°	g. NaCl per 100 g. H ₂ O	Sp. gr.	t°	g. NaCl per 100 g. H ₂ O	Sp. gr.
0.35	35.75	1.2090	61.70	37.28	1.1823
15.20	35.84	1.2020	75.65	37.82	1.1764
30.05	36.20	1.1956	90.50	38.53	1.1701
45.40	36.60	1.1891	107	39.65	1.1631

(Berkeley, Phil. Trans. Roy. Soc. 1904, 203. A. 189.)

Sat. NaCl+Aq. at 25° contains 26.5% NaCl. (Foote, Am. Ch. J. 1906, 35. 239.)

100 g. H₂O dissolve 35.80 g. NaCl at 25°. (Cameron, Bell and Robinson, J. phys. Ch. 1907, 11. 396.)

100 g. NaCl+Aq. sat. at 15° contains 26.3 g. NaCl; at 30°, 26.47 g. (Schreinemakers, Arch. néer. Sc. 1910, (2) 15. 81.)

5.456 g. mol. are contained in 1 l. NaCl+Aq sat. at 25°. (Herz, Z. anorg. 1911, 73. 274.)

5.40 g. mol. are contained in 1 l. NaCl+Aq sat. at 30°. (Masson, Chem. Soc. 1911, 99. 1136.)

26.47 g. NaCl are contained in 100 g. NaCl+Aq. sat. at 30°. (Cocheret, Dissert. 1911.)

35.79 g. NaCl are sol. in 100 g. H₂O at room temp. (Frankforter, J. Am. Chem. Soc. 1914, 36. 1106.)

100 mol. H₂O dissolve at:
19.3° 29.7° 40.1° 54.5°
11.04 11.06 11.15 11.35 mol. NaCl.
(Sudhaus, Miner. Jahrb. Beil. Bd. 1914, 37. 18.)

Solubility of NaCl in H₂O at 24.5° at varying pressures.

S = g. NaCl in 100 g. solvent.
P = pressure in atmospheres.

P	S	100 g. of solution cont. g. NaCl
1	35.90	26.42
250	36.25	26.61
500	36.55	26.77
1000	37.02	27.02
1500	37.36	27.20

(Cohen, Inouye and Euwen, Z. phys. 1910, 75. 257.)

Sp. gr. of NaCl+Aq containing 15% NaCl = 1.15° (Francœur); 1.116 at 15° (Soubeiran); 1.110° (Coulier); 1.111 at 15° (Baudin, C. R. 63. 937).
Sp. gr. of NaCl+Aq saturated at 15° is 1.2046 (Michel and Krafft); at 17.5° is 1.2046 (Karnes); is 1.205 (Anthon).

Sp. gr. of NaCl+Aq.

% NaCl	Sp. gr.	% NaCl	Sp. gr.	% NaCl
5	1.037	15	1.112	25
10	1.074	20	1.154	26.43

(Dahlmann, J. B. 7. 321.)

Sp. gr. of NaCl+Aq at 20°.

% NaCl	Sp. gr.	% NaCl	Sp. gr.
1	1.0066	15	1.10
2	1.0133	16	1.11
3	1.0201	17	1.12
4	1.0270	18	1.13
5	1.0340	19	1.14
6	1.0411	20	1.14
7	1.0483	21	1.15
8	1.0556	22	1.16
9	1.0630	23	1.17
10	1.0705	24	1.18
11	1.0781	25	1.19
12	1.0857	26	1.19
13	1.0934	27	1.20
14	1.1012

(Schiff, A. 110. 76.)

Sp. gr. of NaCl+Aq at 19.5°.

% NaCl	Sp. gr.	% NaCl	Sp. gr.
6.402	1.0460	22.631	1.11
12.265	1.0895	26.530	1.14
17.533	1.1303

(Kremers, Pogg. 95. 120.)

Sp. gr. of NaCl+Aq at 15°.

% NaCl	Sp. gr.	% NaCl	Sp.gr.
1	1.00725	15	1.11146
2	1.01450	16	1.11938
3	1.02174	17	1.12730
4	1.02899	18	1.13523
5	1.03624	19	1.14315
6	1.04366	20	1.15107
7	1.05108	21	1.15931
8	1.05851	22	1.16755
9	1.06593	23	1.17580
10	1.07335	24	1.18404
11	1.08097	25	1.19228
12	1.08859	26	1.20098
13	1.09622	26.395	1.20433
14	1.10384

(Gerlach, Z. anal. 8. 279.)

Sp. gr. of NaCl+Aq at 18°.

% NaCl	Sp. gr.	% NaCl	Sp. gr.
5	1.0345	25	1.1898
10	1.0707	26	1.1982
15	1.1087	26.4	1.2014
20	1.1477

(Kohlrausch, W. Ann. 1879. 1.)

gr. of NaCl+Aq at 20°, containing n mols. H₂O to 1 mol. NaCl.

n	Sp. gr.	n	Sp. gr.
2.5	1.15292	100	1.02069
25	1.08207	200	1.00965
100	1.04227

(Marignac, J. B. 1870. 110.)

gr. of NaCl+Aq at 0°. NaCl=g. NaCl to 100 g. H₂O; d°=sp. gr. at 0°; d^T=maximum sp. gr.; T=temp. of maximum.

NaCl	d°	d ^T	T
0	1.00000	1.000130	+ 4°
0.5	1.003925	1.003988	+ 3
1	1.007634	1.007666	+ 1.77
2	1.015366	1.015367	— 0.58
3	1.023530	1.023583	— 3.24
4	1.030669	1.030890	— 5.63
5	1.045975	1.046952	—11.07

(Rosetti, A. ch. (4) 17. 382.)

Sp. gr. of NaCl+Aq at 20°. x=mols. NaCl to 100 mols. H₂O.

x	Sp gr.	x	Sp. gr.
0.5	1.01145	4.0	1.08408
1.0	1.02255	5.0	1.10276
2.0	1.04393

(Nicol, Phil. Mag. (5) 16. 122.)

Sp. gr. of NaCl+Aq at 0°. S=weight of salt in 100 g. of solution of the given sp. gr.; S₁=No. mols. of salt contained in 100 mols. of the solution.

S	S ₁	Sp. gr.
23.0821	8.627	1.1821
19.1932	6.769	1.1502
14.3415	4.898	1.1111
9.4120	3.097	1.0722
5.1536	1.644	1.0394

(Charpy, A. ch. (6) 29. 23.)

Sp. gr. of NaCl+Aq.

G.-equivalents NaCl per liter	t°	Sp. gr. t°/t°
0.005028	18.549	1.0002119
0.01005	18.550	1.0004258
0.02005	18.538	1.000848
0.04983	18.509	1.002101
0.09873	18.525	1.004143
0.19388	18.542	1.008093
0.28999	18.559	1.012053
0.47574	18.558	1.019627
0.49860	18.06	1.02054
4.9860	17.85	1.18783
0.00259	14.07	1.0001108
0.005178	14.076	1.0002210
0.010318	14.097	1.0004401
0.12580	14.097	1.005315
0.25019	14.076	1.010505

(Kohlrausch, W. Ann. 1894, 53. 26.)

Sp. gr. of NaCl+Aq at 18°/18°.

g.-equivalents of NaCl in 1 liter of solution	Sp. gr.
0.005	1.0002104
0.010	1.0004206
0.020	1.0008476
0.050	1.002109
0.100	1.004205

(Tammann, Z. phys. Ch., 1895, 18. 93.)

Sp. gr. of NaCl+Aq sat. 18.0°, when p = per-
cent strength of solution; d = observed
density; and w = volume conc. in grams
per cc. ($\frac{pd}{100} = w$.)

p	d	w
25.37	1.1928	0.30263
21.25	1.1592	0.24637
17.35	1.1277	0.19503
13.25	1.0958	0.14518
9.34	1.0665	0.09960
4.810	1.0332	0.04969
2.991	1.0202	0.03052
2.593	1.0173	0.2638
1.746	1.0111	0.01765

(Barnes. J. Phys. Chem. 1898, 2. 544.)

Sp. gr. of NaCl+Aq at 20.5°.

Normality of NaCl+Aq	G. NaCl in 100 g. of solution	Sp. gr. 20.5°/4°
3.97	20.22	1.1489
2.96	15.56	1.1124
2.01	10.90	1.0775
0.97	5.49	1.0373

(Oppenheimer. Z. phys. Ch. 1898, 27. 451.)

Sp. gr. of NaCl+Aq at 15°.

Per cent NaCl	Sp. gr.
0	1.00000
5	1.03624
10	1.07335
15	1.11146
20	1.15107
25	1.19228
Sat. at 15°	1.20433

(H. C. Hahn, J. Am. Chem. Soc. 1898, 20. 622.)

Sp. gr. of NaCl+Aq at 18°/4°.

g. NaCl in 100 g. of solution	Sp. gr.
0.19560	1.0001
0.097952	0.9994
0.065410	0.99918
0.048977	0.99905

(Jahn, Z. phys. Ch. 1900, 33. 572.)

Sp. gr. 20°/4° of a normal solution of NaCl
= 1.03866. (Haigh, J. Am. Chem. Soc.
1912, 34. 1151.).

Sp. gr. of sat. NaCl+Aq at t°.

t°	g. NaCl sol. in 100 g. H ₂ O	Sp. gr.
-10°	32.90	1.200
0	35.63	1.210
10	35.69	1.205
20	35.82	1.202
30	36.03	1.198
40	36.32	1.193
50	36.67	1.189
60	37.06	1.184
70	37.51	1.178

(Tschernaj, J. Russ. Phys. Chem. Soc. 1912
44. 1565.)

Sp. gr. of dil. NaCl+Aq at 20.004°.
Conc. = g. equiv. NaCl per l. at 20.004°
Sp. gr. compared with H₂O at 20.004° = 1

Conc.	Sp. gr.
0.0000	1.000,000,0
0.0001	1.000,004,2
0.0002	1.000,008,5
0.0005	1.000,021,3
0.0010	1.000,042,7
0.0020	1.000,085,1
0.0040	1.000,169,6
0.0050	1.000,211,7
0.0100	1.000,423,3

(Lamb and Lee, J. Am. Chem. Soc. 1913, 35
1686.)

The saturated solution boils at 108°
(Kremers.)

NaCl+Aq containing 42.9 pts. NaCl to 100
pts. H₂O boils at 106.8° (Griffiths); contain-
ing 41.2 pts. NaCl to 100 pts. H₂O boils
108.2° (Legrand); containing 40.38 p
NaCl to 100 pts. H₂O boils at 109.73° (G
Lussac); containing 38.7 pts. NaCl to 100 p
H₂O forms a crust at 108.3°; highest point
served, 108.8° (Gerlach, Z. anal. 28. 426.)

Boiling-point of NaCl+Aq.

% NaCl	B.-pt. according to	
	Bischof	G. Karsten
5	101.50°	101.10°
10	103.03	102.38
15	104.63	103.83
20	106.26	105.46
25	107.93	107.27
29.4	107.9-108.99	..

% NaCl	B.-pt. according to	
	Legrand	Gerlach
5	100.80°	100.9°
10	101.75	101.9
15	103.00	103.3
20	104.60	105.3
25	106.60	107.6

+Aq containing pts. NaCl to H_2O . G=according to Gerlach 26. 438; L=according to Lech. (2) 59. 431).

L	B.-pt.	G	L
4.4	105.5°	27.5	29.8
7.7	106	29.5	31.8
10.8	106.5	31.5	33.9
13.4	107	33.5	35.8
15.9	107.5	35.5	37.7
18.3	108	37.5	39.7
20.7	108.4		41.2
23.1	108.5	39.5	..
25.5	108.8	40.7	..
27.7

Cl+Aq containing g. NaCl in

6 11 0 14 9 16 1 18.8
2° 103 104 2 104 8 106 1

3 24 0 28.0 28 7
1 107 7 108 7 109 6

d, Analyst, 1893, 18. 142.)

dissolved in 15 pts. H_2O , heat the temp. is 15°, but much less; at 100° there is neither absorption of heat. (Berthelot,)

mixed with 100 pts. H_2O at a temp. 2.5°. (Rüdorff, B. 2.

l with 100 pts. snow at -1° -21.3°. (Rüdorff, Pogg. 113.

Freezing-point of NaCl + Aq is lowered by gram NaCl up to 10 g. i.e. the freezing-point sinks proportionally, 2 H_2O , 0.342° for every salt. (Rüdorff, Pogg. 113.

e. HCl+Aq.

aCl in HCl+Aq at 0° NaCl = Cl (in milligrams) dissolved of liquid; HCl = mols HCl (in g) dissolved in 10 ccm of liquid

HCl	Sum of mols.	Sp. gr
1	54.5	1.2045
1.85	54.05	1.2025
5.1	53.6	1.196
9.275	53.275	1.185
5.05	53.00	1.1725
0.75	54.95	1.141
6.35	62.45	1.1159

Bull. Soc. (2) 45. 654.)

Solubility in HCl+Aq at 10-10.5°.

g. per 100 cc.		G. per 100 cc.	
HCl	NaCl	HCl	NaCl
0.0	35.77	26.06	30.89
9.84	33.76	94.77	20.01
12.76	33.19	102.1	19.04
15.68	32.71	120.6	16.03
20.78	31.77		..

(Enklaar, R. t. c. 1901, 20. 183.)

Solubility in HCl+Aq at 30°.

Composition of the solution

% by wt. HCl	% by wt. NaCl
0	26.47
6.93	16.16
12.50	9.35
17.35	4.52
35.60	0.11

(Schreinemakers, Z. phys. Ch. 1909, 63. 85.)

Solubility in HCl+Aq.

Conc. = concentration of HCl g. mol. per 1,000 g. H_2O .

NaCl = wt. NaCl dissolved in 1,000 g. H_2O .

t°	Conc.	NaCl	Molecular solubility
0	0	357.75	6.13
	0.25	341.70	5.85
	0.50	324.45	5.56
	1.00	291.20	4.99
25	0	360.80	6.18
	0.25	344.50	5.90
	0.50	329.05	5.64
	1.00	298.10	5.10

(Armstrong and Eyre, Proc. R. Soc. 1910, (A) 84. 127.)

Solubility in HCl+Aq at 30°.

G. mols. per l.

HCl	NaCl	Sp. gr. 30°	HCl	NaCl	Sp. gr. 30°
0.0	5.400	1.2018	3.052	2.463	1.1427
0.4575	4.932	1.1906	4.152	1.628	1.1289
0.969	4.386	1.1801	5.950	0.630	1.1188
1.786	3.589	1.1633	7.205	0.268	1.1258
2.412	2.412	1.1512			

(Masson, Chem. Soc. 1911, 29. 1132.)

Solubility in HCl + Aq at 25°.	
Millimols HCl in 10 ccm.	Millimols NaCl in 10 ccm.
...	54.56
6.07	48.50
10.32	44.67
15.90	37.82
21.17	32.97
32.83	23.43

(Herz, Z. anorg. 1912, 73. 274.)

Moderately dil. H₂SO₄ or HNO₃ + Aq precipitate NaCl from NaCl + Aq. (Karsten.)
Sol. in H₂SO₄. (Walden, Z. anorg. 1902, 29. 384.)

Solubility of NaCl in NH₄OH + Aq at 30°.
(G. in 1 l. sat. solution.)

Sp. gr.	NH ₃	NaCl	Sp. gr.	NH ₃	NaCl
1.1735	29.535	293.38	1.1406	72.07	283.38
1.1656	40.655	292.5	1.1395	72.715	283.06
1.160	47.26	289.7	1.1301	81.855	277.49
1.1494	60.78	286.5	1.205	97.49	270.57

(Hempel and Tedesco, Z. anorg. 1911, 24. 2467.)

Solubility of NaCl in NH₄Cl + Aq at t°.

t°	G. per 100 g. H ₂ O		Sp. gr.
	NH ₄ Cl	NaCl	
0	0	356.3	...
	146.1	286.4	1.185
15	0	357.6	1.200
	57.3	326.4	1.191
	118.9	300	1.183
	186.4	271.6	1.176
	198.8	266.8	1.175
30	0	360.3	...
	255.4	249	1.166
45	0	365	...
	322.1	233.9	...

(Fedotieff, Z. phys. Ch. 1904, 49. 168.)
See also under NH₄Cl.

Solubility in NaOH + Aq at 0°. NaCl = mols. NaCl (in milligrams) in 10 ccm. solution; Na₂O = mols. Na₂O (in milligrams) in 10 ccm. solution.

NaCl	Na ₂ O	Na ₂ O + NaCl	Sp. g.
54.7	0	54.7	1.207
49.375	4.8	54.175	1.221
47.212	6.725	53.937	1.225
42.375	10.406	52.781	1.236
39.55	14.78	54.33	1.249
24.95	30.5	55.45	1.286
19.3	37.875	57.175	1.314
9.408	53.25	62.66	1.362

(Engel, C. R. 112. 1130.)

Solubility in NaOH + Aq at 20°.

G. NaOH in 1 liter	G. NaCl in 1 liter	Sp. gr.	deg. Baumé
10	308	1.200	23.5
20	308	1.210	24.0
30	306	1.215	25.5
40	302	1.225	26.4
50	297	1.230	26.9
60	286	1.235	27.4
70	277	1.240	27.9
80	269	1.245	28.4
90	261	1.250	28.8
100	253	1.250	28.8
110	244	1.252	29.0
120	236	1.252	29.0
130	229	1.260	29.7
140	221	1.265	30.2
150	213	1.270	30.6
160	205	1.275	31.1
170	197	1.275	31.1
180	189	1.280	31.5
190	181	1.285	32.0
200	173	1.290	32.4
210	165	1.295	32.8
220	159	1.295	32.8
230	152	1.300	33.3
240	146	1.303	33.5
250	139	1.305	33.7
260	134	1.310	34.2
270	129	1.315	34.6
280	124	1.320	35.0
290	118	1.325	35.4
300	112	1.330	35.8
310	107	1.333	36.0
320	101	1.335	36.2
330	96	1.340	36.6
340	90	1.345	37.0
350	85	1.350	37.4
360	80	1.355	37.8
370	76	1.360	38.2
380	71	1.365	38.6
390	66	1.370	39.0
400	61	1.375	39.4
410	56	1.380	40.0
420	52	1.385	40.2
430	48	1.390	40.6

y in NaOH + Aq at 20°.—Continued.

in	G. NaCl in 1 liter	Sp. gr.	deg. Baumé
	45	1.395	41.0
	42	1.400	41.5
	39	1.405	41.9
	37	1.410	42.0
	34	1.415	42.3
	32	1.420	42.6
	30	1.425	43.0
	28	1.430	43.5
	27	1.435	43.7
	27	1.440	44.0
	26	1.445	44.3
	26	1.450	44.6
	25	1.450	44.6
	24	1.455	45.0
	23	1.460	45.5
	23	1.465	45.9
	22	1.470	46.2
	21	1.475	46.5
	20	1.480	46.8
	19	1.485	47.0
	18	1.490	47.5

eler, Z. Elektrochem, 1900, 7. 360.)

solubility in Na₂O + Aq at 30°.

tion of the solution		Solid phase
t.	% by wt. NaCl	
	26.47	NaCl
7	21.49	"
2	13.62	"
8	4.36	"
1	2.40	"
5	1.12	"
2	0.97	NaCl + NaOH, H ₂ O
	0	NaOH, H ₂ O

emakers, Z. phys. Ch. 1909, 66. 85.)

presence of other salts increases the
y of NaCl in H₂O.in sat. NH₄Cl + Aq with pptn. of

When the reaction is complete,
tion has sp. gr. 1.1788, and contains
mixed salts, or 100 pts. H₂O dissolve
sa. mixed salts, viz., 26.36 pts. NaCl
.06 pts. NH₄Cl. (Karsten.) (See
[H₂Cl].)

2 sat. BaCl₂ + Aq with pptn. of BaCl₂
state of equilibrium is reached, when
H₂O at 17° dissolve 38.6 pts. of mixed
which 4.1 pts. are BaCl₂. (Karsten.)
ler BaCl₂.)

in sat. CaCl₂ + Aq (Vauquelin,
Chim. 13. 95.)

more sol. in hot than in cold H₂O
ng MgCl₂ or CaCl₂; but NaCl is pptd.
t. NaCl + Aq when that solution is
ith MgCl₂ or CaCl₂ + Aq. (Fuchs and
benbach, 1826.) (See under MgCl₂.)

Less sol. in conc. CaCl₂ + Aq than in H₂O.
(Hermann.)

Solubility of NaCl + CaCl₂ in H₂O at 25°.
G. per 100 g. H₂O.

NaCl	CaCl ₂	Sp. gr. 25°/25°	Solid phase
0	84	CaCl ₂ , 6H ₂ O
1.846	78.49	1.4441	" + NaCl
1.637	58.48	1.3651	NaCl
1.799	53.47	1.3463	"
7.77	36.80	1.2831	"
10.70	30.08	1.2653	"
18.85	19.53	1.2367	"
32.48	3.92	1.2080	"
35.80	0	1.2030	"

(Cameron, Bell and Robinson, J. phys. Chem.
1907, 11. 396.)

Solubility of NaCl in NaHCO₃ sat. with
CO₂ at t°.

t°	G. per 1000 g. H ₂ O	
	NaHCO ₃	NaCl
0	6.0	356.3
"	7.7	350.1
15	0.0	357.6
"	10.0	354.6
30	0.0	360.3
"	13.9	358.1
45	0.0	356.0
"	0.23	361.5

(Fedotieff, Z. phys. Ch. 1904, 49. 170.)

Sol. in sat. KClO₃ + Aq; the solution can
then dissolve more KClO₃. (Marguerite,
C. R. 36. 305.)

In solution containing Na, K, Cl and NO₃
ions, the solubility—relations between the
four salts NaCl, KCl, NaNO₃ and KNO₃
have been studied. (Uyeda, Z. anorg. 1911,
71. 2.)

Sol. in sat. NH₄NO₃ + Aq, without causing
pptn. (Karsten.)

Sol. in sat. NH₄NO₃ + Aq, from which solu-
tion it is not pptd. by salts which would cause
its pptn. in aqueous solution (Marguerite,
C. R. 36. 307.)

Sol. in sat. Ba(NO₃)₂ + Aq without causing
pptn.

Insol. in Ca(NO₃)₂ + Aq.

Sol. in Mg(NO₃)₂ + Aq with pptn. of small
portion of Mg(NO₃)₂.

Sol. in sat. KNO₃ + Aq, the mixed solution
having the power to dissolve more KNO₃, and
the solubility of the KNO₃ apparently in-
creasing in the same ratio as the amount of
NaCl present. (Fourcroy and Vauquelin,
Ann. de Chim. 11. 130.)

Sol. in sat. KNO₃ + Aq; the solution thus

obtained at 18.13° contains 40.34% of the mixed salts, or 100 pts. H₂O dissolve 67.72 pts. of the mixed salts, viz., 38.25 pts. NaCl and 29.45 pts. KNO₃. (Karsten.)

Solubility of NaCl in KNO₃+Aq at 25°. KNO₃=g. KNO₃ in 100 cc. of solution. NaCl=g. mol. per l.

KNO ₃	NaCl	KNO ₃	NaCl
0	5.44	12	5.28
4	5.52	16	5.21
8	5.45	20	5.15

(Ritzel, Z. Krist. 1911, 49. 152.)
(See also under KNO₃.)

Solubility of NaCl in NaNO₃+Aq at 15.5°.

Sp. gr.	g. per 100 cc. sat. solution		
	NaNO ₃	NaCl	H ₂ O
1.2025	0	31.78	88.47
1.2305	7.53	27.89	87.63
1.2580	13.24	26.31	86.25
1.2810	21.58	23.98	82.66
1.3090	28.18	22.30	80.42
1.3345	33.80	20.40	79.25
1.3465	37.88*	19.40*	77.37
1.3465	37.64*	19.67*	77.34

*Solutions sat. with both salts.
(Bodländer, Z. phys. Ch. 1891, 1. 361.)

Sol. in sat. NaNO₃+Aq with pptn. of NaNO₃.

Solubility of NaCl in NaNO₃+Aq.
Conc.=concentration of NaNO₃ in g. mol. per 1,000 g. H₂O.
NaCl=g. NaCl dissolved in 1,000 g. H₂O.

t°	Conc.	NaCl	Molecular solubility
0	0	359.65	6.16
	0.25	355.90	6.09
	0.50	351.20	6.02
	1	342.15	5.86
25	0	362.95	6.20
	0.25	356.65	6.11
	0.50	352.30	6.03
	1	343.65	5.88
	2	325.50	5.58

(Armstrong and Eyre, Proc. R. Soc. 1910, A. 84. 127.)
(See also under NaNO₃.)

Sol. in sat. KCl+Aq with elevation of temp. (Vauquelin.)
100 g. H₂O sat. with KCl dissolve 0.494 gram-equivalent NaCl at 25°. (Fuler, Z. phys. Ch. 1904, 49. 315.)

Solubility in KCl+Aq at t°.

t°	Sat. solution contains	
	% NaCl	% KCl
-20	21.3	5.7
-10	21.3	6.7
0	21.3	7.7
+10	21.3	8.6
20	21.3	9.6
30	21.3	10.6
40	21.3	11.5
50	21.3	12.5
60	21.3	13.5
70	21.3	14.4
80	20.7	15.3
90	19.9	17.8
100	18.8	19.8
110	17.2	22.4
120	16.5	24.1
130	16.4	25.1
140	16.4	26.1
150	16.4	27.1
160	16.4	28.0
170	16.4	29.0
180	16.4	30.0

(Étard. A. ch. 1894, (7) 2. 277.)
(See under KCl.)

100 pts. NaCl+NaI+Aq sat. at 18-19° contain 62.33 pts. of the two salts. (Hauer.)

Sol. in sat. Al₂(SO₄)₃+Aq with no ppt. (Vauquelin.)

Sol. in sat. (NH₄)₂SO₄+Aq with pptn. of considerable amt. of (NH₄)₂SO₄+Aq. (Vauquelin.)

Sol. in sat. CuSO₄+Aq.
100 pts. H₂O dissolve 36.71 pts. NaCl and 7.19 pts. K₂SO₄ at 15°, and solution has sp. gr. 1.24. (Page and Keightey.)

NaCl is sol. in K₂SO₄+Aq, and vice versa, without separation of a salt.

100 pts. H₂O dissolve 7.03 pts. K₂SO₄ and 37.60 pts. NaCl, when warmed and cooled to 14°. (Rüdorff.)

Solubility of NaCl and K₂SO₄ in H₂O at t°. 100 pts. H₂O contain pts. NaCl, K₂SO₄, and KCl.

t°	Pts. NaCl	Pts. K ₂ SO ₄	Pts. KCl
10	33.43	8.10	3.18
20	34.01	8.90	3.06
30	34.56	9.56	2.95
40	35.16	10.38	2.81
50	35.77	11.07	2.64
60	36.40	11.93	2.72
70	36.64	12.82	3.30
80	36.04	12.26	5.06
90	35.86	12.42	6.96
100	35.63	12.56	8.79

(Precht and Wittgen, B. 18. 1006.)

. in cold sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ at first without, afterwards Na_2SO_4 separates out. (Karsten.)

solubility in $\text{Na}_2\text{SO}_4 + \text{Aq}$ containing 7.45 g. Na_2SO_4 in 100 g. of the solution.

t°	g. NaCl in 100 g. of the solution
14.80	23.30
17.90	23.33
24.85	23.45
25.60	23.485
27.75	23.525
32.18	23.55
34.28	23.68

Gerardin and Marquis, C. R. 1903, 136. 684.)
also under Na_2SO_4 .

1. in sat. $\text{ZnSO}_4 + \text{Aq}$ with separation of O_2 , ZnSO_4 . (Karsten.)
sol. in liquid CO_2 . (Büchner, Z. phys. 1906, 54. 674.)
moderately sol. in liquid NH_3 . (Franklin, Ch. J. 1898, 20. 829.)
2 pts. NaCl are sol. in 1 pt. hydrazine at -13° . (de Bruyn, R. t. c. 1899, 18. 297.)
10 g. hydroxylamine dissolve 14.7 g. NaCl at -15° . (de Bruyn, Z. phys. Ch. 1892, 10.

10 g. 95% formic acid dissolve 5.8 g. at -1° . (Aschan, Ch. Ztg. 1913, 37. 1117.)
solubility in alcohol.

100 pts. alcohol of 0.900 sp. gr. dissolve 5.8 pts. NaCl ; 72 sp. gr. dissolve 3.67 pts. NaCl ; of 0.834 sp. gr. dissolve 0.5 pt. NaCl . (Kirwan.)

pts. alcohol containing given % by weight of absolute alcohol dissolve pts. NaCl at 25° .

	Pts. NaCl	% alcohol	Pts. NaCl	% alcohol	Pts. NaCl
0	35.70	33.4	16.08	66.9	5.95
1	30.49	41.8	13.28	75.2	3.75
2	24.84	50.2	11.28	83.6	1.59
3	19.30	58.5	7.96

(Kopp, A. 40. 206.)

pts. alcohol of 75% by weight dissolve at:
 14° 15.2° 38° 71.5°
0.661 0.700 0.736 1.033 pts. NaCl .

pts. alcohol of 95.5% by weight dissolve at:
 15° 77.2°
0.174 0.171 pts. NaCl .
(Wagner, A. 64. 293.)

0 pts. alcohol containing % alcohol by weight dissolve pts. NaCl at 15° , or 100 pts. solution contain % NaCl .
20 30 40 % alcohol,
53 22.55 17.51 13.25 pts. NaCl ,
12 18.4 14.9 11.7 % NaCl ,
50 60 80 % alcohol,
9.77 5.93 1.22 pts. NaCl ,
8.9 5.6 1.2 % NaCl .
(Schiff, A. 118. 365.)

Solubility of NaCl in alcohol increases with the temperature.
100 pts. (by weight) of alcohol of 0.9282 sp. gr. (50.5% by weight) dissolve at:
 4° 10° 13° 23° 32°
10.9 11.1 11.43 11.9 12.3 pts. NaCl ,
 33° 44° 51° 60°
12.5 13.1 13.8 14.1 pts. NaCl .
(Gerardin, A. ch. (4) 5. 146.)

Solubility in alcohol at 13° .

Sp. gr.	100 ccm. contain in g.		
	Alcohol	Water	Salt
1.2030	0	88.70	31.60
1.1348	11.81	78.41	23.26
1.1144	15.99	74.64	20.81
1.0970	19.39	71.45	18.86
1.0698	24.95	65.80	16.23
1.0295	32.33	57.96	12.66
0.9880	40.33	49.34	9.13
0.9445	49.28	38.54	5.93
0.9075	57.91	29.37	3.47
0.8700	63.86	21.62	1.52
0.8400	72.26	11.24	0.50

(Bodländer, Z. phys. Ch. 7. 317.)

Solubility in ethyl alcohol + Aq at 30° .

wt. % alcohol	g. NaCl per 100 g.		wt. % alcohol	g. NaCl per 100 g.	
	solution	H_2O		solution	H_2O
0	26.50	36.05	50	9.34	20.60
5	24.59	34.29	60	6.36	16.96
10	22.56	32.57	70	3.36	12.75
20	19.05	29.40	80	1.56	7.95
30	15.67	26.53	90	0.43	4.30
40	12.45	23.70

(Taylor, J. phys. Ch. 1897, 1. 723.)

Solubility in ethyl alcohol + Aq at 40° .

wt. % alcohol	g. NaCl per 100 g.		wt. % alcohol	g. NaCl + 100 g.	
	solution	H_2O		solution	H_2O
0	26.68	36.38	50	9.67	21.42
5	24.79	34.69	60	6.65	17.82
10	22.90	33.00	70	3.87	13.10
20	19.46	30.20	80	1.69	8.68
30	16.02	27.25	90	0.50	5.10
40	12.75	24.37

(Taylor, l. c.)

Solubility of NaCl in ethyl alcohol+Aq at 28°.

% C ₂ H ₅ OH	% H ₂ O	% NaCl	% C ₂ H ₅ OH	% H ₂ O	% NaCl
0	73.53	26.47	45.35	45.35	9.3
3.8	71.6	24.6	56.2	37.5	6.3
7.7	69.7	22.6	67.4	28.9	3.7
16.1	64.6	19.3	78.8	19.7	1.5
25.3	58.9	15.8	89.6	10	0.4
35.0	52.5	12.5			

(Fontein, Z. phys. Ch. 1904, 73. 212.)

Solubility of NaCl in ethyl alcohol+Aq at 25°.

Conc. = concentration of alcohol in g. mol. per 1,000 g. H₂O.
NaCl = g. in 1,000 g. H₂O.

Conc.	NaCl	Molecular solubility
0	359.65	6.16
0.25	355.15	6.08
0.50	349.65	5.98
1	337.80	5.79
3	301.60	5.16

(Armstrong and Eyre, Proc. R. Soc. 1910, (A) 84. 127.)

100 pts. absolute methyl alcohol dissolve 1.41 pts. at 18.5°; 100 pts. absolute ethyl alcohol dissolve 0.065 pt. at 18.5°. (de Bruyn, Z. phys. Ch. 10. 782.)

100 pts. wood-spirit of 40% (by weight) dissolve 13.0 pts. NaCl. (Schiff, A. 118. 365.)

100 g. NaCl+CH₃OH contain 0.1 g. NaCl at the critical temp. (Centnerszwer, Z. phys. Ch. 1910, 72. 437.)

Solubility of NaCl in methyl alcohol+Aq. at 25°.

Conc. = concentration of alcohol in g. mol. per 1,000 g. H₂O.
NaCl = g. in 1,000 g. H₂O.

t°	Conc.	NaCl	Molecular solubility
0°	0	357.75	6.13
	0.25	355.20	6.08
	0.50	353.10	6.05
	1	347.45	5.95
25°	0	362.95	6.20
	0.25	359.40	6.14
	0.50	357.60	6.11
	1	353.20	6.04
	3	336.25	5.75

(Armstrong and Eyre, Proc. R. Soc. 1910, (A) 84. 127.)

Traces dissolve in propyl alcohol. (Schlamp Z. phys. Ch. 1894, 14. 276.)

Solubility of NaCl in propyl alcohol+Aq. at 25°.

Conc. = concentration of alcohol in g. mol. per 1,000 g. H₂O.
NaCl = g. NaCl in 1,000 g. H₂O.

t°	Conc.	NaCl	Molecular solubility
0	0	357.75	6.13
	0.25	351.20	6.01
	0.50	345.55	5.91
25	0	362.95	6.20
	0.25	355.75	6.10
	0.50	350.20	6.00

(Armstrong and Eyre, Proc. R. Soc. 1910, A 84. 127.)

Solubility of NaCl in propyl alcohol+Aq at 23-25°.

% NaCl	% C ₃ H ₇ OH	% H ₂ O	% NaCl	% C ₃ H ₇ OH	% H ₂ O
0.55	87.7	11.75	14.38	5.39	80.23
2.23	51.57	46.20	15.42	5.11	79.47
3.55	18.99	77.46	16.38	4.47	79.14
3.90	14.78	81.32	18.08	3.83	78.09
5.27	12.77	81.96	20.12	3.27	76.61
8.04	9.49	82.47	22.35	2.64	75.01
10.49	7.79	81.72	24.50	2.13	75.37
12.20	6.57	81.23	24.90	2.3	72.80

(Frankforter and Frary, J. phys. Ch. 1912, 17. 402.)

100 g. sat. solution of NaCl in 99.6% propyl alcohol contain 0.04 g. NaCl at 25°. (Frankforter and Frary.)

Insol. in fusel oil. (Gooch, Am. Ch. J. 2 53.)

Solubility of NaCl in amyl alcohol+Aq. Liquid phases conjugated at 28°.

Upper layer			Lower layer		
NaCl	amyl alcohol	H ₂ O	NaCl	amyl alcohol	H ₂ O
0.05	95.45	4.5	26.36	0.22	73.42
"	94.1	5.9	19	0.4	80.4
"	92.9	7.1	12.7	0.8	86.5
"	91.6	8.4	6.2	1.5	92.3
0	90.2	9.8	0	2.3	97.7

(Fontein, Z. phys. Ch. 1910, 73. 226.)

At room temp. 1 pt. by weight is sol. in:
75 pts. methyl alcohol D₁₅ 0.7990.
566 " ethyl " D₁₅ 0.8100.
3000 " propyl " D₁₅ 0.8160.
(Rohland, Z. anorg. 1898, 18. 325.)

100 g. methyl alcohol dissolve 1.31 g. NaCl at 25°.

ethyl alcohol dissolve 0.065 g. NaCl

propyl alcohol dissolve 0.012 g. NaCl

isoamyl alcohol dissolve 0.002 g. 5°.

and Bissett, Chem. Soc. 1913, 103. 1909.)

of NaCl in ethyl+amyl alcohol at 28°.

Upper layer		Lower layer		
amyl alcohol	% ethyl alcohol	% NaCl	% amyl alcohol	% ethyl alcohol
45	0	26.35	0.22	0
6	9.5	25.30	0.25	1.9
4	19.1	24.02	0.3	9.5
9	30.9	22.64	0.4	6.9
0	38.7	21.19	0.5	10.3
6	44.8	19.26	1.3	15.2
7	41.5	15.81	3.69	22.1

solution: amyl alcohol, 10%; alcohol, 32.5%; NaCl, 11%; H₂O,

in, Z. phys. Ch. 1910, 73. 244.)

ppts. NaCl from NaCl+Aq.

sol. in a mixture of equal pts. of alcohol and ether. (Berzelius.)

NaCl treated with above mixture by 0.5 mg. to the liquid. (Lawrence A. J. Sci. (2) 16. 57.)

of a mixture of 1 pt. 96% alcohol 98% ether dissolve 0.11 pt. NaCl. . 98. 205.)

acetone. (Krug and M'Elroy, J. 6. 184; Eidmann, C. C. 1899, II.

bility in acetone+Aq at 20°.

acetone in 100 ccm. of the solvent. millimols NaCl in 100 ccm. of the

A	NaCl
	537.9
	464.6
	394.8
	330.1
phases	lower layer 308.5
	upper layer 7.7
	7.3
	5.6
	4.3
	...

l Knoch, Z. anorg. 1904, 41. 318.)

Solubility in acetone+Aq at 20°.

% NaCl	% H ₂ O	% acetone
25.9	73.06	1.04
24.19	71.18	4.03
25.06	72.00	2.94
20.85	66.78	12.37
20.17	66.01	13.82
18.32	63.16	18.52
20.44	66.19	13.37
17.89	62.21	19.90
0.32	13.92	85.76
0.19	10.82	88.99
0.15	9.62	90.23
0.12	8.94	90.94

(Frankforter and Cohen, J. Am. Chem. Soc. 1914, 36. 1127.)

Solubility in glycol at 14.8°=31.7%. (de Coninck, Belg. Acad. Bull. 1905. 275.)

Sol. in glycerine. (Pelouze.)

Solubility of NaCl in glycerine+Aq at 25°.

G=g. glycerine in 100 g. glycerine+Aq.

NaCl=millimols NaCl in 100 cc. of the solution.

G	NaCl	Sp. gr.
0	545.6	1.1960
13.28	501.1	1.2048
25.98	448.4	1.2133
45.36	370.2	1.2283
54.23	333.9	1.2381
83.84	220.8	1.2696
100	167.1	1.2964

(Herz and Knoch, Z. anorg. 1905, 45. 267.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3602.)

Solubility of NaCl in solutions of HgCl₂ in ethyl acetate.

100 mols. ethyl acetate dissolve			
Mols. HgCl ₂	Mols. NaCl	Mols. HgCl ₂	Mols. NaCl
40.0	20.0	18.0	5.1
38.1	19.6	16.4	4.3
36.0	19.2	14.1	3.8
34.9	18.5	13.2	2.9
34.8	18.3	12.4	2.3
32.1	13.8	12.0	1.6
28.0	9.1	12.2	1.3
22.8	7.0	12.9	0.8
22.9	7.0

(Linebarger, Am. Ch. J. 1894, 16. 215.)

Solubility of NaCl in urea + Aq at 25°.

% urea	% NaCl	% urea	% NaCl
0	31.80	18	27.65
5	30.63	23	27.24
9.6	29.05	28	26.56
III	28.46

(Ritsel, Z. Kryst. Min. 1911, 49. 152.)

Solubility of NaCl in urea + Aq at (?)°.

g = g. urea in 100 cc. of solution.

sol. = increase of solubility of NaCl in g. per 100 cc. of the solution.

g.	sol.	g.	sol.
5	0.044	30	0.709
10	0.124	35	0.910
15	0.234	40	1.134
20	0.372	45	1.379
25	0.529	50	1.602

(Fastert, N. Jahrb. Min. Beil. Bd. 1912, 23. 286.)

Solubility of NaCl in formamide + Aq at 25°.

% HCONH ₂	% NaCl	% HCONH ₂	% NaCl
0	31.80	11	29.11
2.3	30.98	15	28.52
5.3	30.86	18.8	27.76
8	30.40

(Ritsel, Z. Kryst. Min. 1911, 49. 152.)

Insol. in anhydrous and in 97% pyridine.
 Very sl. sol. in 95% pyridine + Aq. Sl. sol. in 93% pyridine + Aq. (Kahlenberg, J. Am. Chem. Soc. 1908, 30. 1107.)

Insol. in oil of turpentine. (T. S. Hunt, Am. J. Sci. (2) 19. 417.)

100 g. H₂O dissolve 236.3 g. sugar + 42.3 g. NaCl at 31.25°, or 100 g. sat. aq. solution contain 62.17 g. sugar + 11.13 g. NaCl (Köhler, Z. Ver. Zuckerind. 1897, 47. 447.)

Solubility of NaCl in glucose + Aq at 25°.

Conc. = concentration of glucose + Aq in g. mol. per 1,000 g. H₂O

Sol. = Solubility in 1,000 g. H₂O

Conc.	Sol.	Molecular solubility
0	361.40	6.18
0.25	364.15	6.22
0.50	364.30	6.23
1.0	369.90	6.32

(Armstrong and Eyre, Proc. R. Soc. 1910, (A) 84. 127.)

Min. Halite.

+2H₂O. Efflorescent below 0°; sl. deliquescent at temps. above 0°. (Fuchs, 1826.)

The solubility in H₂O at 12.25° corresponds to 32.9 pts. of NaCl per 100 pts. H₂O (Matignon, C. R. 1909, 143. 551.)

Sodium stannic chloride, 2NaCl, 6H₂O.

See Chlorostannate, sodium.

Sodium thallic chloride, 3NaCl, 12H₂O.

Very sol. in H₂O. (Pratt, Am. 1895, (3) 49. 404.)

Sodium uranium chloride, 2NaCl, U

Non volatile and not hygroscopic. (San, C. R. 1896, 122. 1089.)

Sol. in H₂O. (Colani, A. ch. 1907 59.)

Sodium uranyl chloride, Na₂(UO₂)Cl

As K salt. (Aloy, Bull. Soc. 1899 264.)

Sodium zinc chloride, 2NaCl, ZnCl₂.

Deliquescent. Easily sol. in H₂O. (ler, Mag. Pharm. 36. 48.)

Sodium zirconium chloride, 2NaCl, Z

(Paykull.)

Sodium chloriodide, NaCl, I + 2H₂O

Easily decomp. by alcohol or ether and Wheeler, Sil. Am. J. 143. 42.)

Sodium fluoride, NaF.

Very sl. sol. in cold, and not mol. in boiling H₂O. (Rose.)

100 pts. H₂O dissolve 4.78 pts. at 11° zehus.)

100 pts. H₂O dissolve 4 pts. at 15°. A. ch. (3) 47. 32.)

Sp. gr. of aqueous solutions containing 100 pts. H₂O:

1.1081	2.2162	3.3243 pts.
1.0110	1.0221	1.0333

Sat. solution has sp. gr. 1.0486. (Z. anal. 27. 277.)

Sp. gr. of solution sat. at 18° = 1.0486. (Mylus and 1897, 30. 1718.)

Solubility of NaF in HF + Aq at

g. per 1000 g. H ₂ O		g. per 1000 g.	
HF	NaF	HF	
0.0	41.7	83.8	
10.0	41.4	129.7	
45.8	22.5	596.4	
56.5	22.7	777.4	

(Ditte, C. R. 1896, 123. 1283)

Easily sol. in liquid HF. (Franz, anorg. 1905, 46. 2.)

Sl. sol. in conc. KC₂H₃O₂ + Aq. (eyer.)

Almost insol. in alcohol. (Berthel, 1. 13.)

Insol. in methyl acetate. (Nuss, 1909, 42. 3793.)

hydrogen fluoride, NaHF_2 .
r difficultly sol. in cold, more easily
 H_2O . (Berselius, Pogg. 1. 13.)

tantalum fluoride.
notantalate, sodium.

tin (stannous) fluoride, 2NaF , 3SnF_2 .
 H_2O . (Wagner, B. 19. 896.)

tin (stannic) fluoride.
notannate, sodium.

tantalum fluoride.
notantalate, sodium.

tellurium fluoride, NaF , TeF_4 .
sp. by H_2O . (Berselius.)

titanium fluoride.
notitanate, sodium.

tungstyl fluoride.
oxytungstate, sodium.

uranium fluoride, NaF , UF_4 (?).
that soluble in H_2O . (Bolton.)

uranyl fluoride.
oxyuranate, sodium.

vanadium sesquifluoride.
ovanadate, sodium.

zinc fluoride, NaF , ZnF_2 .
 H_2O . (R. Wagner.)

zirconium fluoride, 5NaF , 2ZrF_4 .
ozirconate, sodium.

fluoride vanadium pentoxide.
oxyvanadate, sodium.

hydrazide, NH_2 , NHNa .
p. by H_2O with explosive violence.
by alcohol. (Schlenk, B. 1915, 48.)

hydride, NaH .
p. by H_2O and by acids.
fused Na or Na amalgam. Insol. in
 H_2 . Insol. in CS_2 , CCl_4 , C_6H_6 and
ene. (Moissan, C. R. 1902, 134.)

Decomp. violently by H_2O .

hydrosulphide, NaSH .
escent. Sol. in H_2O and alcohol.
) . Difficultly sol. in H_2O . (Damoi-
C. 1885. 36.)

hydroxide, NaOH .
eliquescent. 100 pts. NaOH under
with H_2O at 16–20° absorb 552 pts.
s. (Mulder.)
ol. in H_2O with evolution of much
l. in 0.47 pt. H_2O . (Bineau, C. R.

Solubility of NaOH in H_2O .

t°	g. per 100 g.		Solid phase
	Solu- tion	H_2O	
— 7.8	8.0	8.7	Ice
—20	16.0	19.1	"
—28	19.0	23.5	Ice + $\text{NaOH} \cdot 7\text{H}_2\text{O}$
—24	22.2	28.5	$\text{NaOH} \cdot 7\text{H}_2\text{O} + \text{NaOH} \cdot 5\text{H}_2\text{O}$
—17.7	24.5	32.5	$\text{NaOH} \cdot 5\text{H}_2\text{O} + \text{NaOH} \cdot 4\text{H}_2\text{O}$
0	29.6	42.0	$\text{NaOH} \cdot 4\text{H}_2\text{O}$
+ 5	32.2	47.5	$\text{NaOH} \cdot 4\text{H}_2\text{O} + \text{NaOH} \cdot 3\frac{1}{2}\text{H}_2\text{O}$
10	34.0	51.5	$\text{NaOH} \cdot 3\frac{1}{2}\text{H}_2\text{O}$
15.5	38.9	63.53	" f. pt.
5	45.5	83.5	$\text{NaOH} \cdot 3\frac{1}{2}\text{H}_2\text{O} + \text{NaOH} \cdot 2\text{H}_2\text{O}$
12	50.7	103.0	$\text{NaOH} \cdot 2\text{H}_2\text{O} + \text{NaOH} \cdot \text{H}_2\text{O}$
20	52.2	109	$\text{NaOH} \cdot \text{H}_2\text{O}$
30	54.3	119	"
40	56.3	129	"
50	59.2	145	"
60	63.5	174	"
64.3	69.0	222.3	" f. pt.
61.8	74.2	288	$\text{NaOH} \cdot \text{H}_2\text{O} + \text{NaOH}$
80	75.8	313	NaOH (?)
110	78.5	365	"
192	83.9	521	"

(Pickering, Chem. Soc. 1893, 63. 890; Mylius and Funk, W. A. B. 1900, 3. 450. Calc. by Seidell, Solubilities, 2d Edition, p. 653.)

100 g. sat. $\text{NaOH} + \text{Aq}$ at 15° contain 46.36 g. NaOH . (de Forcrand, C. R. 149. 1344.)

Sp. gr. and b-pt of $\text{NaOH} + \text{Aq}$.

% Na_2O	Sp. gr.	B.-pt.	% Na_2O	Sp. gr.	B.-pt.
4.7	1.06	100.56°	31.0	1.44	120.00°
9.0	1.12	101.11	34.0	1.47	123.89
13.0	1.18	102.78	36.8	1.50	129.44
16.0	1.23	104.44	41.2	1.56	137.78
19.0	1.29	106.66	46.6	1.63	148.89
23.0	1.32	108.89	53.8	1.72	204.44
26.0	1.36	112.78	63.6	1.85	315.56
29.0	1.40	116.66	77.8	2.00	red heat.

(Dalton.)

Sp. gr. of $\text{NaOH} + \text{Aq}$ at 15°.

% Na_2O	Sp. gr.	% Na_2O	Sp. gr.	% Na_2O	Sp. gr.
0.302	1.0040	10.879	1.1630	21.154	1.3053
0.601	1.0081	11.484	1.1734	21.758	1.3125
1.209	1.0163	12.088	1.1841	21.894	1.3143
1.813	1.0246	12.692	1.1948	22.363	1.3198
2.418	1.0330	13.297	1.2058	22.967	1.3273
3.022	1.0414	13.901	1.2178	23.572	1.3349
3.626	1.0500	14.506	1.2280	24.176	1.3426
4.231	1.0587	15.110	1.2392	24.780	1.3505
4.835	1.0675	15.714	1.2453	25.385	1.3586
5.440	1.0764	16.319	1.2515	25.989	1.3668
6.044	1.0855	16.923	1.2578	26.594	1.3751
6.648	1.0948	17.528	1.2642	27.200	1.3836
7.253	1.1042	18.132	1.2708	27.802	1.3923
7.857	1.1137	18.730	1.2775	28.407	1.4011
8.462	1.1233	19.341	1.2843	29.011	1.4101
9.066	1.1330	19.954	1.2912	29.616	1.4193
9.670	1.1428	20.550	1.2982	30.220	1.4285
10.275	1.1528

(Tünnerman, N. J. Pharm. 18. 2.)

Sp. gr. of NaOH + Aq.

% NaOH	Sp. gr.	% NaOH	Sp. gr.	% NaOH	Sp. gr.
2.07	1.02	14.73	1.16	28.16	1.30
4.02	1.04	16.73	1.18	29.96	1.32
5.89	1.06	18.71	1.20	31.67	1.34
7.69	1.08	20.66	1.22	32.40	1.35
9.43	1.10	22.58	1.24	33.08	1.36
11.10	1.12	24.47	1.26	34.41	1.38
12.81	1.14	26.33	1.28

(Richter.)

Sp. gr. of NaOH + Aq at 15°.

%	Sp. gr. if % is NaOH	Sp. gr. if % is NaOH	%	Sp. gr. if % is NaOH	Sp. gr. if % is NaOH
1	1.015	1.012	32	1.450	1.351
2	1.020	1.023	33	1.462	1.363
3	1.043	1.035	34	1.475	1.374
4	1.067	1.046	35	1.488	1.384
5	1.074	1.059	36	1.500	1.395
6	1.089	1.070	37	1.515	1.405
7	1.104	1.081	38	1.530	1.415
8	1.119	1.092	39	1.543	1.426
9	1.132	1.103	40	1.558	1.437
10	1.145	1.115	41	1.570	1.447
11	1.160	1.126	42	1.583	1.456
12	1.175	1.137	43	1.597	1.468
13	1.190	1.148	44	1.610	1.478
14	1.205	1.159	45	1.623	1.488
15	1.219	1.170	46	1.637	1.499
16	1.233	1.181	47	1.650	1.508
17	1.245	1.192	48	1.663	1.519
18	1.258	1.202	49	1.678	1.529
19	1.270	1.213	50	1.690	1.540
20	1.285	1.225	51	1.705	1.550
21	1.300	1.236	52	1.719	1.560
22	1.315	1.247	53	1.730	1.570
23	1.329	1.258	54	1.745	1.580
24	1.341	1.269	55	1.760	1.591
25	1.355	1.279	56	1.770	1.601
26	1.369	1.290	57	1.785	1.611
27	1.381	1.300	58	1.800	1.622
28	1.395	1.310	59	1.815	1.633
29	1.410	1.321	60	1.830	1.643
30	1.422	1.332	70	...	1.748
31	1.438	1.343			

(Gerlach, Z. anal. 8. 279, calculated from Schiff, A. 107. 300.)

Sp. gr. of NaOH + Aq at 15°.

% NaOH	Sp. gr.	% NaOH	Sp. gr.
0.61	1.0070	4.0	1.0435
0.9	1.0105	4.32	1.0473
1.0	1.0107	4.64	1.0511
1.2	1.0141	4.96	1.0549
1.6	1.0177	5.29	1.0588
2.0	1.0213	5.58	1.0627
2.36	1.0249	5.87	1.0667
2.71	1.0286	6.21	1.0706
3.0	1.0318	6.55	1.0746
3.35	1.0360	6.76	1.0787
3.67	1.0397	7.31	1.0827

Sp. gr. of NaOH + Aq at 5°—Continued

% NaOH	Sp. gr.	% NaOH	Sp. gr.
7.66	1.0868	24.81	1.2746
8.0	1.0909	25.3	1.2800
8.34	1.0951	25.8	1.2857
8.68	1.0992	26.31	1.2906
9.0	1.1030	26.83	1.2973
9.42	1.1077	27.31	1.3032
9.74	1.1120	27.8	1.3093
10.0	1.1158	28.31	1.3151
10.5	1.1195	28.83	1.3211
10.97	1.1250	29.38	1.3272
11.42	1.1294	30.0	1.3335
11.84	1.1339	30.57	1.3396
12.24	1.1383	31.22	1.3456
12.64	1.1423	31.85	1.3521
13.0	1.1474	32.47	1.3585
13.55	1.1520	33.0	1.3642
13.86	1.1566	33.69	1.3714
14.5	1.1631	34.38	1.3780
14.75	1.1662	35.0	1.3856
15.0	1.1697	35.65	1.3913
15.5	1.1755	36.25	1.3961
15.91	1.1803	36.86	1.4040
16.38	1.1852	37.47	1.4116
16.77	1.1901	38.13	1.4187
17.22	1.1950	38.8	1.4267
17.67	1.2000	39.39	1.4328
17.12	1.2050	40.0	1.4410
18.58	1.2101	40.75	1.4472
19.0	1.2148	41.41	1.4545
19.58	1.2202	42.12	1.4619
20.0	1.2250	42.83	1.4694
20.59	1.2308	43.66	1.4770
21.0	1.2361	44.38	1.4845
21.42	1.2414	45.27	1.4922
22.0	1.2462	46.15	1.5000
22.64	1.2522	46.87	1.5079
23.15	1.2576	47.60	1.5156
23.67	1.2632	48.81	1.5236
24.24	1.2687	49.02	1.531

(Hager, Comm. 1883.)

The sp. gr. increases or diminishes for each degree as follows

% NaOH	Corr.
40-50	0.00045
30-39	0.0004
20-29	0.0003
10-19	0.0002

(Hager Comm. 1883.)

Sp. gr. of NaOH + Aq at 15°.

% NaOH	Sp. gr.	% NaOH	Sp. gr.
2.5	1.0280	20	1.2962
5	1.0568	25	1.2833
10	1.1131	30	1.3374
15	1.1790		

(Kohlrausch, W. Ann. 1878.)

r. of NaOH+Aq at 20° containing NaOH to 100 mols. $H_2O = 1.04712$. (Phil. Mag. (5) **16**, 122.)

Sp. gr. of NaOH+Aq at 15°.

	Sp. gr.	% NaOH	Sp. gr.
	1.069	25	1.353
	1.139	30	1.426
	1.210	35	1.500
	1.281		

Adjumenta Varia, Leipsic, **1876**.)

Sp. gr. of NaOH+Aq at 15°.

p. gr.	% NaOH	Sp. gr.	% NaOH	Sp. gr.
99180	17	1.188707	34	1.373453
110611	18	1.199783	35	1.383815
121920	19	1.210861	36	1.394092
133109	20	1.221933	37	1.404279
144317	21	1.233062	38	1.414363
155463	22	1.244119	39	1.424353
166602	23	1.255134	40	1.434299
177733	24	1.266092	41	1.444161
188856	25	1.277063	42	1.453929
199969	26	1.287990	43	1.463623
111069	27	1.298877	44	1.473249
122165	28	1.309708	45	1.482850
133250	29	1.320496	46	1.492406
144353	30	1.331213	47	1.501927
155450	31	1.341879	48	1.511412
166538	32	1.352472	49	1.520868
177619	33	1.362991	50	1.530282

ing, Phil. Mag. 1894, (5) **37**, 373.)

r. of a N solution of NaOH+Aq at $=1.0418$. (Loomis, W. Ann 1896, **60**.)

r. of NaOH+Aq

at 20°/20° 1.0968 3.67 3.82 1.0416 1.0464
ne and Rohland, Z. phys. Ch. 1896, **19**, 272.)

of NaOH+Aq at t°. H_2O at 4°=1.
solutions contained a small amount Na_2CO_3 .

% NaOH	% Na_2CO_3	Sp. gr.
22.57	0.61	1.2312
20.04	0.48	1.2026
17.04	0.35	1.1692
14.16	0.38	1.1374
10.92	0.36	1.1020
22.81	0.55	1.2207
14.01	0.42	1.1232

neider and Walter, M. 1905, **26**, 691.)

Sat. NaOH+Aq boils at 215.5°. (Griffiths.)

Sat. NaOH+Aq boils at 310°. (Gerlach, Z. anal. **26**, 427.)

NaOH+Aq of 1.500 sp. gr. contains 36.8% NaOH and boils at 130°.

B.-pt. of NaOH+Aq containing pts. NaOH to 100 pts. H_2O .

B.-pt.	Pts. NaOH	B.-pt.	Pts. NaOH
105°	17	210°	425.5
110	30	215	475.5
115	41	220	526.3
120	51	225	583.3
125	60.1	230	645.2
130	70.1	235	714.3
135	81.1	240	800
140	93.5	245	888.8
145	106.5	250	1000
150	120.4	255	1142.8
155	134.5	260	1333.3
160	150.8	265	1534
165	168.8	270	1739.1
170	187	275	2000
175	208.3	280	2353
180	230	285	2857
185	254.5	290	3571.4
190	281.7	300	4651.1
195	312.3	305	6451.6
200	345	310	10526.3
205	380.9	314	22222.2

(Gerlach, Z. anal. **26**, 463.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**, 829.)

Sp. gr. of Na_2CO_3 +NaOH+Aq at 11.5°. H_2O at 4°=1.

% Na_2CO_3	% NaOH	Sp. gr.
3.845	14.10	1.196
3.171	13.63	1.182
2.204	12.51	1.164
1.642	10.17	1.136
0.2686	16.64	1.186

(Wegscheider and Walter, M. 1905, **26**, 693.)

Sp. gr. of Na_2CO_3 +NaOH+Aq at t°. H_2O at 4°=1.

t°	% Na_2CO_3	% NaOH	Sp. gr.
60°	15.38	10.63	1.2621
	13.79	9.52	1.2302
	12.10	8.29	1.1952
	9.965	6.86	1.1594
	9.47	6.70	1.1521
	7.69	5.22	1.1158
80°	15.26	11.14	1.2510
	9.48	6.93	1.1417

(Wegscheider and Walter, M. 1905, **26**, 692.)

Solubility of NaOH + Na₂SbS₄ at 30°.

% NaOH	% Na ₂ SbS ₄	Solid phase
0	27.1	Na ₂ SbS ₄ · 9H ₂ O
9.9	13.	"
24.8	5.9	"
32.9	10.5	"
42.6	16.4	"
47.2	17.7	" + NaOH, H ₂ O
49.5	9.1	NaOH, H ₂ O
54.3	0	"

(Donk, Chem. Weekbl. 1908, 5. 529, 629, 767.)

Easily sol. in alcohol or wood spirit; sol. in fusel-oil. Sol. in an aqueous solution of mannite. (Favre, A. ch. (3) 11. 76.)

Easily sol. in glycerine.

Sol. to a certain extent in ether.

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate (Naumann, B. 1904, 37. 3602); benzonitrile. (Naumann, B. 1914, 47. 1370.)

+H₂O. 100 g. solution in H₂O sat. at 25° contain 42 g. Na₂O. (Schreinemakers, Arch. Néer. Sc. 1910, (2) 15. 81.)

Mpt. 64.3°. (See above.)

+1½H₂O. (Cripps, Pharm. J. Trans. (3) 14. 833.)

+2H₂O. See above.

+3½H₂O. Deliquescent. Sol. in H₂O with absorption of much heat. Melts at 6°. (Hermes.)

Mpt. 15.5°. (See above.)

+4, 5 and 7H₂O. (See above.)

The composition of the hydrates formed by NaOH at different dilutions is calculated from determinations of the lowering of the fr.-pt. produced by NaOH and of the conductivity and sp. gr. of NaOH + Aq. (Jones, Am. Ch. J. 1905, 34. 336.)

Sodium perhydroxide, NaO₂H.

"Natrlyl hydroxide."

Decomp. by H₂O. Sol. in cold alcoholic acetic acid. (Tafel, B. 1894, 27. 2300.)

Sodium iodide, NaI, and +2H₂O.

Solubility of NaI and of NaI + 2H₂O in H₂O differ. Below 65°, NaI + 2H₂O usually separates out, and above that temp. NaI separates.

Solubility of NaI in 100 pts. H₂O at t°.

t°	Pts. NaI	t°	Pts. NaI	t°	Pts. NaI
71.3	294.4	92.4	300.2	124.7	317.5
74.1	295.3	97.1	300.3	132.5	317.3
81.6	296.8	101.7	302.5	138.1	319.2
86.4	298.3	110.7	306.2

Solubility is represented by a straight line of the formula $S = 264.19 + 0.3978t$.

Solubility of NaI + 2H₂O in 100 pts. at t°.

t°	Pts. NaI	t°	Pts. NaI	t°	Pts. NaI
-17	149.4	15	173.7	45	215.6
-15	150.3	20	178.7	50	227.8
-5	155.4	25	184.2	55	241.2
0	158.7	30	190.3	60	256.5
5	163.6	35	197.0	65	275.4
10	168.6	40	205.1

(Coppet, A. ch. (5) 50. 424.)

If solubility S = pts. NaI in 100 pts. solution, $S = 61.3 + 0.1712t$ from 0° to 80°; $S = 75 + 0.0258t$ from 80° to 160°. (Étard, C. R. 98. 1432.)

NaI + 2H₂O is sol. in 0.55 pt. H₂O at 15°. (Eder, Dingl. 221. 89.)

100 pts. NaI + Aq at 18–19° contain 62.6 pts. NaI. (v. Hauer, J. pr. 98. 137.)

100 pts. H₂O dissolve at:

0°	20°	40°	60°
158.7	178.6	208.4	256.4 pts. NaI
80°	100°	120°	140°
303	312.5	322.5	333.3 pts. NaI

(Kremers, Pogg. 97. 14.)

Transition pt. for NaI + 2H₂O to NaI is 64.3°, and sat. solution containing 74.4% NaI. (Panfiloff, J. Russ. Phys. Chem. Soc. 1893, 25. 162.)

100 g. H₂O dissolve 172.4 g. NaI at 15°, and sp. gr. of sat. solution = 1.335. (Greenish, Pharm. J. 1900, 65. 190.)

100 g. solution of NaI + 2H₂O sat. at 30° contains 65.5 g. anhyd. NaI. (Corbett, Dissert. 1910.)

Sp. gr. of NaI + Aq at 19.5° containing:

5	10	15	20	25	30 % NaI
1.040	1.082	1.128	1.179	1.234	1.294

35	40	45	50	55	60 % NaI
1.360	1.432	1.510	1.60	1.70	1.81

(Gerlach, Z. anal. 8. 285.)

Sat. solution boils at 141°.

Sol. in liquid SO₂. (Walden, B. 1899, 22. 2864); POCl₃. (Walden, Z. anorg. 1900, 25. 212.)

Very easily sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Sol. in 12.0 pts. absolute alcohol; in 300 pts. ether. (Eder, Dingl. 221. 89.)

Sol. in 3 pts. 90% alcohol. (Hager.)

100 pts. absolute methyl alcohol dissolve 77.7 pts. NaI at 22.5°; ethyl alcohol, 43.1 pts. (de Bruyn, Z. phys. Ch. 10. 783.)

Very sol. in abs. methyl alcohol and is not pptd. therefrom on the addition of a large volume of abs. ether, while wet ether produces immediate separation. (Loeb, J. Am. Chem. Soc. 1905, 27. 1020.)

Solubility of NaI in ethyl alcohol.
NaI in 100 g. alcohol.)

	NaI	t°	NaI
0	43.77	200	42.3
10	44.25	220	38.5
20	44.50	230	36.2
30	45.0	240	32.7
40	45.1	250	26.2
50	45.2	255	21.0
60	45.0	260	10.8
70	44.3	261.5*	8.6

critical temp. of solution.
(Tyrer, Chem. Soc. 1910, 97. 626.)

g. sat. solution of NaI in ethyl alcohol
° contains 30.9 g. (Cocheret, Dissert,

solubility in ethyl alcohol+Aq at 30°.

NaI	% alcohol	Solid phase
52	0	NaI, 2H ₂ O
	3.40	"
2	18.5	"
	18.8	"
8	28.5	"
35	41.7	"
5	53.2	"
91	54.7	"
49	55.37	NaI, 2H ₂ O+NaI
65	59.24	NaI
24	61.78	"
90	68.70	"

(Cocheret, Dissert. 1911.)

room temp. 1 pt. by weight is sol in:
.2 pts. methyl alcohol D¹⁵ 0.7990.
.7 " ethyl " D¹⁵ 0.8100.
.8 " propyl " D¹⁵ 0.8160.

Rohland, Z. anorg. 1898, 18. 325.)

g. methyl alcohol dissolve 90.35 g.
t 25°.
g. ethyl alcohol dissolve 46.02 g. NaI at
g. propyl alcohol dissolve 28.22 g.
t 25°.
g. isoamyl alcohol dissolve 16.30 g.
t 25°.

er and Bissett, Chem. Soc. 1913, 103.
1909.)

Solubility in methyl alcohol+ethyl alcohol
at 25°.

P = % methyl alcohol in the solvent.
G = g. NaI in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution at 25°.

P	G	S 25°/4°
0.00	3.515	1.0806
4.37	3.768	1.1029
10.40	3.971	1.1123
41.02	4.598	1.1742
80.69	5.744	1.2741
84.77	5.892	1.2886
91.25	6.110	1.3056
100.00	6.322	1.3250

(Herz and Kuhn, Z. anorg. 1908, 60. 154.)

Solubility in mixtures of methyl and propyl
alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. NaI in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	6.322	1.3250
11.11	5.845	1.2853
23.8	5.464	1.2528
65.2	4.071	1.138
91.8	2.914	1.0420
93.75	2.649	1.0178
100.00	2.411	0.9968

(Herz and Kuhn, Z. anorg. 1908, 60. 156.)

Solubility in mixtures of propyl and ethy
alcohol at 25°.

P = % propyl alcohol in the solvent.
G = g. NaI in 10 ccm. of the solution.
S = Sp. gr. of the sat. solution.

P	G	S 25°/4°
0	3.515	1.0806
8.1	3.460	1.0732
17.85	3.405	1.0720
56.6	2.841	1.0276
88.6	2.613	1.0130
91.2	2.588	1.0104
95.2	2.474	1.0020
100	2.411	0.9968

(Herz and Kuhn, Z. anorg. 1908, 60. 159.)

Sol. in normal propyl alcohol. (Loeb, J.
Am. Chem. Soc. 1905, 27. 1020.)
28.74 g. are sol. in 100 g. propyl alcohol.
(Schlamp, Z. phys. Ch. 1894, 14. 276.)
Sol. in methyl acetate. (Naumann, B.
1909, 42. 3789.)
Sol. in ethyl acetate. (Casaseca, C. R. 30.
821.)
Insol. in ethyl acetate. (Naumann, B.
1910, 43. 314.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Solubility in organic solvents at t° .

C=pts. by wt. of NaI in 100 ccm. of the sat. solution.

L=no. of liters which at the saturation temp. hold in solution 1 mol. NaI.

Solvent	t°	C	L
Furfurool	25	25.10	0.597
Acetonitrile	25	18.43	0.813
"	0	22.09	0.679
Propionitrile	25	6.230	2.406
"	0	9.091	1.649
Nitromethane	25	0.478	31.36
"	0	0.339	44.27

(Walden, Z. phys. Ch. 1906, 55. 718.)

Very sol. in acetone. (Walden.)

+5H₂O. Transition point of NaI+5H₂O to NaI+2H₂O is -13.5° , and sat. solution contains 60.2% NaI. (Panfiloff, J. Russ. Phys. Chem. Soc. 1893, 25. 162.)

Sodium tin (stannous) iodide, NaI, SnI₂.

Very sol. in H₂O. When treated with little H₂O, NaI is dissolved out, but a larger amt. of H₂O dissolves it completely. (Boullay, A. ch. (2) 34. 375.)

Sodium zinc iodide, 2NaI, ZnI₂+3H₂O.

Deliquescent.

NaZnI₂+2H₂O. Very hygroscopic. (Ephraim, Z. anorg. 1910, 67. 383.)

Sodium nitride,

Decomp. by heat. (Franz Fisher, B. 1910, 43. 1468.)

Sodium suboxide, Na₂O.

Decomp. by H₂O. (de Forcrand, C. R. 1898, 127. 365.)

Sodium oxide, Na₂O.

Very deliquescent, and sol. in H₂O with evolution of heat.

See Sodium hydroxide.

Sodium peroxide, Na₂O₂.

Deliquescent, and very sol. in H₂O with partial decomp.

Solution decomp. on boiling.

Cryst. with 2H₂O, and 8H₂O. (Fairley, Chem. Soc. 1877. 125.)

Forms hydrate Na₂O₂(OH)₄+4H₂O.

Easily sol. in H₂O or dil. acids without decomp. (Schöne, A. 193. 241.)

Sodium peroxide carbonate, Na₂CO₄.

Easily decomp. (Woffenstein, B. 41. 285.)

Sodium peroxide dicarbonate, Na₂C₂O₄.

Easily decomp. (Woffenstein, B. 41. 287.)

Sodium trioxide, NaO₃.

Sol. in H₂O forming a solution of N (Joannis, C. R. 1893, 116. 1371.)

Sodium trioxide carbonate, Na₂CO₃.

(Woffenstein, B. 1908, 41. 296.)

Sodium trioxide dicarbonate, NaHCO₃.

Two isomeric modifications. (W. stein, B. 1908, 41. 390.)

Sodium trioxide hydrate, NaO.OH.

See Sodium perhydroxide.

Isomeric with Tafel's sodyl hydroxide: Na.OH. (B. 27, 2297.)

Insol. in alcohol.

Very unstable. (Woffenstein, B. 41. 290.)

Sodium phosphide, NaP₃.

Easily decomp. by H₂O. (Hugot, C. R. 1895, 121. 208.)

Sodium hydrogen phosphide, NaH₂P.

Decomp. by H₂O. (Joannis, C. R. 119. 558.)

Na₂H₂P₂. Decomp. by acids and (Hugot, C. R., 1898, 126. 1721.)

Sodium selenide, Na₂Se.

Very deliquescent. Decomp. by (Uelsmann, A. 116. 127.)

Insol. in liquid NH₃; sol. in air free to a colorless liquid. (Hugot, C. R. 129. 299.)

Cryst. with 16H₂O, 9H₂O, and 10H₂O. (Fabre, C. R. 102. 613.)

+10H₂O. Very sol. in H₂O; very un- in the air. (Clever, Z. anorg. 1895, 10.

Sodium diselenide, Na₂Se₂.

(Jackson, B. 7. 1277.)

Sodium triselenide, Na₂Se₃.

Sol. in H₂O. (Mathewson, J. Am. C. Soc. 1907, 29. 873.)

Sodium hexaselenide, Na₂Se₆.

Sol. in H₂O. (Mathewson, J. Am. C. Soc. 1907, 29. 873.)

Sodium monosulphide, Na₂S.

Sol. in H₂O. Much less sol. in alcohol in H₂O. Insol. in ether. (Roussin.)

I_2O . Tr. pt. from $\text{Na}_2\text{S} + 5\frac{1}{2}\text{H}_2\text{O}$, 94° .

$\frac{1}{2}\text{H}_2\text{O}$.

sat. solution contains:—

28.48% anhydrous salt at 50°

29.27 " " " 55°

29.92 " " " 60°

31.38 " " " 70°

33.95 " " " 80°

37.20 " " " 90°

le from 48.9 – 91.5° ; stabile from 91.5 –

rano and Fornaini, C. C. 1908, I. 5.)

$\frac{1}{2}\text{O}$. Less efflorescent than with $9\text{H}_2\text{O}$.
 H_2O and alcohol.

sat. solution contains:—

26.7 % anhydrous salt at 50°

28.1 " " " 60°

30.22 " " " 70°

32.95 " " " 80°

36.42 " " " 90°

st. to $\text{Na}_2\text{S} + 5\frac{1}{2}\text{H}_2\text{O}$, 91.5° .

(Parravano and Fornaini.)

I_2O . Efflorescent. Much less sol. in
than H_2O . When dissolved in H_2O ,
sinks from $+22$ to -6.1° . (Finger,
28. 635.)

sat. solution contains:—

3.34% anhydrous Na_2S at -10°

3.36 " " " $+10^\circ$

4.36 " " " 15°

5.30 " " " 18°

6.2 " " " 22°

7.73 " " " 28°

9.09 " " " 32°

9.98 " " " 37°

11.19 " " " 45°

st. to $\text{Na}_2\text{S} + 5\frac{1}{2}\text{H}_2\text{O}$, 48.9° .

(Parravano and Fornaini.)

1 disulphide, Na_2S_2 .

in H_2O and alcohol.

I_2O . Not efflorescent.

1 trisulphide, Na_2S_3 .

in H_2O with decomp.

st. with $3\text{H}_2\text{O}$ from an alcoholic solu-
(Böttger, A. 223. 355.)

1 tetrasulphide, $\text{Na}_2\text{S}_4 + 6\text{H}_2\text{O}$.

deliquescent, and sol. in H_2O . Diffi-
sol. in absolute alcohol. Insol. in ether.
e.)

I_2O . Efflorescent. (Böttger.)

1 pentasulphide, $\text{Na}_2\text{S}_5 + 6\text{H}_2\text{O}$.

in H_2O . (Schöne.)

in alcohol.

I_2O . (Böttger.)

tion is easily decomp. by warming.
Chem. Soc. 37. 461.)

Sodium tellurium sulphide.

See Sulphotellurate, sodium.

Sodium stannic sulphide.

See Sulphostannate, sodium.

Sodium yttrium sulphide, Na_2S , Y_2S_3 .

Decomp. by dil. acids, even by $\text{HC}_2\text{H}_3\text{O}_2 +$
Aq. (Duboin, C. R. 107. 243.)

Sodium zinc sulphide, Na_2S , 3ZnS .

Not so stable as the corresponding K salt.
(Schneider, J. pr. (2) 8. 29.)

Sodium sulphoselenide, $\text{Na}_2\text{SSe}_2 + 5\text{H}_2\text{O}$.

Hydroscopic, and decomp. in the air. (Mes-
singer, B. 1897, 30. 806.)

Sodium telluride, Na_2Te .

Sol. in H_2O . (Demarçay, Bull. Soc. (2)
40. 99.)

Sodium tritelluride, Na_2Te_3 .

Sol. in H_2O .

Sol. in liquid NH_3 . (Hugot, C. C. 1899,
II. 580.)

Stannic acid, H_2SnO_3 .

Insol. in H_2O . Sol. in HCl , and $\text{H}_2\text{SO}_4 +$
Aq, even when dil. (Fremy.) Easily sol. in
acids, from which solution it may be pptd. by
dilution or boiling. While moist it is sol. in
 $\text{HNO}_3 + \text{Aq}$, but gradually separates on stand-
ing, and coagulates at once when heated to
 50° . If NH_4NO_3 be added to the solution, it
remains clear at ord. temp. (Berzelius.)

Easily sol. in $\text{HNO}_3 + \text{Aq}$, when previously
treated with $\text{NH}_4\text{OH} + \text{Aq}$. (Thénard.)

Easily sol. in $\text{KOH} + \text{Aq}$, but addition of
large excess ppts. K_2SnO_3 , insol. in $\text{KOH} +$
Aq.

Easily sol. in $\text{NaOH} + \text{Aq}$, and not pptd. by
an excess of that reagent. (Barfoed, J. B.
1867. 267.)

Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$ or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$.

Completely sol. in $\text{K}_2\text{CO}_3 + \text{Aq}$, but not in
 $\text{Na}_2\text{CO}_3 + \text{Aq}$.

Insol. in alkali hydrogen carbonates or
 $\text{NH}_4\text{Cl} + \text{Aq}$.

Sol. in alkali sulphides + Aq. (Berzelius.)

Sol. in triethyltoluanyl ammonium hy-
drate + Aq.

Not pptd. by $\text{NH}_4\text{OH} + \text{Aq}$ in presence of
Na citrate + Aq.

$\text{SnO}_2, 2\text{H}_2\text{O}$. (Weber, Pogg. 122. 358.)

"*α-Orthostannic acid*." Easily sol. in
 $\text{HCl} + \text{Aq}$. (Neumann, M. 12. 515.)

$\text{H}_{10}\text{Sn}_5\text{O}_{15}$ (?)

Metastannic acid. Insol. in H_2O , HNO_3 ,
or $\text{H}_2\text{SO}_4 + \text{Aq}$. Insol. in $\text{HCl} + \text{Aq}$, but
converted thereby into metastannic chloride,
which dissolves after excess of HCl has been
removed. (Fresenius.) Insol. in $\text{HCl} + \text{Aq}$ of
sp. gr. 1.1. (Barfoed.) Sol. in large amount

of conc. $\text{HCl} + \text{Aq.}$ (Allen, Chem. Soc. (2) 10. 274.)

In contact with $\text{HCl} + \text{Aq.}$ metastannic acid is converted into stannic acid. (Barfoed.)

Insol. in $\text{HNO}_3 + \text{Aq.}$ even after treatment with $\text{NH}_4\text{OH} + \text{Aq.}$

Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$

Sol. in KOH or $\text{NaOH} + \text{Aq.}$ with formation of metastannates, which are insol. in dil. $\text{NaOH} + \text{Aq.}$ but sol. in H_2O or $\text{KOH} + \text{Aq.}$ therefore $\text{KOH} + \text{Aq.}$ dissolves metastannic acid, while $\text{NaOH} + \text{Aq.}$ does not, but if the clear solution in $\text{KOH} + \text{Aq.}$ is treated with a large excess of that reagent, a further pptn. occurs. (Barfoed, J. pr. 101. 368.)

Insol. in $\text{K}_2\text{CO}_3 + \text{Aq.}$ (Rose); alkali carbonates $+ \text{Aq.}$ (Fremy.)

Insol. in $\text{NH}_4\text{Cl} + \text{Aq.}$ even after long boiling.

Sol. in $\text{Fe}(\text{NO}_3)_3 + \text{Aq.}$ containing HNO_3 . (Lepèz and Storch, W. A. B. 98, 2b. 270.)

Also in $\text{Cr}(\text{NO}_3)_3 + \text{Aq.}$ but not in $\text{Ce}(\text{NO}_3)_3$, $\text{Al}(\text{NO}_3)_3$, $\text{Co}(\text{NO}_3)_2 + \text{Aq.}$ etc. (L. and S.)

A colloidal metastannic acid sol. in H_2O can be obtained. (Lepèz and Storch.)

According to Weber (Pogg. 122. 358), stannic and metastannic acids are only different hydrates of same oxide, and it is not a case of allotropic modification.

Colloidal. H_2SnO_3 in colloidal state can be obtained in aqueous solution containing 5.164 g. SnO_2 in a litre. This solution is coagulated by $\text{HNO}_3 + \text{Aq.}$ only when in great excess; easily by dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ but not by conc. $\text{HCl} + \text{Aq.}$ $\text{NH}_4\text{OH} + \text{Aq.}$ in large excess causes coagulation; also NH_4Cl , NaOH , NaCl , Na_2SO_4 , etc. (Schneider, Z. anorg. 5. 83.)

Parastannic acid, $\text{H}_2\text{Sn}_2\text{O}_{11} + 3\text{H}_2\text{O}$.

(Engel, C. R. 1897, 125. 711.)

Stannates.

Stannates of alkali metals are sol. in H_2O ; others are insol. All metastannates, excepting Na , K , and NH_4 salts, are insol. in H_2O . (Fremy, A. ch. (3) 12. 474.)

Ammonium stannate, $(\text{NH}_4)_2\text{O}$, 2SnO_2 .

Sol. in H_2O . Insol. in dil. $\text{NH}_4\text{OH} + \text{Aq.}$ (Berzelius.)

$+ x\text{H}_2\text{O}$. (Moberg, 1838.)

Ammonium cupric stannate, $(\text{NH}_4)_2\text{O}$, $\text{CuSnO}_3 + 2\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in acids. (Ditte, C. R. 96. 701.)

Barium stannate, $\text{BaSnO}_3 + 6\text{H}_2\text{O}$.

Ppt. Sol. in $\text{HCl} + \text{Aq.}$ (Moberg.)

$\text{Ba}_2\text{SnO}_4 + 10\text{H}_2\text{O}$. Insol. in H_2O . Sol. in acids. (Ditte, C. R. 95. 641.)

Calcium stannate, $\text{CaSnO}_3 + 4\text{H}_2\text{O}$.

Ppt. (Moberg.)

$+ 5\text{H}_2\text{O}$. Insol. in H_2O . Sol. in acids. (Ditte, C. R. 98. 701.)

2CaO , SnO_2 . (Zulkowski, Chem. 1901, 24. 422.)

Cobaltous stannate, $\text{CoSnO}_3 + 6\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in acids. (Ditte.)

Cupric stannate, $\text{CuSnO}_3 + 3\text{H}_2\text{O}$.

(Moberg.)

$+ 4\text{H}_2\text{O}$. Insol. in H_2O . (Ditte.)

Cuprous stannous stannate, Cu_2O , $3\text{SnO}_2 + 5\text{H}_2\text{O}$.

Slowly decomp. by dil. acids, and $\text{NH}_4\text{OH} + \text{Aq.}$; completely decomp. by conc. acids. (Lenssen, J. pr. 79. 90.)

Gold (aurous) stannate.

See Gold purple.

Lead stannate, $\text{RbSn}(\text{OH})_6$.

Ppt. (Bellucci, Chem. Soc. 1905, (2) 40.)

Lithium stannate hexatungstate, $2\text{Li}_2\text{O}$, 8SnO_2 , $6\text{WO}_3 = \text{Li}_2\text{SnO}_3$, $\text{Li}_2\text{W}_6\text{O}_{18}$.

Insol. in H_2O . (Knorre, J. pr. (2) 49.)

Magnesium stannate.

Ppt. (Moberg.)

Manganous stannate.

Ppt. (Moberg.)

Mercurous stannate, $\text{Hg}_2\text{SnO}_3 + 5\text{H}_2\text{O}$.

Ppt.

Mercuric stannate, $\text{HgSnO}_3 + 6\text{H}_2\text{O}$.

Ppt. (Moberg, J. pr. 28. 231.)

Nickel stannate, $\text{NiSnO}_3 + 5\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in acids. (Ditte, C. R. 96. 701.)

Platinous sodium stannous stannate, 2P , Na_2O , SnO , $\text{SnO}_2(?)$.

(Schneider, Pogg. 136. 105.)

Platinous stannous stannate, PtO , 2SnO_2 .

Decomp. by conc. alkalis. (Schneider, Pogg. 136. 105.)

Potassium stannate, $\text{K}_2\text{SnO}_3 + 3\text{H}_2\text{O}$.

100 pts. H_2O dissolve 106.6 pts. at 100°; solution has sp. gr. = 1.618; 100 pts. dissolved at 20°, solution has sp. gr. = 1.55. (Ordway, Sill. Am. J. (2) 40. 173.)

sol. in conc. KOH + Aq.
 1 KCl + Aq. (Freymy.)
 1 alcohol.
 From aqueous solution by the addition of any soluble salt, especially those of NH₄ (Freymy); by NH₄Cl, but not by NaCl (Ordway).
 Insol. in acetone. (Naumann, B. 1904, 37.)

1 *metastannate*, K₂O, 10SnO₂,
 SnO₂ + 3H₂O. Sol. in H₂O. Solution becomes turbid on heating. (Rose.)
 3SnO₂ + 5H₂O. Sol. in H₂O, but loses water by drying. (Freymy, A. ch. (3) 23.)

3SnO₂ + 4H₂O. Completely sol. in H₂O. Sol. in alcohol. (Freymy, A. ch. (3) 23.)

3SnO₂ + 3H₂O. Deliquescent.

stannate, Ag₂SnO₃.
 Insol. in H₂O. Unacted upon by NH₄OH + Aq. (Ditte.)

argentous), *stannous stannate* (?),
 SnO, 3SnO₂ + 3H₂O (?).
 Insol. in H₂O. Dil. HNO₃ + Aq slowly dissolves all SnO₂; HNO₃ + Aq rapidly.
 Insol. in boiling conc. H₂SO₄. (Schulze, Z. anorg. 7. 257.)

stannate, Na₂SnO₃ + 3H₂O.
 Easily sol. in cold than in hot H₂O.

2 pts. H₂O at 20° and 100°. (Marshall, Z. anorg. 3. H₂O dissolve 67.4 pts. at 0°, 61.3 at 15°, and solutions have sp. gr. = 1.4728 at 15.5°. (Ordway, Sill. Am. J. 73.)
 From Na₂SnO₃ + Aq by salts of K, Na, or Ag.
 Insol. in acetone. (Naumann, B. 1904, 37.)

1. (Prandtl, B. 1907, 40. 2129.)
 2. (Haefely, J. B. 1857. 650.)
 3. (Jones, C. C. 1865. 607.)
 4. O. Very efflorescent. (Scheurer-Bull. Soc. (2) 8. 389.)

metastannate, Na₂O, 9SnO₂ + 3H₂O.
 1. H₂O. Insol. in NaOH + Aq or in KOH + Aq. (Barfoed, J. B. 1867. 267.)
 2. 5SnO₂. Very difficultly sol. in H₂O. (A. ch. (3) 23. 399.)
 3. in KOH + Aq.
 4. (Haefely, Chem. Gaz. 1855. 59.)

stannate vanadate,
 3SnO₂, 3Na₂VO₄ + 32H₂O.
 3SnO₂, 4Na₂VO₄ + 48H₂O.
 3SnO₂, 5Na₂VO₄ + 64H₂O.
 3SnO₂, 6Na₂VO₄ + 80H₂O. (Prandtl, B. 1907, 40. 2128.)

Strontium stannate, 3SrO, 2SnO₂ + 10H₂O.

Ppt. Insol. in H₂O. Sol. in acids. (Ditte, C. R. 95. 641.)

SrSn(OH)₆. (Belluci, Chem. Soc. 1905, 88 (2) 40.)

Tin (stannous) stannate, SnO, 6SnO₂ + 5H₂O.

Insol. in H₂O. Decomp. by HNO₃ + Aq into metastannic acid. (Schiff, A. 120. 53.)

Sol. in HCl + Aq, and in KOH + Aq.

Tin (stannous) metastannate, SnO, 7SnO₂.

SnO, 6SnO₂ + 9H₂O. Sol. in KOH + Aq or in HCl + Aq. (Freymy.)
 + 4H₂O. (Schiff.)

Zinc stannate, ZnSnO₃ + 2H₂O.

Ppt. (Moberg, 1838.)

3ZnO, 2SnO₂ + 10H₂O. Insol. in H₂O. Sol. in acids. (Ditte.)

Perstannic acid, H₂Sn₂O₇.

See Perstannic acid.

Stannophosphomolybdic acid.

Ammonium stannophosphomolybdate,

3(NH₄)₂O, 4SnO₂, 3P₂O₅, 16MoO₃ + 28H₂O.

Quite insol. even in boiling H₂O. (Gibbs, Am. Ch. J. 7. 392.)

Stannophosphotungstic acid.

Ammonium stannophosphotungstate,

2(NH₄)₂O, 2SnO₂, P₂O₅, 22WO₃ + 15H₂O.

Precipitate. Sl. sol. in boiling H₂O. (Gibbs, Am. Ch. J. 7. 319.)

Stannosulphuric acid.

See Sulphate, stannic.

Stibine.

See Hydrogen antimonide.

Strontium, Sr.

Decomp. by H₂O with violence. Dil. H₂SO₄, and HCl + Aq decomp. and dissolve; cold H₂SO₄ attacks slowly. Fuming HNO₃ has scarcely any action even when boiling. (Franz, J. pr. 107. 253.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 829.)

Sol. in excess of liquid NH₃ at -60° forming Sr(NH₂)₂. (Roederer, C. R. 1905, 140. 1252.)

Strontium amalgam, SrHg₁₂.

Stable below 30°. Above 30° the composition of the amalgam varies. Can be cryst. from Hg at any temp. below 30°. (Kerp, Z. anorg. 1900, 25. 68.)

Strontium amide, Sr(NH₂)₂.

(Roederer, Bull. Soc. 1906, (3) 35. 715.)

Strontium arsenide, Sr_3As_2 .
Decomp. by H_2O . (Lebeau, C. R. 1899, 129. 47.)

Strontium azoimide, SrN_6 .
Hydroscopic.
45.83 pts. are sol. in 100 pts. H_2O at 16° .
0.095 " " " " 100 " abs. alcohol at 16° .
Insol. in pure ether. (Curtius, J. pr. 1898, (2) 58. 287.)

Strontium boride, SrB_4 .
Sol. in fused oxidizing agents; not decomp. by H_2O ; insol. in aq. acids; sl. sol. in conc. H_2SO_4 ; sol. in dil. and conc. HNO_3 . (Moissan, C. R. 1897, 125. 633.)

Strontium bromide, SrBr_2 , and $+6\text{H}_2\text{O}$.
100 pts. H_2O dissolve at:

0°	20°	38°	59°	83°	110°
87.7	99	112	133	182	250 pts. SrBr_2 .

(Kremers, Pogg. 103. 65.)
Sat. $\text{SrBr}_2 + \text{Aq}$ contains at:

-11°	-1°	$+7^\circ$	18°
43.1	46.85	48.2	51.7% SrBr_2 ,
20°	93°	97°	107°
51.8	68.5	68.7	69.8% SrBr_2 .

(Étard, A. ch. 1894, (7) 2. 540.)
Sp. gr. of $\text{SrBr}_2 + 19.5^\circ$ containing:

5	10	15	20	25 % SrBr_2
1.046	1.094	1.146	1.204	1.266
30	35	40	45	50 % SrBr_2 .
1.332	1.41	1.492	1.59	1.694

(Kremers, Pogg. 99. 444; calculated by Gerlach, Z. anal. 8. 285.)
Somewhat sol. in absolute alcohol. (Löwig.)
Solubility of anhydrous SrBr_2 in alcohol is practically constant between 0° and 40° , 100 ccm. of abs. alcohol dissolving about 64.5 g. of the anhydrous salt and forming a solution having a sp. gr. = 1.210 at 0° . (Fonzes-Diacon, Chem. Soc. 1895, 68 (2) 223.)
Much more sol. than BaBr_2 in boiling amyl alcohol.
Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)
Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Strontium stannic bromide.
See Bromostannate, strontium.

Strontium bromide ammonia, $2\text{SrBr}_2, \text{NH}_3$.
Sol. in H_2O . (Rammelsberg, Pogg. 55. 238.)

Strontium bromide hydrazine, $\text{SrBr}_2, 3\text{N}_2\text{H}_4$.
Very sol. in H_2O . (Franzen, Z. anorg. 1908, 60. 290.)

Strontium bromofluoride, $\text{SrF}_2, \text{SrBr}_2$.
Decomp. by H_2O . (Defacqz, A. ch. 1904, (8) 1. 356.)

Strontium carbide, SrC_2 .
Easily decomp. by H_2O and dil. acids. (Moissan, Bull. Soc. 1894, (3) 11. 1008.)

Strontium carbonyl, $\text{Sr}(\text{CO})_2$.
(Roederer, Bull. Soc. 1906, (3) 25. 725.)

Strontium chloride, SrCl_2 , and $+6\text{H}_2\text{O}$.
Deliquescent in moist air.
Sol. in 1.5 pts. H_2O at 15° . and 0.8 pt. at boiling (Dumas); in 1.996 pts. H_2O at 15° (Gerlach).
1 pt. anhydrous SrCl_2 is sol. in 2.27 pts. H_2O at 0° in 1.88 pts. at 20° ; in 1.54 pts. at 40° ; in 1.15 pts. at 60° . in 1.08 pts. at 80° ; in 0.98 pt. at 100° . (Kremers, Pogg. 103. 66.)
100 pts. H_2O dissolve 106.2 pts. $\text{SrCl}_2 + 6\text{H}_2\text{O}$ at 0° , and 205.8 pts. at 40° . (Tilden, Chem. Soc. 45. 409.)

Solubility in 100 pts. H_2O at t°

t°	Pts. SrCl_2	t°	Pts. SrCl_2	t°	Pts. SrCl_2
0	44.2	41	67.4	81	92.7
1	44.5	42	68.2	82	93.1
2	44.8	43	68.9	83	93.4
3	45.2	44	69.7	84	93.7
4	45.6	45	70.4	85	94.1
5	46.0	46	71.2	86	94.5
6	46.5	47	72.0	87	94.9
7	46.9	48	72.8	88	95.4
8	47.4	49	73.6	89	95.8
9	47.8	50	74.4	90	96.2
10	48.3	51	75.3	91	96.7
11	48.8	52	76.1	92	97.2
12	49.4	53	77.0	93	97.9
13	49.9	54	77.9	94	98.2
14	50.4	55	78.7	95	98.5
15	51.0	56	79.6	96	99.4
16	51.5	57	80.4	97	100.0
17	52.1	58	81.3	98	101.6
18	52.7	59	82.2	99	101.3
19	53.3	60	83.1	100	101.9
20	53.9	61	84.0	101	102.6
21	54.5	62	84.9	102	103.3
22	55.1	63	85.8	103	104.0
23	55.7	64	86.6	104	104.7
24	56.3	65	87.5	105	105.4
25	56.9	66	88.4	106	106.1
26	57.5	66.5	88.8	107	106.9
27	58.1	67	88.9	108	107.6
28	58.7	68	89.1	109	108.4
29	59.3	69	89.3	110	109.1
30	60.0	70	89.6	111	109.9
31	60.6	71	89.8	112	110.7
32	61.3	72	90.1	113	111.4
33	61.9	73	90.3	114	112.2
34	62.5	74	90.6	115	113.0
35	63.2	75	90.9	116	113.8
36	63.9	76	91.2	117	114.6
37	64.6	77	91.5	118	115.5
38	65.3	78	91.8	118.8	116.4
39	66.0	79	92.1	...	
40	66.7	80	92.4	...	

(Mulder, Scheik. Verhandel. 1864. 118.)

100 pts. H₂O dissolve 52.4 pts. SrCl₂ at 18°. (Gerardin.)

Sat. SrCl₂+Aq contains % SrCl₂ at t°.

t°	% SrCl ₂	t°	% SrCl ₂
- 17	26.5	75	46.5
- 11	28.6	80	47.1
- 5	29.3	92	47.5
- 1	30.8	98	49.6
+ 2	31.3	104	50.7
7	31.7	105	50.7
18	33.7	118	52.0
21.5	34.7	132	52.5
35	37.8	144	54.7
44.5	39.8	153	55.7
54	42.8	175	60.5
55	43.8	215	64.1
59	47.7	222	65.4
64	46.4	250	67.3
70	46.1

(Étard, A. ch. 1894, (7) 2. 535.)

SrCl₂+Aq sat. at 8° has sp. gr. =1.379. (Anthon, A. 211.)

Sp. gr. of SrCl₂+Aq.

Pts. SrCl ₂ to 100 pts. H ₂ O	Sp. gr.	Pts. SrCl ₂ to 100 pts. H ₂ O	Sp. gr.
9.81	1.0823	41.04	1.3114
20.12	1.1632	51.69	1.3816
30.57	1.2401

(Kremers, Pogg. 99. 444.)

Sp. gr. of SrCl₂+Aq at 15°.

% SrCl ₂	Sp. gr.	% SrCl ₂	Sp. gr.
5	1.0453	25	1.2580
10	1.0929	30	1.3220
15	1.1439	33	1.3633
20	1.1989

(Gerlach, Z. anal. 8. 283.)

Sp. gr. of SrCl₂+Aq at 24.7°. a=no. of molecules in grms. dissolved in 1,000 g. H₂O; b=sp. gr. when a=SrCl₂+6H₂O, ½ mol. SrCl₂+6H₂O=133.5 g.; c=sp. gr. when a=SrCl₂, ½ mol.=79.5 g.

a	b	c	a	b	c
1	1.063	1.067	7	1.304	1.401
2	1.118	1.130	8	1.330	...
3	1.166	1.190	9	1.354	...
4	1.207	1.247	10	1.376	...
5	1.243	1.301	11	1.396	...
6	1.275	1.352

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of SrCl₂+Aq at 18°.

%SrCl ₂	Sp. gr.	% SrCl ₂	Sp. gr.
5	1.0443	20	1.2023
10	1.0932	22	1.2259
15	1.1456

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of SrCl₂+Aq at 0°. S=pts. SrCl₂ in 100 pts. solution.

S	Sp. gr.	S	Sp. gr.
31.8193	1.3609	18.2629	1.1915
27.7170	1.3086	12.9997	1.1284
23.2300	1.2515	6.7243	1.0637

(Charpy, A. ch. (6) 29. 24.)

Sat. SrCl₂+Aq boils at 114° (Kremers); 118.8° (Mulder); 117.45°, and contains 117.5 pts. SrCl₂ to 100 pts. H₂O (Legrand); forms a crust at 115.5°, and contains 120.7 pts. SrCl₂ to 100 pts. H₂O; highest temp. observed, 119°. (Gerlach, Z. anal. 26. 436.)

B.-pt. of SrCl₂+Aq containing pts. SrCl₂ to 100 pts. H₂O. G=according to Gerlach (Z. anal. 26. 442); L=according to Legrand (A. ch. (2) 59. 436.)

B.-pt.	G	L	B.-pt.	G	L
101°	11	16.7	110°	71.4	68.9
102	20.5	25.2	111	76.5	74.1
103	28.9	32.1	112	81.6	79.6
104	36.2	37.9	113	87	85.3
105	43.2	43.4	114	93.1	91.2
106	49.6	48.8	115	99.5	97.5
107	55.4	54.0	116	105.9	104.0
108	60.8	59.0	117	112.3	110.9
109	66.2	63.9	117.5

Melts in its crystal H₂O at 112°. (Tilden, Chem. Soc. 45. 409.)

Sp. gr. of SrCl₂+Aq at 25°.

Concentration of SrCl ₂ +Aq.	Sp. gr.
1-normal	1.0676
1/2- " "	1.0336
1/4- " "	1.0171
1/8- " "	1.0084

(Wagner, Z. phys. Ch. 1890, 5. 40.)

SrCl₂+Aq containing 3.24% SrCl₂ has sp. gr. 20°/20°=1.0284.

SrCl₂+Aq containing 7.08% SrCl₂ has sp. gr. 20°/20°=1.0638.

(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 279.)

Sp. gr. of SrCl ₂ +Aq at 20°.	
g. mols. SrCl ₂ per l.	Sp. gr.
0.01	1.0012284
0.02937	1.0038396
0.03987	1.0053832
0.05017	1.007028
0.07077	1.009560
0.10	1.013205
0.25	1.034433
0.50	1.068379
0.75	1.101760
1.00	1.135423

(Jones and Pearce, Am. Ch. J. 1907, 38. 697.)

Conc. HCl+Aq ppts. part of the SrCl₂ from SrCl₂+Aq. (Hope.)

Solubility of SrCl₂ in HCl+Aq at 0°. SrCl₂=
½ mols. SrCl₂ (in milligrammes) dis-
solved in 10 ccm. of liquid; HCl=mols.
HCl (in milligrammes) dissolved in
10 ccm. of liquid.

SrCl ₂	HCl	Sum of mols.	Sp. gr.
55	0	55.0	1.334
48.2	6.1	54.3	1.3045
41.25	12.75	54.00	1.2695
30.6	23.3	53.9	1.220

(Engel, Bull. Soc. (2) 45. 655.)

Solubility of SrCl₂ in HCl+Aq at 0°.

Mg. mols. per 10 cc. solution		Sp. gr. of solution	G. per 100 cc. solution	
$\frac{\text{SrCl}_2}{2}$	HCl		SrCl ₂	HCl
51.6	0	1.334	40.9	0.0
44.8	6.1	1.304	35.5	2.22
37.85	12.75	1.269	30.0	4.65
27.2	23.3	1.220	21.56	8.49
22.0	28.38	1.201	17.44	10.35
14.0	37.25	1.167	11.09	13.58
4.25	52.75	1.133	3.37	19.23

(Engel, A. ch. 1888, (6) 13. 376.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Sol. in 6 pts. alcohol of 0.833 sp. gr. at 15°. (Vau-
quelin.)

Sol. in 24 pts. absolute alcohol at 15° and in 19 pts. at
boiling. (Bucholz.) Sol. in 2.5 pts. of boiling alcohol.

Anhydrous SrCl₂ is sol. in 111.6–116.4 pts.
alcohol of 99.3% at 14.5°, and in 26.2 pts. of
the same alcohol at boiling. (Fresenius, A.
59. 127.)

100 pts. alcohol of given sp. gr. at 0° dis-
solve pts. SrCl₂ at 18°.

0.990	0.985	0.973	0.966	0.953 sp. gr.
49.81	47.0	39.6	35.9	30.4 pts. SrCl ₂ ,
0.939	0.909	0.846	0.832 sp. gr.	
26.8	19.2	4.9	3.2 pts. SrCl ₂ .	

Insol. in absolute alcohol. (Gerardin, A.
ch. (4) 5. 156.)

100 pts. absolute methyl alcohol dissolve
63.3 pts. SrCl₂+6H₂O at 6°; ethyl alcohol,
3.8 pts. (de Bruyn, Z. phys. Ch. 18. 787.)

Sl. sol. in boiling amyl alcohol. (Browning,
Sill. Am. J. 144. 459.)

100 g. 95% formic acid dissolve 23.8 g. SrCl₂
at room temp. (Aschan, Ch. Ztg. 1912, 37.
1117.)

Absolutely insol. in acetic ether. (Com.
C. R. 102. 363.)

Very sl. sol. in acetone. (Krug and
M'Elroy.)

Sol. in acetone. (Eidmann, C. C. 188,
II. 1014.)

Insol. in methyl acetate. (Naumann, B.
1909, 42. 3790.)

Insol. in benzonitrile. (Naumann, B. 1914
47. 1370.)

Sl. sol. in anhydrous pyridine. Sol. in
97%, 95% and 93% pyridine+Aq. (Kalle-
berg, J. Am. Chem. Soc. 1908, 30. 1107.)

+2H₂O. Tr. pt. from +6H₂O is 615°.
(Richards and Churchill, Z. phys. Ch. 188,
28. 313.)

+6H₂O. See above.

Strontium thallic chloride, SrCl₃, 2TiCl₃-
6H₂O.

(Gewecke, A. 1909, 366. 223.)

Strontium tin (stannous) chloride, SrCl₂
SnCl₂+4H₂O.

Sol. in H₂O. (Poggiale, C. R. 20. 1183.)

Strontium tin (stannic) chloride.

See Chlorostannate, strontium.

Strontium uranium chloride, SrCl₂, UCl₄

Decomp. by H₂O. (Aloy, Bull. Soc. 1909,
(3) 21. 265.)

Strontium zinc chloride, SrZnCl₄+4H₂O.

Very sol. in H₂O. (Ephraim, Z. anorg.
1910, 67. 380.)

Strontium chloride ammonia, SrCl₂, 8NH₃

Decomp. by H₂O. (Rose, Pogg. 29. 153.)

Strontium chloride hydrazine, SrCl₂, 2N₂H₄
+H₂O.

Hydroscopic. (Franzen, Z. anorg. 1906,
60. 289.)

Strontium chloride hydroxylamine, 2SrCl₂.
5NH₂OH+2H₂O.

As Ca comp. (Antonow, J. Russ. Phys.
Chem. Soc. 1905, 37. 482.)

Strontium hydrogen chloride hydroxylamine,
2SrCl₂, 3HCl, 9NH₂OH+H₂O.

(Antonow, J. Russ. Phys. Chem. Soc. 1905,
37. 482.)

Strontium chlorofluoride, SrF₂, SrCl₂.

Decomp. by H₂O, by very dil. HCl, HNO₃,
or acetic acid, by hot dil. or conc. H₂SO₄.

ol. in conc. HCl or HNO₃. Insol. in, and
ot decomp. by cold or boiling alcohol.
Defacqz, A. ch. 1904, (8) 1. 355.)

strontium fluoride, SrF₂.

Somewhat sol. in H₂O. (Fr. Röder.)
1 l. H₂O dissolves 113.5 mg. SrF₂ at 0.26°;
17.3 mg. at 17.4°; 119.3 mg. at 27.4°.
Kohlrausch, Z. phys. Ch. 1908, 64. 168.)
Insol in HF + Aq. (Berzelius.)
Boiling HCl + Aq dissolves; sl. attacked by
oiling HNO₃ + Aq; decomp. by hot H₂SO₄.
Poulenc, C. R. 116. 987.)

strontium stannic fluoride.

See Fluostannate, strontium.

strontium titanium fluoride.

See Fluotitanate, strontium.

strontium fluoiodide, SrF₂, Srl₂.

Decomp. by cold H₂O, more rapidly by hot
I₂O. Decomp. by dil. HCl, dil. HNO₃, dil.
I₂SO₄ or conc. H₂SO₄, also by alcohol and by
ether, if not absolute. (Defacqz, A. ch. 1904,
8) 1. 358.)

strontium hydride, SrH.

Decomp. by H₂O or HCl + Aq. (Winkler,
3. 24. 1976.)
SrH₂. Decomp. by H₂O. (Gautier, C. R.
.902, 134. 100.)

strontium hydroselenide.

Sol. in H₂O.

strontium hydrosulphide, SrS₂H₂.

Sol. in H₂O; decomp. by boiling.

strontium hydroxide, SrO₂H₂, and +8H₂O.

Deliquescent.
Sol. in 50 pts. cold, and 2.4 pts. boiling H₂O (Bu-
hols); in 50 pts. H₂O at 15.56° (Dalton); in 51.4 pts.
I₂O at 15.56°, and 2 pts. at 100° (Hope); in 52 pts. H₂O
t 15°, and 2.4 pts. at 100° (Berzelius); in 48 pts. H₂O
t 18.75° (Abl).
100 pts. H₂O at 20° dissolve 1.49 pts. SrO. (Bineau,
. R. 41. 509.)
00 pts. aqueous solution of SrO₂H₂ contain
pts. SrO and pts. SrO₂H₂ + 8H₂O at t°.

t°	Pts. SrO	Pts. SrO ₂ H ₂ +8H ₂ O	t°	Pts. SrO	Pts. SrO ₂ H ₂ +8H ₂ O
0	0.35	0.90	55	2.54	6.52
5	0.41	1.05	60	3.03	7.77
10	0.48	1.23	65	3.62	9.29
15	0.57	1.46	70	4.35	11.16
20	0.68	1.74	75	5.30	13.60
25	0.82	2.10	80	6.56	16.83
30	1.00	2.57	85	9.00	23.09
35	1.22	3.13	90	12.00	30.78
40	1.48	3.80	95	15.15	38.86
45	1.78	4.57	100	18.60	47.71
50	2.13	5.46

Scheibler, J. pharm. Chim. 1883, (5) 8. 540.)
Sol. in cold NH₄Cl + Aq. (Rose.)

Solubility in Sr(NO₃)₂ + Aq at 25°.

Sp. gr. 25°/25°	G. SrO as Sr(OH) ₂ in 100 g. H ₂ O	G. Sr(NO ₃) ₂ in 100 g. H ₂ O
1.481	0.0	79.27
*1.506	1.76	81.06
1.490	1.71	74.27
1.450	1.55	66.88
1.419	1.51	63.71
1.403	1.47	60.37
1.381	1.41	56.30
1.359	1.34	52.90
1.327	1.27	46.97
1.317	1.20	44.03
1.291	1.14	40.83
1.267	1.11	37.81
1.239	1.03	32.41
1.217	1.01	28.80
1.206	0.96	26.58
1.178	0.95	23.83
1.148	0.91	17.96
1.126	0.87	16.21
1.108	0.84	12.78
1.079	0.81	8.96
1.059	0.79	6.29
1.033	0.78	4.45

*Solution is sat. with respect to both sub-
stances.
(Parsons and Perkins, J. Am. Chem. Soc.
1910, 32. 1388.)

Sol. in methyl alcohol. At room temp. 1
l. contains 31.5 g. SrO. (Neuberg and Re-
wald, Biochem. Z. 1908, 9. 540.)
Insol. in acetone. (Eidmann, C. C. 1899,
II. 1014.)
Sol. in an aqueous solution of cane sugar.
(Hunton, Phil. Mag. (3) 11. 156.)

Solubility in H₂O containing 10 g. sugar at t°.

t°	g. SrO ₂ H ₂ + 8H ₂ O	t°	g. SrO ₂ H ₂ + 8H ₂ O
3	3.10	24	4.79
15	3.79	40	9.70

(Sidersky, C. C. 1886. 57.)
+8H₂O. 0.0835 mol. is sol. in 1 l. H₂O at
25°. (Rothmund, Z. phys. Ch. 1909, 69. 539.)

Solubility in organic compds. + Aq at 25°.

Solvent	Mol. SrO ₂ H ₂ + 8H ₂ O sol. in 1litre
water	0.0835
0.5-N methyl alcohol	0.0820
" ethyl alcohol	0.0744
" propyl alcohol	0.0708
" tert. amyl alcohol	0.0630
" acetone	0.0692
" ether	0.0645

Solubility in organic compds. + Aq at 25°.—
Continued.

Solvent	Mol. $\text{SrO}_2\text{H}_2 + 8\text{H}_2\text{O}$ sol. id 1 litre
0.5-N glycol	0.0922
" glycerine	0.1094
" mannitol	0.1996
" urea	0.0820
" ammonia	0.0785
" diethyl amine	0.0586
" pyridine	0.0694

(Rothmund, Z. phys. Ch. 1909, **69**. 539.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329.)

See also Strontium oxide.

Strontium iodide, SrI_2 , and +6, or $7\text{H}_2\text{O}$.

100 pts. H_2O dissolve at:

0°	20°	40°	70°	100°
164	179	196	250	370 pts. SrI_2 .

(Kremers, Pogg. **103**. 65.)

Sat. aq. solution contains at:

—20°	—10°	—3°	+7°	11°	18°	38°
60.0	60.3	62.2	63.0	63.4	63.5	64.8% SrI_2 ,
52°	63°	77°	81°	97°	105°	120 175°
66.0	68.5	70.5	74.0	79.2	79.4	80.8 85.6% SrI_2 .

(Étard, A. ch. 1894, (7) **2**. 543.)

Sp. gr. of SrI_2 + Aq at 19.5° containing:

5	10	20	30	% SrI_2 ,
1.045	1.091	1.200	1.330	

40	50	60	65	% SrI_2 .
1.491	1.695	1.955	2.150	

(Kremers, Pogg. **103**. 67; calculated by Gerlach, Z. anal. **8**. 285.)

Sat. solution in abs. ethyl alcohol contains at:

—20°	+4°	39°	82°
2.6	3.1	4.3	4.7% SrI_2 .

(Étard, A. ch. 1894, (7) **2**. 565.)

Strontium periodide, $\text{SrI}_2 + 15\text{H}_2\text{O}$.

(Mosnier, A. ch. 1897, (7) **12**. 399.)

SrI_4 . (Herz and Bulla, Z. anorg. 1911, **71**. 255.)

Strontium stannous iodide.

Very sol. in H_2O . (Boullay.)

Strontium zinc iodide, $\text{SrZnI}_4 + 9\text{H}_2\text{O}$.

Hydroscopic. (Ephraim, Z. anorg. 1910, **67**. 385.)

Strontium nitride, Sr_2N_3 .

Decomp. H_2O violently, but not alcohol. (Maquenne, A. ch. (6) **29**. 225.)

Strontium oxide, SrO .

Decomp. by H_2O to SrO_2H_2 , which see.

Sol. in 160 pts. H_2O at 15.56° (Dalton); in 50 pts. at 100° (Dalton); in 130 pts. at 20° (Bineau); in 40 pts. cold, and 20 pts. hot H_2O (Dumas).

Very sl. sol. in alcohol. Insol. in ether.

1 l. methyl alcohol dissolves 11.2 g. SrO . (Neuberg and Rewald, Biochem. Z. 1908, **2**. 540.)

Insol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329; Eidmann, C. C. **1899**, II. 1014.)

Sol. in cane-sugar + Aq.

Solubility in H_2O containing 10 g. sugar at t°.

t°	g. SrO	t°	g. SrO
8	1.21	24	1.87
15	1.48	40	3.55

(Sidérsky, C. C. **1888**. 57.)

See also Strontium hydroxide.

Strontium peroxide, SrO_2 .

Sl. sol. in H_2O . Easily sol. in acids and NH_4Cl + Aq. Insol. in NH_4OH + Aq. (Corroy, Chem. Soc. (2) **11**. 812.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329; Eidmann, C. C. **1899**, II. 1014.)

Strontium oxybromide, SrBr_2 , $\text{SrO} + 9\text{H}_2\text{O}$.

Not hygroscopic; sol. in H_2O . (Tassily, C. R. 1895, **120**. 1339.)

Strontium oxychloride, SrCl_2 , $\text{SrO} + 9\text{H}_2\text{O}$.

Very easily decomp. by H_2O and alcohol (André, A. ch. (6) **3**. 76.)

Strontium oxyiodide, 2SrI_2 , $5\text{SrO} + 30\text{H}_2\text{O}$.

Not hygroscopic; sol. in H_2O . (Tassily, C. R. 1895, **120**. 1339.)

Strontium oxysulphide, $\text{Sr}_2\text{OS}_4 + 12\text{H}_2\text{O}$.

Decomp. by H_2O .

Insol. in alcohol, ether, and CS_2 . (Schäfer.)

Mixture of SrS_2O_8 and SrS_2 . (Geuther, A. **224**. 178.)

Strontium phosphide, Sr_3P_2 .

Crystallized. Sol. in dil. acids; insol. in conc. acids; decomp. by H_2O . Insol. in organic solvents at ord. temp. (Jaboin, C. R. 1899, **129**. 764.)

Strontium selenide, SrSe .

Sl. sol. in H_2O . (Fabre, C. R. **102**. 1400.)

Strontium silicide, SrSi_2 .

Decomp. by H_2O . (Bradley, C. N. 1900, **82**. 150.)

Strontium sulphide, SrS .

Insol. in H_2O with decomp. into SrO , H_2 , and H_2S .
Sol. in acetone, (Eidmann, C. C. 1899, 1; Naumann, B. 1904, 37. 4329.)
Sol. in methyl acetate. (Naumann, B. 1904, 37. 3790.)

Strontium tetrasulphide, SrS_4 .

Deliquescent, and sol. in H_2O and H_2SO_4 .
Aqueous solution decomp. on air.
with 2, or 6 H_2O . (Schöne, Pogg. 117.)

Strontium pentasulphide, SrS_5 .

Known only in solution.

Strontium stannic sulphide.

Sulphostannate, strontium.

Sulphuric acid.

Sulphaluminate, $4\text{Ag}_2\text{S}$, $5\text{Al}_2\text{S}_3$.

Reich, Real. Ac. Linc. 1912, (5) 21, II.

Sulphamic acid, HOSO_2NH_2 .

Imidosulphonic acid.

Ammonium sulphamate, 2NH_3 , SO_3 .

(Worin.)

Ammonium imidosulphonate, which see. (Worin.)

Ammonium sulphamate, acid, 3NH_3 , 2SO_3 .

(Worin.)

Is basic ammonium imidosulphonate, which see. (Berglund.)

Sulphamate, basic, 2BaO , 3SO_3 , 2NH_3 .

What sol. in H_2O , easily in $\text{HCl} + \text{Aq}$. (Jacquelin, A. ch. (3) 8. 304.)

$\text{O}_6(\text{NH}_2)_2$. Sl. sol. in H_2O . Decomp. on heating with H_2O . (Woronin, J. B. 1860.)

Strontium imidosulphonate. (Berglund.)

Sulphamide, $\text{SO}_2(\text{NH}_2)_2$.

Sol. in H_2O . (Regnault, A. ch. 69. 267.)
Sol. in alcohol, ether, etc. (Traube, B. 1904, 37. 3790.)

Sol. in H_2O .
Sol. in abs. alcohol.

Sol. in dry ether. (Divers and Ogawa, Soc. 1902, 81. 504.)

Sol. in liquid NH_3 . (Franklin and L. Am. Ch. J. 1902, 28. 95.)

Sol. in alcohol; very sol. in H_2O . (Hantzsch, J. 1904, 34. 3436.)

Silver sulphamide, $\text{SO}_2(\text{NHAg})_2$.

Sl. sol. in cold H_2O . Sol. in HNO_3 , and $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Traube, B. 26. 607.)

3NH_3 , 2SO_3 . (Jacquelin.)

Is basic ammonium imidosulphonate, which see. (Berglund.)

Sulphamidic acid.

(Freymy.)

See Imidosulphonic acid.

Sulphaminoplatinous acid.

Ammonium tetrasulphaminoplatinite,

$[\text{Pt}(\text{SO}_2\text{NH}_2)_4](\text{NH}_4)_2 + 6\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . (Ramberg, B. 1912, 45. 1512.)

Potassium ———, $[\text{Pt}(\text{SO}_2\text{NH}_2)_4]\text{K}_2 + 2\text{H}_2\text{O}$.

Very sl. sol. in cold H_2O . (Ramberg.)

Sodium ———.

Easily sol. in cold H_2O . (Ramberg.)

Sulphammonic, and Metasulphammonic acids.

(Freymy.)

See Nitrilosulphonic acid.

Monosulphammonic acid.

(Claus.)

See Amidosulphonic acid.

Disulphammonic acid.

(Claus.)

See Imidosulphonic acid.

Trisulphammonic acid.

(Claus.)

See Nitrilosulphonic acid.

Tetrasulphammonic acid.

(Claus.)

Does not exist. See Nitrilosulphonic acid.

Sulphammonium, $\text{S}(\text{NH}_2)_2$, 2NH_3 .

Sol. in liquid NH_3 .

Sol. in abs. alcohol and anhydrous ether. (Moissan, C. R. 1901, 132. 517.)

Sulphantimonic acid.

Sulphantimonates.

The alkali sulphantimonates are sol. in H_2O , but the solutions decomp. on the air; most of the other sulphantimonates are insol. in H_2O ; all sulphantimonates are insol. in alcohol. (Rammelsberg.)

Ammonium sulphantimonate, $(\text{NH}_4)_3\text{SbS}_4$.Sbl. in H_2O .

Sol. in dil. acids with decomp. (Stanek, Z. anorg. 1898, 17. 122.)

 $+4\text{H}_2\text{O}$. (Stanek.)Solubility of $(\text{NH}_4)_3\text{SbS}_4 + 4\text{H}_2\text{O}$ in H_2O at t° .

t°	$(\text{NH}_4)_3\text{SbS}_4$	Solid phase
- 1.9	9.9	Ice
- 5	20.0	"
- 8	30.2	"
-13.5	41.6	Ice + $(\text{NH}_4)_3\text{SbS}_4 \cdot 4\text{H}_2\text{O}$
0	41.6	$(\text{NH}_4)_3\text{SbS}_4 \cdot 4\text{H}_2\text{O}$
+20	47.7	"
30	54.5	"

(Donk, Chem. Weekbl, 1908, 5. 529.)

Solubility of $(\text{NH}_4)_3\text{SbS}_4$ in alcohol at 10° .Solid phase, $(\text{NH}_4)_3\text{SbS}_4 + 4\text{H}_2\text{O}$.

$\text{C}_2\text{H}_5\text{OH}$	$(\text{NH}_4)_3\text{SbS}_4$	$\text{C}_2\text{H}_5\text{OH}$	$(\text{NH}_4)_3\text{SbS}_4$
0	43.2	43.1	8.7
5.1	35.9	53.1	4.1
19.1	23.1	93.3	0

(Donk, l. c.)

Antimonyl sulphantimonate, $(\text{SbO})_2\text{SbS}_4$.Sol. in HCl . (Rammelsberg, Pogg. 1841, 52. 236.)**Barium sulphantimonate, $\text{Ba}_2(\text{SbS}_4) + 3\text{H}_2\text{O}$.**Sol. in H_2O . Insol. in alcohol.**Barium potassium sulphantimonate,** $\text{KBaSbS}_4 + 6\text{H}_2\text{O}$.Easily sol. in H_2O .

Decomp. by acids. (Glatzel, Z. anorg. 1911, 72. 100.)

Bismuth sulphantimonate.

Ppt.

Cadmium sulphantimonate.

Ppt. (Rammelsberg, Pogg. 52. 236.)

Calcium sulphantimonate, $\text{Ca}_2(\text{SbS}_4)_2$.Partially sol. in H_2O . Insol. in alcohol.**Cobaltous sulphantimonate, $\text{Co}_2(\text{SbS}_4)_2$.**Ppt. Decomp. by $\text{HCl} + \text{Aq}$. (Rammelsberg, Pogg. 52. 236.)**Cupric sulphantimonate, $\text{Cu}_2(\text{SbS}_4)_2$.**

Ppt. (Rammelsberg, Pogg. 52. 226.)

Iron (ferrous) sulphantimonate.

Ppt.

Iron (ferric) sulphantimonate, $\text{Fe}_2(\text{SbS}_4)_2$

(Rammelsberg, Pogg. 52. 234.)

Lead sulphantimonate, $\text{Pb}_2(\text{SbS}_4)_2$.Ppt. Decomp. by $\text{KOH} + \text{Aq}$. (Rammelsberg, Pogg. 52. 223.)**Lithium sulphantimonate, $\text{Li}_2\text{SbS}_4 + 8\frac{1}{2}\text{H}_2\text{O}$** 100 g. sat. solution in H_2O contains anhyd. Li_2SbS_4 .Solubility in alcohol at 30° .

$\%$ alcohol	Li_2SbS_4	Solid phase
0	50.8	$\text{Li}_2\text{SbS}_4 \cdot 8\frac{1}{2}\text{H}_2\text{O}$
13.3	46.3	"
51.9	30.7	"
54.8	29.9	"
58.4	30.8	$\text{Li}_2\text{SbS}_4 \cdot 8\frac{1}{2}\text{H}_2\text{O} + \text{H}_2\text{O}$
58.6	32.3	"
65.26	29.31	Li_2SbS_4
74.3	24.1	"
79.5	20.5	"

(Schreinemakers and Jacobs, Ch. W. 1910, 72. 213.)

 $+9\text{H}_2\text{O}$. Very sol. in H_2O . (Brink Dissert. 1891.) $+10\text{H}_2\text{O}$. Solubility of $\text{Li}_2\text{SbS}_4 + 10\frac{1}{2}\text{H}_2\text{O}$ at t° .

t°	Li_2SbS_4	Solid phase
- 1.7	7.1	Ice
- 3.2	12.8	"
- 5.1	17.5	"
-10.8	23.2	"
-15.9	28.5	"
-26.2	35.3	"
-42	40.4	Ice + $\text{Li}_2\text{SbS}_4 \cdot 10\frac{1}{2}\text{H}_2\text{O}$
0	45.5	$\text{Li}_2\text{SbS}_4 \cdot 10\frac{1}{2}\text{H}_2\text{O}$
+10	46.9	"
30	50.1	"
50	51.3	"

(Donk, Chem. Weekbl. 1908, 5. 6)

At 10° , 100 g. sat. $\text{Li}_2\text{SbS}_4 + 10\frac{1}{2}\text{H}_2\text{O}$ % alcohol contain 41.8 g. Li_2SbS_4 ; alcohol, 36.5 g. Li_2SbS_4 . (Donk, l. c.)**Magnesium sulphantimonate, $\text{Mg}_2(\text{SbS}_4)_2$** Deliquescent. Sol. in H_2O . Decm. alcohol.**Mercurous sulphantimonate, $(\text{Hg}_2)_2(\text{SbS}_4)_2$**

Ppt.

Mercuric sulphantimonate, $\text{Hg}_2(\text{SbS}_4)_2$

Ppt. (Rammelsberg, Pogg. 52. 229.)

ric sulphantimonate chloride,
 $\text{g}_3(\text{SbS}_4)_2, 3\text{HgCl}_2, 3\text{HgO}$.
l. in acids, except aqua regia. (Ram-
erg.)

sulphantimonate, $\text{Ni}_2(\text{SbS}_4)_2$.
Decomp. by hot $\text{HCl} + \text{Aq}$. (Ram-
erg, Pogg. 52. 226.)

ium sulphantimonate, K_2SbS_4 .
in H_2O .

ubility of K_2SbS_4 in H_2O at t° .

	K_2SbS_4	Solid phase
	9.5	Ice
	17.1	"
	24.2	"
	35.4	"
	42.9	"
	48.8	"
	52.6	"
	59.6	"
	62	Ice + $\text{K}_2\text{SbS}_4, 6\text{H}_2\text{O}$
	65.5	$\text{K}_2\text{SbS}_4, 6\text{H}_2\text{O}$
	69.1	"
	75.4	$\text{K}_2\text{SbS}_4, 5\text{H}_2\text{O}$
	76.2	"
	77.1	"
	77.7	$\text{K}_2\text{SbS}_4, 3\text{H}_2\text{O}$
	79.2	"

, Chem. Weekbl. 1908, 5. 529, 629, 767.)

ility of K_2SbS_4 in $\text{KOH} + \text{Aq}$ at 25° .

	KOH	Solid phase
	0	$\text{K}_2\text{SbS}_4, 5\text{H}_2\text{O}$
	3.4	$\text{K}_2\text{SbS}_4, 3\text{H}_2\text{O}$
	11.0	"
	16.1	K_2SbS_4
	25.5	"
	40.5	"
	46.9	$\text{K}_2\text{SbS}_4 + \text{KOH}, 2\text{H}_2\text{O}$
	49.9	$\text{KOH}, 2\text{H}_2\text{O}$
	56.3	"

(Donk.)

lity of K_2SbS_4 in alcohol + Aq at 10° .

H	K_2SbS_4	Solid phase
	0	$\text{K}_2\text{SbS}_4, 5\text{H}_2\text{O}$
	0	"
	69.2	"
	76.1	"

no liquid layers are formed.

Composition of the liquid layers.

Alcohol layer		H_2O layer	
$\text{C}_2\text{H}_5\text{OH}$	K_2SbS_4	% alcohol	K_2SbS_4
85	0	1.1	67.4
54.7	2.2	3.4	49.0
46.9	4.2	3.8	45.6
16	27.4		
....	31.1	12.7

(Donk, l. c.)

+ $4\frac{1}{2}\text{H}_2\text{O}$. Deliquescent. Sol. in H_2O ; more sol. than the Na salt.
+ 3, 5, and $6\text{H}_2\text{O}$. See Donk above.
 $2\text{K}_2\text{S}, \text{Sb}_2\text{S}_3$. Decomp. by cold H_2O . (Ditte, C. R. 102. 168.)
 $\text{K}_2\text{S}, 2\text{Sb}_2\text{S}_3 + 3\text{H}_2\text{O}$. Sl. sol. in H_2O . (Ditte.)
 $\text{K}_2\text{S}, \text{Sb}_2\text{S}_3$. Decomp. by H_2O . (Ditte.)
 $\text{K}_2\text{S}, 2\text{Sb}_2\text{S}_3$. (Ditte.)

Silver sulphantimonate, Ag_2SbS_4 .
Insol. in H_2O or acids. Decomp. by $\text{KOH} + \text{Aq}$. (Rammelsberg, Pogg. 52. 218.)

Sodium sulphantimonate, $\text{Na}_2\text{SbS}_4 + 9\text{H}_2\text{O}$.
(Schlippe's salt.) Sol. in 2.9 pts. H_2O at 15° . Aqueous solution is precipitated by alcohol. (Rammelsberg.)
Sol. in 3 pts. cold H_2O . (van den Corput.)
Sol. in 4 pts. cold H_2O . (Duflos.)
Sol. in 1 pt. boiling H_2O . (Duflos.)

Solubility of $\text{Na}_2\text{SbS}_4 + 9\text{H}_2\text{O}$ in H_2O at t° .

t°	Na_2SbS_4	Solid phase
— 0.1	0.5	Ice
— 0.65	4	"
— 0.9	5.7	"
— 1.26	7.8	"
— 1.45	9.2	"
— 1.75	11.2	"
0	11.3	$\text{Na}_2\text{SbS}_4, 9\text{H}_2\text{O}$
15	19.3	"
30	27.1	"
38	32	"
49.6	38.9	"
59.6	45	"
69.6	50.7	"
79.5	57.1	"

(Donk, Chem. Weekbl. 1908, 5. 529, 629, 767.)

Solubility of $\text{Na}_3\text{SbS}_4 + \text{Na}_2\text{S}_2\text{O}_3$ in H_2O .		
Na_3SbS_4	$\text{Na}_2\text{S}_2\text{O}_3$	Solid phase
$t^\circ = 10^\circ$.		
11.8	0	$\text{Na}_3\text{SbS}_4, 9\text{H}_2\text{O}$
4.4	4.9	"
0.8	14.6	"
0.1	27.3	"
0.0	33.6	$\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$
$t^\circ = 30^\circ$.		
19.9	7.7	$\text{Na}_3\text{SbS}_4, 9\text{H}_2\text{O}$
12.5	16.4	"
4.2	37.7	"
1.	43.8	"
1	47	"
1	47.8	$\text{Na}_3\text{SbS}_4 + \text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$
0	45.8	$\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$

(Donk, l. c.)

Solubility of Na_3SbS_4 in alcohol + Aq at t° .
Solid phase, $\text{Na}_3\text{SbS}_4 + 9\text{H}_2\text{O}$.

$t = 0^\circ$		$t = 30^\circ$		$t^\circ = 65^\circ$	
% $\text{C}_2\text{H}_5\text{OH}$	% Na_3SbS_4	% $\text{C}_2\text{H}_5\text{OH}$	% Na_3SbS_4	% $\text{C}_2\text{H}_5\text{OH}$	% Na_3SbS_4
0	11.8	5.	19.3	0	47.9
3.7	8.2	10.3	14.6	4.7	39.3
12.7	3.2	24.8	6.4	8	36.5
29.	0.9	46	1.2	* 54.1	4.1
60.8	0	76.2	0	81.	0

* Two layers are formed.

Composition of above layers.

Alcohol layer		H_2O layer	
% alcohol	% Na_3SbS_4	% alcohol	% Na_3SbS_4
54.1	4.1	8.0	36.5
40.4	10.2	14.3	27.8
33.5	14.1	18.8	24.1
....	27.2	18.0

(Donk, l. c.)

Solubility of Na_3SbS_4 in methyl alcohol + Solid phase, $\text{Na}_3\text{SbS}_4 + 9\text{H}_2\text{O}$.			
$t = 0^\circ$		$t = 30^\circ$	
% CH_3OH	% Na_3SbS_4	% CH_3OH	% Na_3SbS_4
3.4	8.6	0	27.1
15.5	2.8	18.1	12.5
23.1	2.1	33.1	5.9
50.3	0.3	65.7	0.1
57	0.1	84.2	0.1
81.7	0.05	91.2	1.2
92	0.2	94.	3.9
95.9	2.0

(Donk, l. c.)

Sodium sulphantimonate thiosulphate,
 $\text{Na}_3\text{SbS}_4, 2\text{Na}_2\text{S}_2\text{O}_3 + 20\text{H}_2\text{O}$.
Efflorescent, and decomp. by H_2O . (Iger, Arch. Pharm. (2) 147. 193.)
No double salt exists. See Donk, $\text{Na}_3\text{SbS}_4 + \text{Na}_2\text{S}_2\text{O}_3$ under Na_3SbS_4 .

Strontium sulphantimonate.
Sol. in H_2O ; pptd. by alcohol.

Uranium sulphantimonate.
Ppt.

Zinc sulphantimonate, $\text{Zn}_3(\text{SbS}_4)_2$.
Ppt.. Sol. in hot $\text{Na}_3\text{SbS}_4 + \text{Aq}$; insol. $\text{ZnSO}_4 + \text{Aq}$. Partially sol. in $\text{KOH} + \text{Aq}$. sol. in hot $\text{HCl} + \text{Aq}$. (Rammelsberg, P. 52. 233.)

Sulphantimonous acid.

Ammonium metasulphantimonite, NH_4SbS_3
Insol. in H_2O . (Rouget, C. R. 1898, 1145.)
+ $2\text{H}_2\text{O}$. Insol. in H_2O and alcohol.
Decomp. in the air. (Stanek, Z. an 1898, 17. 119.)

Ammonium orthosulphantimonite, $(\text{NH}_4)_2\text{Sb}_2\text{S}_5$.
Easily decomp. Stable only in press of $(\text{NH}_4)_2\text{S}$. Sol. in H_2O . Insol. in alcoh by which it is pptd. from aqueous soluti (Pouget, A. ch. 1899, (7) 18. 536.)

Ammonium parasulphantimonite, $(\text{NH}_4)_3\text{Sb}_2\text{S}_7$.
Stable in the air.
Insol. in H_2O .
Decomp. by acids. (Stanek, Z. an 1898, 17. 120.)
Stable; cryst. from hot solutions. (Pou C. R. 1898, 126. 1145.)

silver orthosulphantimonite, Ag_2SbS_3 .
 . by H_2O . (Pouget, A. ch. 1899, 1.)

basulphantimonite, BaSb_2S_4
 H_2O .
 . H_2O . (Pouget, A. ch. 1899, .)

orthosulphantimonite, $\text{Ba}_2\text{Sb}_2\text{S}_4$
 O .
 . in the air and by H_2O . Some-
 n $\text{BaS} + \text{Aq}$. (Pouget, C. R. 1898, .)

orthosulphantimonite, $\text{Ba}_2\text{Sb}_2\text{S}_4$
 O .
 . by H_2O .
 nsol. in $\text{BaS} + \text{Aq}$. (Rouget.)

phantimonite, $\text{Ba}_2\text{Sb}_2\text{S}_4 + 10\text{H}_2\text{O}$.
 om aq. solution of ortho and pyro-
 ts. (Pouget.)
 . $+16\text{H}_2\text{O}$. (Pouget, A. ch. 1899, .)

phantimonite basic, $\text{Ca}(\text{OH})\text{SbS}_2$.
 H_2O .
 conc. HCl . (Pouget, A. ch. 1899, .)

pyrosulphoantimonite, $\text{Ca}_2\text{Sb}_2\text{S}_4$
 H_2O .
 H_2O without decomp. (Pouget, , 126. 1793.)

orthosulphantimonite, $\text{Co}_2\text{Sb}_2\text{S}_4$.
 ouget, A. ch. 1899, (7) 18. 554.)

basulphantimonite, CuSbS_2 .
 mixture of HNO_3 and tartaric acid
 ation of S.
 $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by hot
 alkali sulphides $+ \text{Aq}$. (Sommer-
 rg. 1898, 18. 430.)
Volsber. ite. Sol. in $\text{HNO}_3 + \text{Aq}$
 ation of S and Sb_2O_3 .

orthosulphantimonite, Cu_2SbS_3 .
 lad, Z. anorg. 1898, 18. 432.)
 sol. in H_2O . Decomp. by H_2O .
 . ch. 1899, (7) 18. 556.)

ulphantimonite, $\text{Cu}_2\text{Sb}_4\text{S}_7$.
erjarite.

osulphantimonite, $\text{Cu}_2\text{Sb}_2\text{S}_4$.
 ouget, A. ch. 1899, (7) 18. 557.)

ad sulphantimonite, Cu_2SbS_3 ,
 SbS_3 .
urnonite. Decomp. by $\text{HNO}_3 +$
qua regia.

Cuprous potassium orthosulphantimonite,
 Cu_2KSbS_3 .
 Ppt.; easily decomp. by H_2O . (Pouget,
 C. R. 1899, 129. 104.)
 $+3\text{H}_2\text{O}$. Ppt., decomp. by H_2O . (Pou-
 get, A. ch. 1899, (7) 18. 556.)

Iron (ferrous) orthosulphantimonite,
 $\text{Fe}_2(\text{Sb}_2\text{S}_4)_2$.
 Ppt. (Pouget, A. ch. 1899, (7) 18. 554.)
 Min. *Berthierite*. Sl. sol. in $\text{HCl} + \text{Aq}$;
 easily sol. in aqua regia.

Lead orthosulphantimonite, $\text{Pb}_2(\text{SbS}_3)_2$.
 Ppt. Very sl. sol. in H_2O . Decomp. by
 H_2O . (Pouget, A. ch. 1899, (7) 18. 553.)
 Min. *Boulangerite*. Completely sol. in
 hot $\text{HCl} + \text{Aq}$; decomp. by $\text{HNO}_3 + \text{Aq}$.

Lead sulphantimonite.
 Sol. in boiling conc. $\text{HNO}_3 + \text{Aq}$. (Four-
 net.)
 $\text{Pb}(\text{SbS}_3)_2$. Min. *Zinckenite*. Decomp.
 by hot $\text{HCl} + \text{Aq}$.
 $4\text{PbS}, \text{Sb}_2\text{S}_3$. Min. *Plagionite*.
 $2\text{PbS}, \text{Sb}_2\text{S}_3$. Min. *Jamesonite*. Decomp.
 by hot $\text{HCl} + \text{Aq}$.
 $4\text{PbS}, \text{Sb}_2\text{S}_3$. Min. *Meneghinite*.
 $5\text{PbS}, \text{Sb}_2\text{S}_3$. Min. *Geokronite*.
 $6\text{PbS}, \text{Sb}_2\text{S}_3$. Min. *Kibrickenite* (?).

Lead potassium orthosulphantimonite,
 PbKSbS_3 .
 Very sl. sol. in H_2O .
 Decomp. by H_2O . (Pouget, A. ch. 1899,
 (7) 18. 554.)

Lead silver sulphantimonite, $(\text{Ag}_2, \text{Pb})_2\text{Sb}_4\text{S}_{11}$.
 Min. *Freieslebenite*.

Lithium orthosulphantimonite, Li_2SbS_3
 $+3\text{H}_2\text{O}$.
 Very deliquescent.
 Very sol. in H_2O . (Pouget, A. ch. 1899,
 (7) 18. 530.)

Lithium parasulphantimonite,
 $\text{Li}_2\text{Sb}_4\text{S}_7 + 3\text{H}_2\text{O}$.
 Ppt. (Pouget, A. ch. 1899, (7) 18. 531.)

Lithium silver orthosulphantimonite,
 $\text{LiAg}_2\text{SbS}_3$.
 Decomp. by H_2O . (Pouget, A. ch. 1899,
 (7) 18. 551.)

Manganous orthosulphantimonite, $\text{Mn}_2\text{Sb}_2\text{S}_4$.
 Ppt. Sl. sol. in H_2O . (Pouget, A. ch
 1899, (7) 18. 553.)

Manganous potassium orthosulphantimonite,
 MnKSbS_3 .
 Sl. sol. in H_2O . Decomp. by H_2O . (Pou-
 get, A. ch. 1899, (7) 18. 553.)

Nickel orthosulphantimonite, $\text{Ni}_3\text{Sb}_2\text{S}_4$.

Ppt. (Pouget, A. ch. 1899, (7) 18. 554.)

Potassium metasulphantimonite, KSbS_3 .

Insol. in cold H_2O . Decomp. by hot H_2O . (Pouget, A. ch. 1899, (7) 18. 513.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O , but decomp. quickly.

Sol. in H_2O . (Stanek, Z. anorg. 1898, 17. 119.)

Potassium orthosulphantimonite, K_2SbS_3 .

Very deliquescent.

Very sol. in H_2O .

Decomp. by acids. (Pouget, A. ch. 1899, (7) 18. 518.)

Potassium sulphantimonite, $\text{K}_2\text{Sb}_2\text{S}_7 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O and not decomp. thereby. (Pouget, A. ch. 1899, (7) 18. 522.)

Decomp. in the air.

Sol. in $\text{K}_2\text{S} + \text{Aq}$. (Stanek, Z. anorg. 1898, 17. 120.)

$2\text{K}_2\text{S}$, Sb_2S_3 . Sol. in H_2O . (Ditte, C. R. 102. 68.)

$x\text{K}_2\text{S}$, $y\text{Sb}_2\text{S}_3$. Deliquescent. When K_2S is in excess, sol. in H_2O ; when Sb_2S_3 is in excess, partially sol. Aqueous solution is decomp. by all acids, even CO_2 , and by K_2CO_3 , Na_2CO_3 , NaHCO_3 , KHCO_3 , $\text{NH}_4\text{HCO}_3 + \text{Aq}$. Insol. in absolute alcohol. (Kohl.)

Potassium hydrogen sulphantimonite, KHSb_2S_7 .

(Pouget, A. ch. 1899, (7) 18. 522.)

Potassium silver orthosulphantimonite, Ag_2KSbS_3 .

Decomp. by boiling H_2O . (Pouget, C. R. 1897, 124. 1519.)

Potassium zinc orthosulphantimonite, KZnSbS_3 .

Decomp. by H_2O . (Pouget, A. ch. 1899, (7) 18. 552.)

Silver orthosulphantimonite, Ag_2SbS_3 .

Ppt. Sl. sol. in H_2O . (Pouget, A. ch. 1899, (7) 18. 547.)

Min. *Pyrargyrite*. Sol. in $\text{HNO}_3 + \text{Aq}$ with residue of S and Sb_2O_3 . $\text{KOH} + \text{Aq}$ dissolves out Sb_2S_3 .

Silver sulphantimonite.

AgSbS_3 . Min. *Miargyrite*.

$5\text{Ag}_2\text{S}$, Sb_2S_3 . Min. *Stephanite*. Easily decomp. by warm $\text{HNO}_3 + \text{Aq}$.

$12\text{Ag}_2\text{S}$, Sb_2S_3 . Min. *Polyargyrite*.

Silver sodium orthosulphantimonite, $\text{Ag}_2\text{NaSbS}_3$.

Decomp. by H_2O . Pouget, A. ch. 1899, (7) 18. 551.)

Sodium metasulphantimonite, NaSbS_3 .

Deliquescent. Decomp. by hot H_2O . When Na_2S is in excess, sol. in H_2O , but partially sol. if Sb_2S_3 is in excess. (Unger, Arch. Pharm. (2) 148. 1.)

Ppt. Insol. in H_2O . (Pouget, C. R. 1898, 126. 1145.)

Sodium orthosulphantimonite, $\text{Na}_2\text{SbS}_3 + 9\text{H}_2\text{O}$.

Decomp. in solution in H_2O . (Pouget, C. R. 1898, 126. 1144.)

Sodium sulphantimonite, $\text{Na}_2\text{Sb}_2\text{S}_7 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Pouget, C. R. 1898, 126. 1145.)

$\text{Na}_2\text{Sb}_2\text{S}_7$. (Pouget, C. R. 1898, 126. 1144.) $4\text{Na}_2\text{S}$, $3\text{Sb}_2\text{S}_3 + 3\text{H}_2\text{O}$. Permanent; sl. in H_2O . Insol. in alcohol and ether. (Kohl.)

Strontium orthosulphantimonite, $\text{Sr}_2\text{Sb}_2\text{S}_7 + 10\text{H}_2\text{O}$.

Sol. in H_2O . (Pouget, C. R. 1898, 126. 1793.)

Strontium pyrosulphantimonite, $\text{Sr}_2\text{Sb}_2\text{S}_7 + 15\text{H}_2\text{O}$.

Sol. in H_2O without essential decomp. (Pouget, C. R. 1898, 126. 1793.)

Zinc orthosulphantimonite, $\text{Zn}_2\text{Sb}_2\text{S}_7$.

Ppt. (Pouget, A. ch. 1899, (7) 18. 552.)

Orthosulpharsenic acid, H_2AsS_4 .

Ppt. Loses H_2S by prolonged boiling with H_2O . (Nilson, J. pr. (2) 14. 145.)

See also Sulphoxyarsenic acid.

Ammonium sulpharsenate, $(\text{NH}_4)_2\text{As}_2\text{S}_7$.

Known only in solution in H_2O . Decomp. on boiling into—

NH_4AsS_4 . Sol. in alcohol.

$(\text{NH}_4)_2\text{AsS}_4$. Sol. in H_2O . Precipitated by alcohol.

$(\text{NH}_4)_2\text{S}$, $12\text{As}_2\text{S}_3$. Ppt. Insol. in H_2O .

Ammonium magnesium sulpharsenate, $(\text{NH}_4)_2\text{S}$, MgS , As_2S_3 .**Ammonium sodium sulpharsenate, $(\text{NH}_4)_2\text{AsS}_4$, Na_2AsS_4 .**

Much more sol. in H_2O than Na_2AsS_4 ; sl. sol. in cold, more sol. in hot alcohol. (Berzelius.)

Barium sulpharsenate, $\text{Ba}(\text{AsS}_4)_2$.

Sol. in H_2O and alcohol. Decomp. by evaporation.

$\text{Ba}_2\text{As}_2\text{S}_7$. Sol. in H_2O in all proportions with decomp. Decomp. by alcohol.

$\text{Ba}_2(\text{AsS}_4)_2$. Sol. in H_2O . Insol. in alcohol.

BaS , $3\text{As}_2\text{S}_3$. Ppt. Insol. in H_2O .

Barium potassium sulpharsenate,
 $\text{KBaAsS}_4 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O .

Decomp. by acids with separation of As_2S_5 .
 (Glatzel, Z. anorg. 1911, 71. 209.)

Barium sulpharsenate sulpharsenite,
 $\text{Ba}_3(\text{AsS}_4)_2, \text{Ba}_2\text{As}_2\text{S}_5 + 4\text{H}_2\text{O}$.

Sl. sol. in cold, more easily in hot H_2O .
 (Nilson.)

Bismuth sulpharsenate, $2\text{Bi}_2\text{S}_3, 3\text{As}_2\text{S}_5$.

Sol. in $\text{Na}_2\text{AsS}_4 + \text{Aq}$.

$\text{Bi}_2\text{S}_3, 3\text{As}_2\text{S}_5$. As above. (Berzelius.)

Cadmium sulpharsenate.

Ppt. (Berzelius, Pogg. 7. 88.)

Calcium sulpharsenate, $\text{Ca}_2\text{As}_2\text{S}_7$.

Sol. in H_2O and alcohol.

$\text{Ca}_2(\text{AsS}_4)_2$. Easily sol. in H_2O . Insol. in alcohol.

+10 H_2O . Easily sol. in H_2O . (Nilson, J. pr. (2) 14. 169.)

$5\text{CaS}, 2\text{As}_2\text{S}_5 + 6\text{H}_2\text{O}$. Easily sol. in H_2O .
 (Nilson, J. pr. (2) 14. 163.)

Cerous sulpharsenate, $\text{Ce}_2\text{As}_2\text{S}_7$.

Ppt.

$\text{Ce}_2(\text{AsS}_4)_2$. Ppt.

$\text{Ce}_4(\text{As}_2\text{S}_7)_2$. Ppt.

Cobaltous sulpharsenate, $\text{Co}_2\text{As}_2\text{S}_7$.

Ppt. Sol. in excess of sodium sulpharsenate + Aq.

Cuprous sulpharsenate, Cu_2AsS_4 .

Ppt. (Preis, A. 257. 201.)

Min. *Enargite*. *Clarite*. Not wholly decomp. by $\text{HCl} + \text{Aq}$. Sol. in $\text{HCl} + \text{Aq}$ with residue of As_2O_3 . Not attacked by $\text{KOH} + \text{Aq}$.

Cupric sulpharsenate, $\text{Cu}_2\text{As}_2\text{S}_7$.

Ppt. Sol. in $(\text{NH}_4)_2\text{S} + \text{Aq}$. Decomp. by $\text{NH}_4\text{OH} + \text{Aq}$. (Berzelius.)

$\text{Cu}_2(\text{AsS}_4)_2$. Ppt. (Preis, A. 257. 201.)

Glucinum sulpharsenate.

Sl. sol. in H_2O .

Gold sulpharsenate, AuAsS_4 .

Sol. in pure H_2O . Insol. in $\text{Na}_2\text{AsS}_4 + \text{Aq}$.
 $2\text{Au}_2\text{S}_3, 3\text{As}_2\text{S}_5$. Sol. in H_2O . (Berzelius.)

Iron (ferrous) sulpharsenate, $\text{Fe}_2\text{As}_2\text{S}_7$.

Ppt. Sol. in $\text{Na}_2\text{AsS}_4 + \text{Aq}$. (Berzelius.)

Iron (ferric) sulpharsenate, $\text{Fe}_4(\text{As}_2\text{S}_7)_3$.

Ppt. Sol. in $\text{Na}_2\text{AsS}_4 + \text{Aq}$. (Berzelius.)

Lead sulpharsenate, $\text{Pb}_2\text{As}_2\text{S}_7$.

Ppt. (Berzelius.)

$\text{Pb}_2(\text{AsS}_4)_2$. Ppt.

Lithium sulpharsenate, Li_2AsS_4 .

Easily sol. in hot, less sol. in cold H_2O . Insol. in alcohol.

$\text{Li}_4\text{As}_2\text{S}_7$. Completely sol. in H_2O . Decomp. by alcohol.

LiAsS_3 . Known only in acid solution.

Magnesium sulpharsenate, $\text{Mg}_2\text{As}_2\text{S}_7$.

Sol. in all proportions of H_2O , and in alcohol.

$\text{Mg}_2(\text{AsS}_4)_2$. Sol. in H_2O . Decomp. alcohol.

$3\text{MgS}, \text{As}_2\text{S}_5$. Nearly insol. in H_2O .

$5\text{MgS}, 2\text{As}_2\text{S}_5 + 15\text{H}_2\text{O}$. Very sol. in H_2O . (Nilson.)

Manganous sulpharsenate, $\text{Mn}_2\text{As}_2\text{S}_7$.

Sl. sol. in H_2O .

$\text{Mn}_2(\text{AsS}_4)_2$. Permanent. Sl. sol. in H_2O .

$6\text{MnS}, \text{As}_2\text{S}_5$. Sl. sol. in H_2O .

Mercurous sulpharsenate, $(\text{Hg}_2)_2\text{As}_2\text{S}_7$.

Ppt.

Mercuric sulpharsenate, $\text{Hg}_2\text{As}_2\text{S}_7$.

Ppt. (Berzelius, Pogg. 7. 29.)

$\text{Hg}_2(\text{AsS}_4)_2$. Ppt. (Preis, A. 257. 200.)

Nickel sulpharsenate, $\text{Ni}_2(\text{AsS}_4)_2$.

Ppt. Not decomp. by $\text{HCl} + \text{Aq}$. Sol. in $\text{Na}_2\text{AsS}_4 + \text{Aq}$. (Berzelius.)

$2\text{NiS}, \text{As}_2\text{S}_5$. As above.

Potassium sulpharsenate, KAsS_3 .

Known only in alcoholic solution.

$\text{K}_4\text{As}_2\text{S}_7$. Deliquescent. Sol. in H_2O , from which alcohol ppts. K_2AsS_4 .

K_2AsS_4 . Deliquescent. Very sol. in H_2O , from which it is precipitated by alcohol.

+ H_2O . Very deliquescent. (Nilson; J. pr. (2) 14. 159.)

Potassium sodium sulpharsenate.

Sol. in H_2O .

Silver sulpharsenate, Ag_2AsS_4 .

Ppt. (Berzelius, Pogg. 7. 29.)

$\text{Ag}_2\text{As}_2\text{S}_7$. Ppt.

Sodium sulpharsenate, NaAsS_3 .

Known only in alcoholic solution.

$\text{Na}_4\text{As}_2\text{S}_7$. Sol. in H_2O . Alcohol ppts. Na_2AsS_4 from H_2O solution.

$\text{Na}_3\text{AsS}_4 + 7\frac{1}{2}\text{H}_2\text{O}$. Easily sol. in H_2O , from which it is precipitated by alcohol.

+8 H_2O . Insol. in alcohol; very sol. in H_2O . (McCay, Z. anal. 1895, 34. 726.)

+9 H_2O . (Nilson, J. pr. (2) 14. 160.)

$\text{Na}_2\text{S}, 12\text{As}_2\text{S}_5$ (?). Insol. in H_2O .

Sodium zinc sulpharsenate, $\text{NaZnAsS}_4 + 4\text{H}_2\text{O}$.

Sol. in hot H_2O with decomp. (Preis, A. 257. 202.)

Strontium sulpharsenate, $\text{Sr}_2(\text{AsS}_4)_2$.

Easily sol. in H_2O ; insol. in alcohol.
 $\text{Sr}_2\text{As}_2\text{S}_7$. Easily sol. in H_2O , from which alcohol ppts. $\text{Sr}_2(\text{AsS}_4)_2$.

Strontium sulpharsenate sulpharsenite, $\text{Sr}_2(\text{AsS}_4)_2, \text{Sr}_2\text{As}_2\text{S}_7 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O . (Nilson, J. pr. (2) 14. 162.)

Thallous sulpharsenate, Tl_2AsS_4 .

Not decomp. by H_2O . Decomp. by dil. acids. Insol. in dil. alkali sulphides. Partially decomp. by boiling with a conc. solution of sodium sulphide. (Hawley, J. Am. Chem. Soc. 1907, 29. 1013.)

Tin (stannous) sulpharsenate.

Ppt.

Tin (stannic) sulpharsenate.

Ppt.

Uranic sulpharsenate, $2\text{U}_2\text{S}_5, \text{As}_2\text{S}_5$.

Ppt. Sol. in $\text{Na}_2\text{AsS}_4 + \text{Aq}$.

Zinc sulpharsenate, $\text{Zn}_2(\text{AsS}_4)_2$.

Ppt. (Berzelius.)

$2\text{ZnS}, \text{As}_2\text{S}_5$. Ppt. (Berzelius.)

$\text{ZnS}, \text{As}_2\text{S}_5$. (Wöhler.)

Disulpharsenic acid.

See Disulphoxyarsenic acid.

Sulpharseniosulphomolybdic acid.**Ammonium sulpharseniosulphomolybdate, $(\text{NH}_4)_4\text{As}_2\text{S}_7(\text{MoS}_4)_2 + 5\text{H}_2\text{O}$.**

Very unstable.

Sol. in H_2O with decomp. (Weinland, Z. anorg. 1897, 15. 49.)

Barium —, $\text{Ba}_2\text{As}_2\text{S}_7(\text{MoS}_4)_2 + 14\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Weinland.)

Potassium —, $\text{KAsS}_4(\text{MoS}_4) + 4\text{H}_2\text{O}$.

(Weinland.)

$\text{K}_4\text{As}_2\text{S}_7(\text{MoS}_4)_2 + 8\text{H}_2\text{O}$. Sol. in H_2O . Decomp. by mineral acids. Insol. in alcohol. (Weinland.)

Sodium —, $\text{NaAsS}_4(\text{MoS}_4) + 6\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in dil. NaOH and $\text{NH}_3 + \text{Aq}$. (Weinland.)

$\text{Na}_4\text{As}_2\text{S}_7(\text{MoS}_4)_2 + 14\text{H}_2\text{O}$. Sol. in H_2O . Decomp. by mineral acids. (Weinland.)

Sulpharseniosulphoxymolybdic acid.**Barium sulpharseniosulphoxymolybdate, $\text{Ba}_2\text{As}_2\text{S}_7(\text{Mo}_2\text{S}_2\text{O}_8)_2 + 12\text{H}_2\text{O}$.**

Sol. in H_2O . (Weinland, Z. anorg. 1897, 15. 60.)

Magnesium sulpharseniosulphoxymolybdate, $\text{Mg}_2\text{As}_2\text{S}_7(\text{Mo}_2\text{S}_2\text{O}_8)_2 + 16\text{H}_2\text{O}$.

Very sol. in H_2O . (Weinland.)

Potassium —, $\text{KAsS}_4(\text{MoSO}_4) + 2\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Weinland.)
 $\text{K}_4\text{As}_2\text{S}_7(\text{Mo}_2\text{S}_2\text{O}_8)_2 + 6\text{H}_2\text{O}$. Very sol. in H_2O . (Weinland.)
 $+ 10\text{H}_2\text{O}$. (Weinland.)

Sodium —, $\text{NaAsS}_4(\text{MoSO}_4) + 5\text{H}_2\text{O}$.

Sl. sol. in cold, very sol. in hot H_2O . (Weinland.)

$\text{Na}_4\text{As}_2\text{S}_7(\text{Mo}_2\text{S}_2\text{O}_8)_2 + 15\text{H}_2\text{O}$. Sol. in H_2O . (Weinland.)

Sulpharsenious acid.**Ammonium sulpharsenite, NH_4AsS_3 .**

Insol. in H_2O . Ppt. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Sl. attacked by boil Aq . (Nilson, J. pr. (2) 14. 42.)

$(\text{NH}_4)_4\text{As}_2\text{S}_5 = 2(\text{NH}_4)_2\text{S}, \text{As}_2\text{S}_5$.
 H_2O , from which alcohol ppts. $(\text{NH}_4)_2\text{AsS}_3 = 3(\text{NH}_4)_2\text{S}, \text{As}_2\text{S}_5$.
 on air; sol. in H_2O . Insol. in alc.
 $(\text{NH}_4)_4\text{As}_2\text{S}_{10}$. Sol. in H_2O . (J. pr. (2) 14. 160.)

Barium sulpharsenite, $\text{Ba}_2\text{As}_2\text{S}_5$.

Sl. sol. in H_2O . Decomp. by alk. $+ 5\text{H}_2\text{O}$. Sl. sol. in H_2O . (Nilson, J. pr. (2) 14. 46.)

$+ 15\text{H}_2\text{O}$. Sl. sol. in cold H_2O .
 $\text{Ba}_2(\text{AsS}_3)_2$. Sl. sol. in H_2O . Ppt. by alcohol.

$+ 14\text{H}_2\text{O}$. Sl. sol. in cold, conc. H_2O . (Nilson.)

$\text{Ba}(\text{AsS}_3)_2 + 2\text{H}_2\text{O}$. Insol. in H_2O . (Nilson, J. pr. (2) 14. 44.)

$\text{BaAs}_2\text{S}_{10}$. Insol. in $\text{HCl} + \text{Aq}$.

Bismuth sulpharsenite, $2\text{Bi}_2\text{S}_3, \text{As}_2\text{S}_5$.

Ppt.

Cadmium sulpharsenite.

Ppt. (Berzelius, Pogg. 7. 146.)

Calcium sulpharsenite, $\text{Ca}_2\text{As}_2\text{S}_5$.

Sol. in H_2O , from which also $\text{Ca}_2(\text{AsS}_3)_2$.

$\text{Ca}_2(\text{AsS}_3)_2$. Sol. in H_2O .
 $+ 15\text{H}_2\text{O}$. Precipitated by alkali.
 $\text{Ca}(\text{AsS}_3)_2 + 10\text{H}_2\text{O}$. Sol. in H_2O . (J. pr. (2) 14. 54.)

$\text{CaAs}_2\text{S}_{10} + 10\text{H}_2\text{O}$ (?). Insol. in H_2O . (Nilson.)

$\text{CaAs}_2\text{S}_{10} + 10\text{H}_2\text{O}$ (?). Sl. sol. in H_2O . (Nilson.)

$\text{Ca}_2\text{As}_2\text{S}_{10} + 25\text{H}_2\text{O}$. Sl. sol. in H_2O . (Nilson.)

Cerous sulpharsenite, $\text{Ce}_2\text{As}_2\text{S}_5$.

Ppt.

sulpharsenite, $2\text{Cr}_2\text{S}_3, 3\text{As}_2\text{S}_3$.

Insol. in $\text{Na}_2\text{S} + \text{Aq}$.

; sulpharsenite, $2\text{CoS}, \text{As}_2\text{S}_3$.

Sol. in excess of sodium sulpharsen-

sulpharsenite,
 $\text{S}_7, 2\text{As}_2\text{S}_3 = \text{Cu}_4\text{As}_4\text{S}_9$.

3innite. Decomp. by hot acids and
q.

$\text{As}_2\text{S}_3 = \text{Cu}_4\text{As}_4\text{S}_9$. Decomp. by
 OH and $\text{K}_2\text{S} + \text{Aq}$. (Sommerlad, Z.
98, 18. 434.)

ulpharsenite, Cu_2AsS_3 .

n H_2O or $\text{HCl} + \text{Aq}$. Sol. in Na_2AsS_3

S_6 . Ppt. (Berzelius.)

1 sulpharsenite, $2\text{GlS}, \text{As}_2\text{S}_3$.

Sol. in acids; partly sol. in NH_4OH

harsenite, $2\text{Au}_2\text{S}_3, 3\text{As}_2\text{S}_3$.

Berzelius.)

rous) sulpharsenite.

Sol. in $\text{Na}_2\text{AsS}_3 + \text{Aq}$. (Berzelius.)

ric) sulpharsenite.

Sol. in excess of a ferric salt, or
 $+ \text{Aq}$. (Berzelius.)

harsenite, $\text{Pb}_2\text{As}_2\text{S}_3$.

Min. *Dufrenoyite*.

$\frac{1}{2}, \frac{1}{2} = \text{PbS}, \text{As}_2\text{S}_3$. Min. *Sartorite*.
 S_7 . Min. *Jordanite*.

sulpharsenites.

ble K salts.

um sulpharsenite, $\text{Mg}_2\text{As}_2\text{S}_3$.

; completely sol. in H_2O . Easily sol.
l. (Berzelius.)

). Sl. sol. in H_2O . (Nilson.)

$\text{S}_3)_2 + 5\text{H}_2\text{O}$. Slowly sol. in both
hot H_2O . (Nilson, J. pr. (2) 14.

$\text{S}_3)_2 + 9\text{H}_2\text{O}$. (Nilson.)

us sulpharsenite, $\text{Mn}_2\text{As}_2\text{S}_3$.

Decomp. by $\text{HCl} + \text{Aq}$.

us sulpharsenite, $(\text{Hg}_2)_2\text{As}_2\text{S}_3$.

Berzelius.)

; sulpharsenite, $\text{Hg}_2\text{As}_2\text{S}_3$.

$\text{S}_3)_2$. Ppt. (Berzelius, Pogg. 7. 149.)

ulpharsenite, $\text{Ni}_2(\text{AsS}_3)_2$.

(Berzelius.)

Platinum sulpharsenite, $\text{Pt}_2\text{As}_2\text{S}_3$.

Ppt.

Potassium sulpharsenite, $\text{K}_4\text{As}_2\text{S}_3$.

Decomp. by H_2O or alcohol. (Berzelius.)
 K_2AsS_3 . Sol. in H_2O . Insol. in alcohol.

(Berzelius.)

$\text{K}_2\text{As}_4\text{S}_7$. Sol. in H_2O and alcohol. (Ber-
zelius.)

K_2AsS_3 . Decomp. by H_2O . (Berzelius.)
 $+ 2\frac{1}{2}\text{H}_2\text{O}$. Not wholly sol. in H_2O . (Nil-
son, J. pr. (2) 14. 30.)

$\text{K}_2\text{As}_4\text{S}_7 + 8\text{H}_2\text{O}$. (Nilson.)

$\text{KAs}_2\text{S}_3 + \text{H}_2\text{O}$. Insol. in H_2O . Slowly at-
tacked by hot $\text{HCl} + \text{Aq}$. Sol. in $\text{KOH} + \text{Aq}$.
(Nilson.)

Silver sulpharsenite, $12\text{Ag}_2\text{S}, \text{As}_2\text{S}_3$.

Ppt. (Sommerlad, Z. anorg. 1898, 18.
428.)

$5\text{Ag}_2\text{S}, \text{As}_2\text{S}_3 = \text{Ag}_5\text{AsS}_4$. (Sommerlad.)

Ag_2AsS_3 . Min. *Proustite*. Sol. in $\text{HNO}_3 +$
 Aq . $\text{KOH} + \text{Aq}$ dissolves out Sb_2S_3 . (Senar-
mont, A. ch. (3) 32. 129; Wöhler, A. 27. 159.)

$2\text{Ag}_2\text{S}, \text{As}_2\text{S}_3$. Partially sol. in $\text{HNO}_3 +$
 Aq . (Berzelius.)

AgAsS_3 . (Berzelius, Pogg. 7. 150.)

Sodium sulpharsenite, $\text{NaAsS}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Attacked by $\text{HCl} + \text{Aq}$ with difficulty.
(Nilson, J. pr. (2) 14. 37.)

$+ 1\frac{1}{2}\text{H}_2\text{O}$. Forms coagulum with cold, sol.
in hot H_2O . (Nilson.)

$\text{Na}_2\text{As}_4\text{S}_7 + 6\text{H}_2\text{O}$. Sol. in much H_2O ; not
easily decomp. by $\text{HCl} + \text{Aq}$. (Nilson.)

$\text{NaAs}_2\text{S}_3 + 4\text{H}_2\text{O}$. Ppt. (Nilson, J. pr. (2)
14. 3.)

Strontium sulpharsenite, $3\text{SrS}, \text{As}_2\text{S}_3 +$
 $15\text{H}_2\text{O}$.

Sol. in $\text{H}_2\text{O} + \text{Aq}$; insol. in alcohol. (Voigt
and Göttling.)

$2\text{SrS}, \text{As}_2\text{S}_3$. Sol. in H_2O ; decomp. by
alcohol.

$+ 15\text{H}_2\text{O}$. (Nilson, J. pr. (2) 14. 53.)

$\text{Sr}(\text{AsS}_3)_2 + 2\frac{1}{2}\text{H}_2\text{O}$. Sl. sol. in H_2O . (Nil-
son.)

Thallous sulpharsenite, TlAsS_3 .

Ppt. Decomp. by $\text{KOH} + \text{Aq}$. (Gunning,
J. B. 1868. 247.)

Above compound is a mixture of As_2S_3 and
 Tl_2S . (Hawley, J. Am. Chem. Soc. 1907,
29. 1012.)

Min. *Lorandite*. (Kuenner and Loezka,
C. C. 1904, II. 844.)

Tin (stannous) sulpharsenite, $\text{Sn}_2\text{As}_2\text{S}_3$.

Ppt.

Tin (stannic) sulpharsenite, SnAs_2S_3 .

Ppt. (Berzelius, Pogg. 7. 147.)

Uranic sulpharsenite, $2\text{U}_2\text{S}_3, \text{As}_2\text{S}_3$.

Ppt.

Zinc sulpharsenite.

Ppt. (Berzelius, Pogg. 7. 145.)

Zirconium sulpharsenite, $2\text{Zr}_2\text{S}_3, \text{As}_2\text{S}_3$.

Ppt. Insol. in solutions of alkali sulpharsenites. Sl. sol. in Na_2S Aq. Not decomp. by acids. (Berzelius.)

"Sulphatammon," $2\text{NH}_3, \text{SO}_3$.

(Rose.)

Is ammonium imidosulphonate, which see. (Berglund.)

"Parasulphatammon," $3\text{NH}_3, 2\text{SO}_3$.

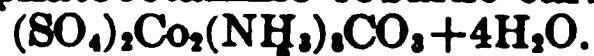
(Rose.)

Is basic ammonium imidosulphonate, which see. (Berglund.)

Sulphatiodic acid.**Potassium sulphatiodate, $\text{K}_2\text{HO}_3\text{SiO}_4$ or $\text{KIO}_3, \text{KHSO}_4$.**

Decomp. by H_2O . (Blomstrand, J. pr. (2) 40. 317.)

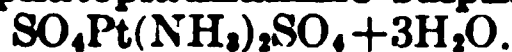
See Iodate sulphate, potassium.

Sulphatooctamine cobaltic carbonate

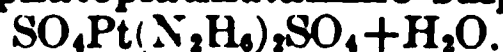
Sol. in H_2C . (Vortmann and Blasberg, B. 22. 2650.)

$(\text{SO}_4)\text{Co}_2(\text{NH}_3)_4(\text{CO}_3)_2 + 3\text{H}_2\text{O}$. Sol. in H_2O . (V. and B.)

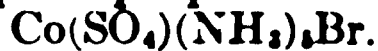
See Carbonatotetramine cobaltic sulphate. (Jørgensen.)

Sulphatoplatinamine sulphate,

Easily sol. in H_2O . Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$.

Sulphatoplatindiamine sulphate,

Insol. in H_2O .

Sulphatopurpureocobaltic bromide,

Sol. in H_2O , from which it is precipitated by conc. $\text{HBr} + \text{Aq}$. (Jørgensen, J. pr. (2) 25. 94.)

— carbonate, $[(\text{SO}_4)\text{Co}(\text{NH}_3)_4]_2\text{CO}_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Vortmann and Blasberg, B. 22. 2648.)

— chloroplatinate, $2\text{Co}(\text{SO}_4)(\text{NH}_3)_4\text{Cl}$, $\text{PtCl}_4 + 2\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . (Jørgensen.)

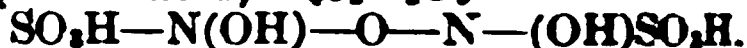
— nitrate, $\text{Co}(\text{SO}_4)(\text{NH}_3)_4(\text{NO}_3)$.

Somewhat sl. sol. in cold H_2O . (Jørgensen.)

Sulphatopurpureocobaltic sulphate,

Very easily sol. in H_2O . (Jørgensen, J. pr. (2) 25. 94.)

$\text{Co}(\text{SO}_4)(\text{NH}_3)_4(\text{HSO}_4) + 2\text{H}_2\text{O}$. Sol. in about 25 pts. of cold H_2O . Sol. in dil., insol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. (Jørgensen.)

Sulphazic acid, $\text{H}_2\text{S}_2\text{N}_2\text{O}_6 =$ 

Known only in its salts. (Raschig, A. 241. 161.)

Potassium sulphazate, $\text{K}_2\text{HS}_2\text{N}_2\text{O}_6 =$ 

Sol. in H_2O , but decomp. on standing. (Raschig, A. 241. 161.)

Sulphazidic acid.

(Fremy.)

See Hydroxylamine monosulphonic acid.

Sulphazilinic acid.

See Oxysulphazotic acid.

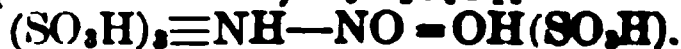
Metasulphazilinic acid.

See Trisulphoxyazotic acid.

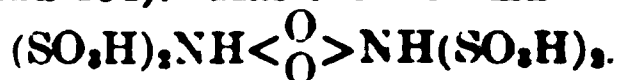
Sulphazinous acid.

(Fremy.)

See Dihydroxylamine sulphonic acid.

Sulphazotic acid, $\text{H}_2\text{N}_2\text{S}_2\text{O}_{14} =$ 

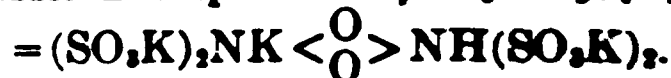
Known only in its salts. (Claus, A. 188. 52 and 194). Has the formula



(Raschig, A. 241. 161.)

Lead potassium sulphazotate.

Insol. in cold, decomp. by hot H_2O . Insol. in alcohol and ether. (Fremy, A. ch. (3) 15. 439.)

Potassium sulphazotate, $\text{K}_2\text{HN}_2\text{S}_2\text{O}_{14} + \text{H}_2\text{O}$ 

Very sol. in hot, less in cold H_2O . (Raschig, A. 241. 161.) Decomp. gradually by boiling. (Claus.) Insol. in alcohol or ether. (Fremy, A. ch. (3) 15. 428.)

True composition is $\text{HON}(\text{SO}_3\text{K})_2\text{KON}(\text{SO}_3\text{K})_2 + \text{H}_2\text{O}$. Potassium hydroxylamine disulphonate. (Divers and Haga, Chem. Soc. 1900, 77. 432.)

Forms basic salt

$(\text{SO}_3\text{K})_2\text{NK}<\overset{\text{O}}{\underset{\text{O}}{\parallel}}>\text{NK}(\text{SO}_3\text{K})_2$, which is easily sol. and decomp. by H_2O . (Raschig.)

Potassium sodium sulphazotate,

Quite easily sol. in H_2O . (Raschig, A. 241. 161.)

hydroxyazotic acid, $\text{ONH}(\text{SO}_2\text{H})_2$, only in its salts. (Claus, A. 158.)	Cuprous lead sulphobismuthite, $\text{Cu}_2\text{S}, 2\text{PbS}, \text{Bi}_2\text{S}_3$. Min. <i>Patrinite</i> . Sol. in $\text{HNO}_3 + \text{Aq}$ with residue of S and PbSO_4 .
hydroxylamic acid.	Lead —, $2\text{PbS}, \text{Bi}_2\text{S}_3$. Min. <i>Cosakite</i> .
hydroxylamine monosulphonic acid.	$2\text{PbS}, 3\text{Bi}_2\text{S}_3$. Min. <i>Chiviatite</i> .
hydroxyazotic acid.	Potassium —, KBiS . Decomp. by H_2O .
hydroxylamine disulphonic acid.	Sol. in $\text{HCl} + \text{Aq}$. (Schneider, Pogg. 1869, 136. 464.)
1. Sulphides of the alkali metals are sol. in water; those of the alkali-earth metals are sol., and are decomp. upon solution in sulphide and hydroxide. Other sulphides are insol. in H_2O . For sulphide, see under the respective element.	<i>Metasulphoboric acid</i> , $\text{B}_2\text{S}_3\text{H}_2\text{S}$. Decomp. by H_2O and alcohol. 1 pt. is sol. in 5 pts. benzene 1 pt. is " " 5 " CS_2 Very sl. sol. in CS_2 at -20° . (Stock, B. 1901, 34. 401.)
Is, SO_2NH_2 . Isosulphamide.	Sulphocarbonic acid.
Amide, $\text{OHSO} \begin{smallmatrix} \text{NSO.OH} \\ \text{NSO.OH} \end{smallmatrix} \text{N}(\text{H})_2$. Sol. in methyl alcohol; sl. sol. in ether; insol. in chloroform and benzene. (Hantzsch, Z. 3440.)	Ammonium cuprous sulphocarbonate, CS_2CuNH_4 . This salt was formerly described as cuprio sulphocarbonate ammonia, $\text{CS}_2\text{Cu}, \text{NH}_3$. (Hofmann, B. 1903, 36. 1146.)
1 sulphimide, $\text{SO}_2\text{N}(\text{NH}_2)_2$. I_2O ; insol. in alcohol. (Traube.)	Cuprous potassium sulphocarbonate, CS_2CuK . Nearly insol. in cold H_2O . Somewhat sol. in hot H_2O , NaOH and $\text{NH}_4\text{OH} + \text{Aq}$. (Hofmann.)
—, $(\text{SO}_2\text{N})_2\text{Ba} + 2\text{H}_2\text{O}$. I_2O . (Traube.)	Cupric sulphocarbonate ammonia, $\text{CS}_2\text{Cu}, \text{NH}_3$. Very sl. sol. in strong $\text{NH}_4\text{OH} + \text{Aq}$; insol. in cold H_2O , sl. sol. in hot H_2O . (Hofmann, Z. anorg. 1897, 14. 295.) Is ammonium cuprous sulphocarbonate. (Hofmann, B. 1903, 36. 1146.)
—, SO_2NK . Sol. in H_2O .	Cuprous sulphocarbonate potassium cyanide, $\text{CS}_2\text{Cu}, 2\text{KCN} + 2\text{H}_2\text{O}$. Sol. in H_2O and dil. alkalies on warming. (Hofmann, B. 1903, 36. 1148.)
—, SO_2NAg . 100–600 pts. cold, more easily in H_2O . Sol. in acids.	Zinc sulphocarbonate ammonia, $\text{CS}_2\text{Zn}, 2\text{NH}_3$. Ppt. (Hofmann, Z. anorg. 1897, 14. 277.)
—, SO_2NNa . Sol. in H_2O .	Sulphochromic acid, $\text{H}_2\text{CrO}_4, \text{SO}_2$. (?) Sol. in H_2O . (Bolley, A. 56. 113.) $(\text{SO}_2)_2\text{Cr}_2\text{O}_7(\text{OH})_2$. Sol. in H_2O . All salts even alkali salts are insol. in H_2O . (Recours, Bull. Soc. 1896, (3) 15. 315.) $[\text{Cr}_2\text{O}_7(\text{OH})_4(\text{SO}_2)_2\text{O}_2]$, $\text{Cr}_2\text{O}(\text{OH})_4(\text{SO}_2)_2$, $(\text{OH})_2(\text{OH})_2$. Sol. in H_2O . (Wyrouboff, Bull. Soc. 1902, (3) 27. 721.)
Ammonium, " NH_3, SO_2 . Sulphamic acid.	
Sulphurous acid.	
Sulphobismuthite, AuBiS_2 . Min. <i>Klaprothite</i> . Completely insol. in $\text{HNO}_3 + \text{Aq}$. Min. <i>Wittichenite</i> . Sol. in $\text{HCl} + \text{HNO}_3 + \text{Aq}$.	

Chromium sulphochromate,

Ppt.; decomp. by boiling H_2O . (Wyrouboff, Bull. Soc. 1902, (3) 27. 720.)

Sulphochromous acid.**Ferrous sulphochromite, FeCr_2S_4 .**

Insol. in H_2O , and nearly so in $\text{HCl} + \text{Aq}$. (Gröger, W. A. B. 81, 2. 531.)

Manganous —, MnCr_2S_4 .

Insol. in H_2O and $\text{HCl} + \text{Aq}$. (Gröger.)

Potassium —, $\text{K}_2\text{Cr}_2\text{S}_4$.

Insol. in H_2O and in hot $\text{HCl} + \text{Aq}$.

Easily sol. in aqua regia. Slowly sol. in cold, rapidly sol. in hot dil. $\text{HNO}_3 + \text{Aq}$. (Milbauer, Z. anorg. 1904, 42. 443.)

$\text{K}_2\text{Cr}_2\text{S}_7$. Stable in the air; sol. in HNO_3 and aqua regia with decomp. (Schneider, J. pr. 1897, (3) 56. 407.)

Silver —, $\text{Ag}_2\text{Cr}_2\text{S}_4$.

Not attacked by $\text{HCl} + \text{Aq}$ even on heating. Decomp. by conc. HNO_3 . (Schneider, J. pr. 1897, (2) 56. 401.)

Sodium —, $\text{Na}_2\text{Cr}_2\text{S}_4$.

Insol. in H_2O . Sl. attacked by dil. HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in cold conc. HNO_3 or aqua regia. Sol. in hot dil. $\text{HNO}_3 + \text{Aq}$. (Gröger.)

Sol. in acids with decomp. (Schneider, J. pr. 1897, (3) 56. 415.)

Zinc —, ZnCr_2S_4 .

Insol. in H_2O ; sol. in traces in boiling conc. HCl or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$; sol. in $\text{HNO}_3 + \text{Aq}$. (Gröger, W. A. B. 81, 2. 531.)

Sulphocyanhydric acid, HSCN .

Sol. in H_2O .

Sat. $\text{HSCN} + \text{Aq}$ has sp. gr. = 1.022. (Porrett, 1814.) $\text{HSCN} + \text{Aq}$ containing 12.7% HSCN has sp. gr. 1.040 at 12.7°. (Hermes, Z. Ch. 1866. 417.)

Sulphocyanides.

Most sulphocyanides are sol. in H_2O , but Cu , Pb , Hg , and Ag sulphocyanides are insol.

Aluminum sulphocyanide, $\text{Al}(\text{SCN})_3$.

Known only in solution.

$\text{Al}(\text{SCN})_2(\text{OH})_4$. Known only in solution (Suida.)

Aluminum potassium sulphocyanide,

Very hygroscopic.

Sol. in H_2O and alcohol. (Rosenheim, Z. anorg. 1901, 27. 302.)

Ammonium sulphocyanide, NH_4SCN

Deliquescent, and very sol. in H_2O . 100 pts. H_2O dissolve 128.1 pts. and 162.2 pts. at 20°.

$\text{NH}_4\text{SCN} + \text{Aq}$ sat. at ord. temp. density of 1.138 and 100 cc. contains NH_4SCN . (Klason, J. pr. 1887, (2)

By dissolving 90 g. NH_4SCN in 90 cc. H_2O at 17°, the temp. falls to -12°. (Ch. 1866. 190.)

133 pts. $\text{NH}_4\text{SCN} + 100$ pts. H_2O lower the temp. 31.2°. (Rüdorff, B. 1866. 190.)

Sol. in liquid SO_2 . (Walden, B. 1864.)

Difficultly sol. in AsBr_3 . (Walsh, anorg. 1902, 29. 374.)

Very easily sol. in liquid NH_3 . (Am. Ch. J. 1898, 20. 826.)

Easily sol. in alcohol.

Easily sol. in acetone. (Krug and

Sl. sol. in benzonitrile. (Naum, 1914, 47. 1369.)

Sol. in methyl acetate. (Naum, 1909, 42. 3789.)

Difficultly sol. in ethyl acetate. (mann, B. 1910, 43. 314.)

Ammonium bismuth sulphocyanide, $(\text{NH}_4)_3\text{Bi}(\text{SCN})_6$.

As K salt. (Rosenheim and Vogt, Z. anorg. 1906, 48. 215.)

Ammonium cadmium sulphocyanide, $(\text{NH}_4)_2\text{Cd}(\text{SCN})_4 + 2\text{H}_2\text{O}$.

Somewhat deliquescent.

Melts in crystal H_2O at 25°.

Insol. in alcohol. (Grossmann, 1885. 2667.)

Ammonium cadmium molybdenyl cyanide, NH_4SCN , $\text{CdMo}(\text{OH})(\text{SCN})_3 + 3\text{H}_2\text{O}$.

(Maas and Sand, B. 1908, 41. 1513)

Ammonium cobaltous sulphocyanide, $(\text{NH}_4)_2\text{Co}(\text{SCN})_4$.

Decomp. in moist air.

Cannot be recryst. from H_2O . (Th. Z. anorg. 1901, 26. 109.)

+4 H_2O . Sol. in H_2O .

Sol. in methyl, ethyl and amyl in acetone and in ether + Aq .

Can be recryst. from H_2O or alcohol out decomp. (Rosenheim and C. anorg. 1901, 27. 289.)

Ammonium iron (ferric) sulphocyanide, $9\text{NH}_4\text{SCN}$, $\text{Fe}(\text{SCN})_3 + 4\text{H}_2\text{O}$.

Deliquescent, and sol. in H_2O . (K. Moraht, A. 260. 207.)

3 NH_4SCN , $\text{Fe}(\text{SCN})_3$. Extremely deliquescent.

Ammonium mercuric sulphocyanide, $2\text{NH}_4\text{SCN}$, $\text{Hg}(\text{SCN})_2$.

Easily sol. in H_2O . (Fleischer, 1885. 228.)

$\text{NH}_4\text{Hg}(\text{SCN})_3$. Insol. in cold; sol. in hot H_2O . (Rosenheim, Z. anorg. 1901, 27. 284.)

Ammonium molybdenyl sulphocyanide,
 $3\text{NH}_4\text{SCN}$, $\text{Mo}(\text{OH})(\text{SCN})_3 + 3\text{H}_2\text{O}$.
(Sand and Maas, B. 1907, 40. 4507.)

Ammonium nickel sulphocyanide,
 $(\text{NH}_4)_4\text{Ni}(\text{SCN})_6 + 4\text{H}_2\text{O}$.
Sol. in H_2O with decomp.
Sl. sol. in cold; easily sol. in hot alcohol.
(Rosenheim, Z. anorg. 1901, 27. 292.)

Ammonium silver sulphocyanide, NH_4SCN ,
 AgSCN .
Decomp. by H_2O .

Ammonium vanadium sulphocyanide,
 $\text{V}(\text{SCN})_3$, $3\text{NH}_4\text{SCN} + 4\text{H}_2\text{O}$.
Sol. in H_2O ; sol. in alcohol; sl. sol. in ether.
(Ciocci, Z. anorg. 1898, 19. 311.)

Ammonium vanadyl sulphocyanide,
 $(\text{NH}_4)_2\text{VO}(\text{SCN})_4 + 5\text{H}_2\text{O}$.
Sol. in H_2O , alcohol, ether, acetone, amyl alcohol and ethyl acetate. (Koppel, Z. anorg. 1903, 36. 290.)

Ammonium zinc sulphocyanide,
 $(\text{NH}_4)_2\text{Zn}(\text{SCN})_4 + 3\text{H}_2\text{O}$.
Easily sol. in H_2O and in alcohol. (Walden, Z. anorg. 1900, 23. 374.)
 $+ 4\text{H}_2\text{O}$. Easily sol. in cold H_2O , acetone, alcohols and ether. (Rosenheim and Huld-schinsky, B. 1901, 34. 3913.)

Ammonium sulphocyanide mercuric bromide,
 NH_4SCN , HgBr_2 .
Very sol. in H_2O .
Sol. in alcohol. (Grossmann, B. 1902, 35. 2945.)
 $2\text{NH}_4\text{SCN}$, $\text{HgBr}_2 + \text{H}_2\text{O}$. Somewhat deliquescent.
Very sol. in H_2O .
Sol. in alcohol. (Grossmann.)

Arsenic sulphocyanide, $\text{As}(\text{SCN})_3$.
Decomp. by H_2O . Insol. in all ordinary solvents. (Miguel, A. ch. (5) 11. 341.)

Barium sulphocyanide, $\text{Ba}(\text{SCN})_2 + 2\text{H}_2\text{O}$.
Deliquescent. Easily sol. in H_2O and alcohol. Boiling solution in alcohol contains 32.8% anhydrous salt. Solution sat. at 20° contains 30%. (Tscherniak, B. 16. 349.)
Cryst. with $3\text{H}_2\text{O}$. (Tscherniak, B. 25. 2627.)

Barium cadmium sulphocyanide,
 $4\text{Ba}(\text{SCN})_2$, $\text{Cd}(\text{SCN})_2 + 10\text{H}_2\text{O}$.
Deliquescent. (Grossmann, B. 1902, 36. 2669.)

Barium caesium cuprous sulphocyanide,
 $\text{Ba}(\text{SCN})_2$, 3CsSCN , 2CuSCN .
Rapidly decomp. by H_2O . (Wells, Am. Ch. J. 1902, 28. 273.)

Barium caesium silver sulphocyanide,
 $\text{Ba}(\text{SCN})_2$, 3CsSCN , 2AgSCN .
100 pts. H_2O dissolve 92 pts. at 19° .
Decomp. by much H_2O . (Wells, Am. Ch. J. 1902, 28. 272.)

Barium cobaltous sulphocyanide,
 $\text{BaCo}(\text{SCN})_4 + 8\text{H}_2\text{O}$.
Ppt. (Rosenheim, Z. anorg. 1901, 27. 290.)

Barium mercuric sulphocyanide,
 $\text{BaHg}(\text{SCN})_4$.
Very sol. in H_2O and in alcohol. (Rosenheim, Z. anorg. 1901, 27. 286.)
 $\text{BaHg}(\text{SCN})_4 + 2\text{H}_2\text{O}$. Ppt. Nearly insol. in cold; easily sol. in hot H_2O . (Rosenheim.)

Barium potassium silver sulphocyanide,
 $\text{Ba}(\text{SCN})_2$, 4KSCN , $2\text{AgSCN} + \text{H}_2\text{O}$.
Very sol. in a little H_2O . Decomp. by much H_2O . (Wells, Am. Ch. J. 1902, 28. 283.)

Barium rubidium silver sulphocyanide,
 $\text{BaRb}_2\text{Ag}_2(\text{SCN})_6 + 2\text{H}_2\text{O}$.
Very sol. in H_2O . (Wells, Am. Ch. J. 1903, 30. 186.)
 $\text{BaRb}_4\text{Ag}_2(\text{SCN})_8 + \text{H}_2\text{O}$. Sol. in H_2O . (Wells.)

Barium silver sulphocyanide, $\text{Ba}(\text{SCN})_2$,
 $2\text{AgSCN} + 2\text{H}_2\text{O}$.
Stable in the air. (Wells, Am. Ch. J. 1902, 28. 269.)

Barium zinc sulphocyanide, $\text{BaZn}(\text{SCN})_4$,
 $+ 3\text{H}_2\text{O}$.
Easily sol. in alcohol. (Walden, Z. anorg. 1900, 23. 374.)

Barium sulphocyanide mercuric bromide,
 $\text{Ba}(\text{SCN})_2$, $2\text{HgBr}_2 + 5\text{H}_2\text{O}$.
Very sol. in H_2O . (Grossmann, Z. anorg. 1903, 37. 420.)

Bismuth sulphocyanide, basic,
 $\text{Bi}(\text{OH})(\text{SCN})_2 + 5\text{H}_2\text{O}$.
(Rosenheim and Vogelgesang, Z. anorg. 1906, 48. 214.)
 $\text{Bi}(\text{SCN})_3$, $2\text{Bi}_2\text{O}_3$. Insol. in H_2O , but when recently pptd. decomp. by boiling therewith. Insol. in $\text{HSCN} + \text{Aq}$. (Meitzendorf.)

Bismuth sulphocyanide, $\text{Bi}(\text{SCN})_3$.
Insol. or sl. sol. in H_2O . Sol. in HNO_3 , HCl , and $\text{HSCN} + \text{Aq}$. (Meitzendorf, Pogg. 56. 83.)

Decomp. by cold H_2O . (Bender, B. 20 723.)

+14 H_2O . Extremely deliquescent.

Decomp. by H_2O . (Rosenheim and Vogelgesang, Z. anorg. 1906, 48. 214.)

Bismuth potassium sulphocyanide,
 $K_3Bi(SCN)_6$.

Decomp. by H_2O . (Rosenheim and Vogelgesang, Z. anorg. 1906, 48. 215.)

Not hygroscopic.

Decomp. by H_2O .

Easily sol. in alcohol. (Vanino, Z. anorg. 1901, 28. 220.)

$Bi(SCN)_3$, 9 $KSCN$. Very hygroscopic.

Decomp. by H_2O .

Sol. in alcohol. (Vanino, Z. anorg. 1901, 28. 221.)

Bismuth sodium sulphocyanide,
 $Na_3Bi(SCN)_6$.

As K salt. (Rosenheim and Vogelgesang, Z. anorg. 1906, 48. 215.)

Boron sulphocyanide, $B(SCN)_3$.

Sol. in benzene and ether. (Cocksedge, Chem. Soc. 1908, (2) 93. 217.)

Cadmium sulphocyanide, $Cd(SCN)_2$.

Sl. sol. in H_2O . Sol. in NH_4OH + Aq with combination.

Cadmium caesium sulphocyanide,
 $CsCd(SCN)_3$.

Recryst. from H_2O . (Wells, Am. Ch. J. 1903, 30. 148.)

$Cs_4Cd(SCN)_6 + 2H_2O$. Very sol. in H_2O . Can be recryst. from conc. solution but decomp. on dilution to $CsCd(SCN)_3$. (Wells.)

Cadmium caesium silver sulphocyanide,
 $Cs_2CdAg_2(SCN)_6$.

(Wells.)

+2 H_2O . (Wells.)

$(Cs_2CdAg_4(SCN)_8 + 2H_2O)$. (Wells.)

$Cs_4Cd_3Ag_{10}(SCN)_{21} + 6H_2O$. (Wells.)

Cadmium mercuric sulphocyanide, $Cd(SCN)_2$,
 $Hg(SCN)_2$.

Very sol. in hot H_2O . (Grossmann, Z. anorg. 1903, 37. 414.)

Cadmium molybdenum sulphocyanide,
 $Cd(SCN)_2$, $Mo(SCN)_4 + 2H_2O$.

(Maas and Sand, B. 1908, 41. 1513.)

+3 H_2O . (Maas and Sand.)

Cadmium molybdenyl potassium sulphocyanide, $KSCN$, $4Cd(SCN)_2$,
 $3Mo(OH)(SCN)_3 + 18H_2O$.

(Maas and Sand, B. 1908, 41. 1513.)

Cadmium molybdenyl sulphocyanide ammonia, $3Cd(SCN)_2$, $2Mo(OH)_3$,
 $13NH_3$.

(Maas and Sand, B. 1908, 41. 1512.)

+2 H_2O . (Maas and Sand.)

Cadmium potassium sulphocyanide,
 $K_2Cd(SCN)_4 + 2H_2O$.

Very sol. in H_2O . (Grossmann, B. 35. 2668.)

Cadmium rubidium sulphocyanide,
 $Rb_2Cd(SCN)_4 + 2H_2O$.

Very sol. in H_2O . (Grossmann, B. 35. 2668.)

Cadmium sodium sulphocyanide,
 $NaCd(SCN)_3 + 3H_2O$.

(Grossmann, B. 1902, 35. 2668.)

Cadmium sulphocyanide ammonia, $Cd(SCN)_2$,
 NH_3 .

Decomp. by H_2O . (Grossmann, B. 35. 2666.)

$Cd(SCN)_2$, 2 NH_3 . Decomp. by pur (Grossmann.)

Cadmium sulphocyanide ammonium bromide,
 $Cd(SCN)_2$, $NH_4Br + H_2O$.

Can be recryst. from H_2O . Deco dil. solution. (Grossmann, Z. anorg. 37. 425.)

$Cd(SCN)_2$, 2 NH_4Br . Easily spl NH_4Br . (Grossmann.)

Cadmium sulphocyanide ammonium chloride,
 $Cd(SCN)_2$, 2 NH_4Cl .

Can be recryst from H_2O . Decomp. solution. (Grossmann, Z. anorg. 1904, 423.)

Cadmium sulphocyanide potassium bromide,
 $Cd(SCN)_2$, $KBr + H_2O$.

Recryst. from H_2O . (Grossmann, Z. 1903, 37. 425.)

$Cd(SCN)_2$, 2 KBr . Recryst. from (Grossmann.)

Cadmium sulphocyanide potassium chloride,
 $Cd(SCN)_2$, 2 KCl .

Recryst. from H_2O . Decomp. in dilution. (Grossmann, Z. anorg. 1903, 37.)

Cadmium sulphocyanide potassium iodide,
 $Cd(SCN)_2$, 2 KI .

Recryst. from H_2O . (Grossmann.)

Cæsium calcium silver sulphocyanide, $2CsSCN$, $Ca(SCN)_2$, $2AgSCN + 2H_2O$.

Recryst. from H_2O . (Wells, Am. Ch. J. 1902, 28. 275.)

Cæsium chromium sulphocyanide.

See Chromisulphocyanide, cæsium.

Cæsium cobaltous sulphocyanide,

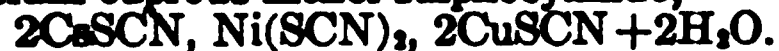
Stable in the air. (Shinn and Wells, Am. Ch. J. 1903, 29. 476.)

Cæsium cobaltous silver sulphocyanide,

Slowly attacked by H_2O ; decomp. by boiling H_2O . Very sl. sol. in CsSCN or $\text{Co}(\text{SCN})_2$ + Aq. (Shinn and Wells, Am. Ch. J. 1903, 29. 478.)

Cæsium cuprous sulphocyanide, CsSCN , CuSCN .

H_2O separates CuSCN . (Roberts, Am. Ch. J. 1902, 28. 262.)

Cæsium cuprous nickel sulphocyanide,

Sl. sol. in H_2O . (Roberts and Wells, Am. Ch. J. 1902, 28. 277.)

Cæsium cuprous strontium sulphocyanide,

As Ba salt. (Wells, Am. Ch. J. 1902, 28. 275.)

Cæsium magnesium silver sulphocyanide,

As Ca comp. (Wells, Am. Ch. J. 1902, 28. 275.)

Cæsium manganous silver sulphocyanide,

Rather sl. sol. in H_2O . (Wells.)

Cæsium mercuric sulphocyanide, CsSCN , $\text{Hg}(\text{SCN})_2$.

Sl. sol. in hot H_2O . (Bristol and Wells, Am. Ch. J. 1902, 28. 260.)

Cæsium mercuric sulphocyanide, 2CsSCN , $\text{Hg}(\text{SCN})_2 + \text{H}_2\text{O}$.

Moderately sol. in H_2O , especially when warm. Recryst. without decomp. (Bristol and Wells, Am. Ch. J. 1902, 28. 260.)

Cæsium nickel silver sulphocyanide,

Slowly decomp. by hot H_2O . (Wells, Am. Ch. J. 1902, 28. 277.)

Cæsium silver sulphocyanide, CsSCN , AgSCN .

Easily forms supersat. solution. (Wells, Am. Ch. J. 1902, 28. 264.)

2CsSCN , AgSCN . Stable in the air. (Wells.)

3CsSCN , AgSCN . Stable in the air. (Wells.)

Cæsium silver strontium sulphocyanide,

As Ba comp. (Wells.)

Cæsium silver zinc sulphocyanide,

(Wells.)

$\text{Cs}_2\text{ZnAg}(\text{SCN})_6$. Ppt. Stable in the air. (Wells.)

$\text{CsZn}_2\text{Ag}_2(\text{SCN})_8$. Decomp. by cold, more rapidly by hot H_2O . (Wells.)

$\text{CsZn}_2\text{Ag}_4(\text{SCN})_{10}$. Slowly decomp. by H_2O . (Wells.)

Cæsium zinc sulphocyanide, $\text{Cs}_2\text{Zn}(\text{SCN})_4 + 2\text{H}_2\text{O}$.

Moderately sol. in H_2O and can be recryst. therefrom. (Wells.)

Calcium sulphocyanide, $\text{Ca}(\text{SCN})_2 + 3\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O and alcohol.

Calcium silver sulphocyanide, $\text{Ca}(\text{SCN})_2$, $2\text{AgSCN} + 2\text{H}_2\text{O}$.

(Wells.)

Calcium stannic sulphocyanide, $\text{CaSn}(\text{SCN})_6 + 7\text{H}_2\text{O}$.

Very sol. in H_2O . Can be recryst. therefrom. Sol. in alcohol and acetone. (Weinland and Bames, Z. anorg. 1909, 62. 258.)

Cerous sulphocyanide, $\text{Ce}(\text{SCN})_3 + 7\text{H}_2\text{O}$.

Deliquescent. Sol. in H_2O and alcohol. (Jolin, Bull. Soc. (2) 21. 534.)

Chromous sulphocyanide with MSCN .

See Chromosulphocyanide, M.

Chromic sulphocyanide, $\text{Cr}(\text{SCN})_3$.

Deliquescent, and sol. in H_2O .

Somewhat sol. in organic solvents. (Speransky, C. C. 1897, I. 141.)

See also Chromisulphocyanhydric acid.

Chromic sulphocyanide with MSCN .

See Chromisulphocyanide, M.

Cobaltous sulphocyanide, $\text{Co}(\text{SCN})_2 + \frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O and alcohol; also in ether. Sol. in liquid SO_2 . (Walden, B. 1899, 32. 2864.)

Sol. in acetone. (Krug and M'Elroy.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

+ $3\text{H}_2\text{O}$. Sol. in H_2O and in alcohol. (Rosenheim and Cohn, Z. anorg. 1901, 27. 288.)

Cobaltous mercuric sulphocyanide, $\text{Co}(\text{SCN})_2$, $\text{Hg}(\text{SCN})_2$.

Very sl. sol. in H_2O and dil. HCl + Aq. Easily sol. in HNO_3 + Aq. (Cleve, J. pr. 91. 227.)

Cobaltous potassium sulphocyanide,
 $\text{CoK}_2(\text{SCN})_4$.Decomp. by H_2O . (Treadwell, Z. anorg. 1901, 26. 109.)+4 H_2O . Sol. in H_2O . Sol. in methyl, ethyl and amyl alcohol, in acetone and in ether+Aq. Can be recryst. from H_2O or alcohol without decomp. (Rosenheim and Cohn, Z. anorg. 1901, 27. 289.)**Cobaltous silver sulphocyanide, $\text{CoAg}(\text{SCN})_2$, +2 H_2O .**Decomp. by H_2O . (Shinn and Wells, Am. Ch. J. 1903, 29. 476.) $\text{Ag}_2\text{Co}(\text{SCN})_4$. Almost insol. in H_2O and in alcohol. (Rosenheim, Z. anorg. 1901, 27. 291.)**Cobaltous sodium sulphocyanide,**
 $\text{Na}_2\text{Co}(\text{SCN})_4 + 8\text{H}_2\text{O}$.Sol. in H_2O and in alcohol. (Rosenheim.)**Cobaltous sulphocyanideammonia, $\text{Co}(\text{SCN})_2$, 2 NH_3 , and $\text{Co}(\text{SCN})_2$, 6 NH_3 .**

(Peters, B. 1908, 41. 3178.)

 $\text{Co}(\text{SCN})_2, 4\text{NH}_3$. Sol. in H_2O and alcohol. (Sand, B. 1903, 36. 1439.)**Cobaltous sulphocyanide mercuric chloride,**
 $2\text{Co}(\text{SCN})_2, 2\text{HgCl}_2$.

(Hantzsch and Shibata, Z. anorg. 1912, 73. 320.)

 $2\text{Co}(\text{SCN})_2, 3\text{HgCl}_2$. Easily decomp. (Hantzsch and Shibata.)**Cuprous sulphocyanide, CuSCN .**1 l. H_2O at 18° dissolves 0.004 mg. mol. or 0.5 mg. CuSCN . (Kohlrausch and Ross, Z. phys. Ch. 1893, 12. 241.)Insol. in dil. acids. Sl. sol. in cold, easily in warm conc. HCl +Aq. Decomp. by conc. H_2SO_4 or HNO_3 +Aq. Sol. with combination in NH_4OH +Aq. Insol. in KSCN +Aq.Less sol. in H_2SO_4 and H_2SO_4 +Aq than in HNO_3 . (Kuhn, Ch. Z. 1908, 32. 1056.)Sol. in $\text{Fe}_2(\text{SO}_4)_3$ +Aq. (Johnson, J. Soc. Chem. Ind. 1889, 8. 603.) KSCN +Aq (85–90 g. in 50 g. H_2O) dissolves 18 g. CuSCN . (Thurnauer, B. 1899, 23. 770.)

Sol. in ether. (Skey, C. N. 1867, 16. 201.)

Cupric sulphocyanide, $\text{Cu}(\text{SCN})_2$.Decomp. by H_2O to cuprous salt. Sol. in warm HCl , H_2SO_4 , or HNO_3 +Aq. Sol. in MSCN +Aq, but solutions decomp. by dilution. Sol. in NH_4OH +Aq.Solubility in NH_4OH +Aq at 25° and at 40° .At 25° .

Sp. gr. $25^\circ/25^\circ$	One gram of solution contains			1000 mols. H_2O dissolve		Composition of solid salt in contact with solution
	g. NH_3	g. $\text{Cu}(\text{SCN})_2$	g. H_2O	Mols. NH_3	Mols. $\text{Cu}(\text{SCN})_2$	
0.99853	0.2147	0.1522	0.6331	358.04	24.09	} $\text{Cu}(\text{SCN})_2, 4\text{NH}_3$
0.99871	0.1655	0.1124	0.7221	242.02	15.60	
1.00703	0.0993	0.0798	0.8209	127.76	9.74	
1.01336	0.0639	0.0659	0.8702	77.51	7.59	
1.01506	0.0535	0.0622	0.8843	64.05	7.04	
1.01705	0.0426	0.0596	0.8978	50.21	6.65	
1.02132	0.0250	0.0511	0.9239	28.55	5.55	} $\text{Cu}(\text{SCN})_2, 2\text{NH}_3$
1.01661	0.0198	0.0408	0.9394	22.27	4.35	
1.00816	0.0079	0.0245	0.9676	18.61	2.54	

At 40° .

	0.1802	0.1976	0.6222	306.28	31.83	} $\text{Cu}(\text{SCN})_2, 4\text{NH}_3$
	0.1398	0.1658	0.6944	213.10	23.93	
	0.0758	0.1299	0.7943	101.00	16.38	
	0.0550	0.1207	0.8243	70.59	14.67	
	0.0435	0.1178	0.8388	54.82	14.07	
	0.0352	0.0876	0.8772	42.53	10.00	
	0.0257	0.0655	0.9088	30.00	7.22	} $\text{Cu}(\text{SCN})_2, 2\text{NH}_3$
	0.0177	0.0418	0.9405	19.86	4.46	
	0.0094	0.0281	0.9625	10.31	2.93	

(Horn, Am. Ch. J. 1907, 37. 471.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Cuprocupric sulphocyanide, $\text{Cu}(\text{SCN})_2$, $\text{Cu}_2(\text{SCN})_2$.Not attacked by hot HCl +Aq. Insol. in KSCN +Aq.

- mercuric sulphocyanide**, $\text{CuHg}(\text{SCN})_4$.
 Insol. in cold H_2O and in alcohol;
 in boiling H_2O . (Rosenheim, Z. anorg.
 7. 288.)
- potassium sulphocyanide**, CuSCN ,
 SCN .
 Lustrous. Decomp. by H_2O . (Thurn-
 1890, 23. 770.)
- potassium sulphocyanide ammonia**, $\text{Cu}_2(\text{SCN})_2$,
 H_2 .
 mp. in the air. (Richards, Z. anorg.
 7. 247.)
 $3\text{SCN})_2$, 5NH_3 . Very unstable in the
 air. (Richards.)
- potassium sulphocyanide ammonia**, $\text{Cu}(\text{SCN})_2$,
 H_2 .
 Insol. in little H_2O , but decomp. by dilution
 into a basic salt. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
 Long standing a small amount dissolves
 with separation of CuSCN . (Litter-
 Arch. Pharm. 1901, 239. 337.)
 Sol. in H_2O . Sol. in H_2O containing
 a small amount of ammonia. (Horn, Am.
 1907, 37. 477.)
 $(\text{SCN})_2$, 4NH_3 . Very unstable in the
 air.
- in H_2O , but decomp. by much H_2O
 into a basic salt. (Horn.)
 pts. N/10 $\text{NH}_4\text{OH} + \text{Aq}$ dissolve 10.4
 hydrous salt at 25° . (Pudschies, Dis-
 1907, 37. 477.)
- potassium sulphocyanide ammonia**, $\text{Cu}(\text{SCN})_2$,
 H_2 .
 mp. in the air. (Kohlestrücker, B. 1904, 37.
 1907, 37. 477.)
- mp. in the air and by H_2O and dil.
 conc. acids; sol. in cold conc. HNO_3 and
 $\text{HCl} + \text{Aq}$. Sol. in boiling conc. HCl .
 (Richards, Z. anorg. 1898, 17. 250.)
- potassium sulphocyanide**, $\text{Di}(\text{SCN})_2 + 6\text{H}_2\text{O}$.
 Lustrous, and sol. in H_2O .
- potassium sulphocyanide**, $\text{Er}(\text{SCN})_3 + 6\text{H}_2\text{O}$.
 Lustrous. Sol. in H_2O . (Höglund.)
- potassium sulphocyanide**, $\text{Gl}(\text{SCN})_2$ (?).
 in H_2O . (Hermes, J. pr. 97. 465.)
- aurous) potassium sulphocyanide**,
 2SCN , KSCN .
 Sol. in H_2O , less in absolute alcohol.
 (J. pr. 94. 16.)
- (aurous) potassium sulphocyanide**
ammonia, $\text{KAu}(\text{SCN})_2$, 5NH_3 .
 mp. in the air. (Hermes, B. 1908, 41. 3178.)
- (auric) potassium sulphocyanide am-**
monia, $\text{KAu}(\text{SCN})_4$, 4NH_3 .
 mp. in the air. (Hermes.)
- Gold (aurous) silver sulphocyanide**, AuSCN ,
 AgSCN .
 Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.
- Gold (auric) potassium sulphocyanide**.
 Sol. in H_2O , alcohol, and ether. (Cleve.)
- Gold (aurous) sulphocyanide ammonia**,
 AuSCN , NH_3 .
 Very sol. in cold, decomp. by hot H_2O .
- Iron (ferrous) sulphocyanide**, $\text{Fe}(\text{SCN})_2 + 3\text{H}_2\text{O}$.
 Very sol. in H_2O , alcohol, or ether.
 Sol. in acetone. (Krug and M'Elroy.)
- Iron (ferric) sulphocyanide**, $\text{Fe}(\text{SCN})_3 + 3\text{H}_2\text{O}$.
 Deliquescent. Very sol. in H_2O , alcohol,
 or ether. Ether extracts the salt from
 $\text{Fe}(\text{SCN})_2 + \text{Aq}$. Decomp. by much H_2O
 if pure. Not decomp. by monobasic acids,
 but conc. H_2SO_4 and H_3PO_4 , also oxalic,
 tartaric, malic, etc., acids destroy the colour.
- Iron (ferric) lithium sulphocyanide**, $\text{Fe}(\text{SCN})_3$,
 $9\text{LiSCN} + 4\text{H}_2\text{O}$.
 More deliquescent than the other ferric
 sulphocyanides. (Krüss and Moraht.)
- Iron (ferrous) mercuric sulphocyanide**,
 $\text{Fe}(\text{SCN})_2$, $\text{Hg}(\text{SCN})_2 + 2\text{H}_2\text{O}$.
 Moderately sol. in hot H_2O . (Cleve, J.
 pr. 91. 227.)
- Iron (ferric) potassium sulphocyanide**,
 $\text{Fe}(\text{SCN})_3$, $3\text{KSCN} + x\text{H}_2\text{O}$.
 Extremely deliquescent, and sol. in H_2O .
 (Krüss and Moraht.)
 $\text{Fe}(\text{SCN})_3$, $9\text{KSCN} + 4\text{H}_2\text{O}$. Hygroscopic.
 Sol. in H_2O without decomp. Insol. in pure
 anhydrous ether, but decomp. by ether con-
 taining traces of H_2O into $\text{Fe}(\text{SCN})_2$ and
 KSCN . (Krüss and Moraht, A. 260. 204.)
- Iron (ferrous) sodium sulphocyanide**,
 $\text{Na}_4\text{Fe}(\text{SCN})_6 + 12\text{H}_2\text{O}$.
 Sol. in H_2O and alcohol. (Rosenheim, Z.
 anorg. 1901, 27. 299.)
- Iron (ferric) sodium sulphocyanide**, $\text{Fe}(\text{SCN})_3$,
 $9\text{NaSCN} + 4\text{H}_2\text{O}$.
 Less deliquescent than the corresponding
 NH_4 or K salt. (Krüss and Moraht.)
 $\text{Na}_4\text{Fe}(\text{SCN})_6 + 12\text{H}_2\text{O}$. (Rosenheim, Z.
 anorg. 1901, 27. 297.)
- Lanthanum sulphocyanide**, $\text{La}(\text{SCN})_3 + 7\text{H}_2\text{O}$.
 Deliquescent; sol. in H_2O . (Cleve.)
- Lead sulphocyanide, basic**,
 6PbO , $\text{Pb}(\text{SCN})_2 + 2\text{H}_2\text{O}$. Ppt.
 $\text{Pb}(\text{SCN})_2$, $\text{PbO} + \text{H}_2\text{O}$. Insol. in H_2O .
 (Strömholm, Z. anorg. 1904, 33. 440.)

Lead sulphocyanide, $\text{Pb}(\text{SCN})_2$.

Nearly insol. in cold, decomp. by boiling H_2O . (Liebig.)

Sl. sol. in H_2O .

4.5×10^{-1} g. are dissolved in 1 liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

Lead sulphocyanide bromide, $\text{Pb}(\text{SCN})_2 \cdot 8\text{PbBr}_2$.

(Grissom and Thorp, Am. Ch. J. 10. 219.)

Lead sulphocyanide chloride, PbSCNCl .

Sl. sol. in cold, easily sol. in hot H_2O . (Murtry, Chem. Soc. 55. 50.)

Sol. in H_2O . (Grissom and Thorp, Am. Ch. J. 10. 229.)

Lead sulphocyanide iodide, $3\text{Pb}(\text{SCN})_2 \cdot \text{PbI}_2$.

Sol. in H_2O . (Grissom and Thorp, Am. Ch. J. 10. 229.)

Lithium sulphocyanide, LiSCN .

Very deliquescent. Sol. in H_2O and alcohol. (Hermes, Z. Ch. 1866. 417.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3789.)

Magnesium sulphocyanide, $\text{Mg}(\text{SCN})_2 \cdot 4\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O and alcohol.

Magnesium stannic sulphocyanide, $\text{MgSn}(\text{SCN})_6 \cdot 6\text{H}_2\text{O}$.

Hygroscopic. Sol. in H_2O , alcohol and acetone. (Weinland and Bames, Z. anorg. 1909, 62. 258.)

Manganous sulphocyanide, $\text{Mn}(\text{SCN})_2 \cdot 3\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O and alcohol.

Mercurous sulphocyanide, $\text{Hg}_2(\text{SCN})_2$.

Insol. in H_2O . Sol. in hot $\text{HCl} + \text{Aq}$. Slowly decomp. by hot aqua regia. Sol. in hot $\text{KSCN} + \text{Aq}$.

Mercuric sulphocyanide, basic, $\text{Hg}(\text{SCN})_2 \cdot 3\text{HgO}$.

Insol. in H_2O . Easily sol. in $\text{HCl} + \text{Aq}$. Insol. in H_2SO_4 or $\text{HNO}_3 + \text{Aq}$. (Fleischer.)

$\text{Hg}(\text{SCN})_2 \cdot 2\text{HgO}$. Insol. in H_2O . Sl. attacked by acids. (Claus, J. pr. 15. 401.)

Mercuric sulphocyanide, $\text{Hg}(\text{SCN})_2$.

Very sl. sol. in cold, much more easily in hot H_2O . Easily sol. in dil. $\text{HCl} + \text{Aq}$. (Crookes, Chem. Soc. 4. 18.)

Solubility in $\text{H}_2\text{O} = 0.00218$ mol. in 1 l. (Grossmann, Z. anorg. 1904, 43. 358.)

More sol. in H_2O than in alcohol. (Peters, B. 1908, 41. 3180.)

Very sl. sol. in H_2O at 25° . Appreci sol. only in boiling H_2O . (Jander, D. 1902.)

Sol. in $\text{Hg}(\text{NO}_3)_2$ or $\text{KSCN} + \text{Aq}$, also $\text{NH}_4\text{Cl} + \text{Aq}$. Sol. in many sulphocyanides + Aq .

Easily sol. in cold HCl , NH_4Cl , KCl , $\text{BaCl}_2 + \text{Aq}$. (Hermes, J. pr. 1866, 1. 477.)

Very sol. in liquid NH_3 . (Franklin, Ch. J. 1898, 20. 829.)

Sl. sol. in benzonitrile. (Naumann, 1914, 47. 1369.)

Mercuric hydrogen sulphocyanide, $\text{Hg}(\text{SCN})_2 \cdot 2\text{HSCN}$.

Easily decomp. (Hermes, Dissert. 1)

Mercuric nickel sulphocyanide, $\text{Hg}(\text{SCN})_2 \cdot \text{Ni}(\text{SCN})_2 \cdot 2\text{H}_2\text{O}$.

Moderately sol. in hot H_2O . (Cleve, pr. 91. 227.)

Very sol. in $\text{MSCN} + \text{Aq}$. (Orloff, 1906, I. 1411.)

Mercuric potassium sulphocyanide, $\text{Hg}(\text{SCN})_2 \cdot \text{KSCN}$.

Sol. in cold, more easily in hot H_2O . in alcohol and ether. Very sol. in $\text{NH}_4\text{KCl} + \text{Aq}$. (Claus.)

$\text{K}_2\text{Hg}(\text{SCN})_4$. Very sol. in H_2O ; also in alcohol.

Insol. in anhydrous ether. (Rosen, Z. anorg. 1901, 27. 285.)

Mercuric rubidium sulphocyanide, $\text{Hg}(\text{SCN})_2 \cdot \text{RbSCN}$.

Sol. in alcohol without decomp. Dec. by H_2O .

$\text{Hg}(\text{SCN})_2 \cdot 2\text{RbSCN} + \frac{1}{2}\text{H}_2\text{O}$. Easily in H_2O without decomp. (Grossmann, 1904, 37. 1259.)

Mercuric sodium sulphocyanide, $\text{Na}_2\text{Hg}(\text{SCN})_2$.

Very hygroscopic. (Rosenheim, Z. anorg. 1901, 27. 286.)

Mercuric zinc sulphocyanide, $\text{Hg}(\text{SCN})_2 \cdot \text{Zn}(\text{SCN})_2$.

Scarcely sol. in cold H_2O . Easily in $\text{HCl} + \text{Aq}$. (Cleve.)

Mercuric sulphocyanide ammonia, $2\text{Hg}(\text{SCN})_2 \cdot 3\text{NH}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Decomp. by H_2O and alcohol.

$\text{Hg}(\text{SCN})_2 \cdot 4\text{NH}_3$. (Peters, B. 1908, 3178.)

Mercuric sulphocyanide ammonium bromide, $\text{Hg}(\text{SCN})_2 \cdot \text{NH}_4\text{Br}$.

Decomp. by H_2O . Sol. in alcohol. (Grossmann, Z. anorg. 1903, 37. 418.)

Mercuric sulphocyanide ammonium chloride,
 $\text{Hg}(\text{SCN})_2, \text{NH}_4\text{Cl}$.

Decomp. by H_2O . Sol. in warm alcohol from which it can be cryst. (Grossmann.)

Mercuric sulphocyanide bromide, HgSCNBr .

Insol. in cold H_2O ; sol. in hot H_2O and in alcohol. (Rosenheim, Z. anorg. 1901, 27. 282.)

Mercuric sulphocyanide chloride, HgSCNCl .

Insol. in cold H_2O .

Sol. in hot H_2O and alcohol. (Rosenheim.)

Mercuric sulphocyanide potassium bromide,
 $\text{Hg}(\text{SCN})_2, 2\text{KBr}$.

Very sol. in H_2O . (Grossmann, Z. anorg. 1903, 37. 418.)

Mercuric sulphocyanide potassium chloride,
 $\text{Hg}(\text{SCN})_2, \text{KCl}$.

Decomp. by H_2O .

Not decomp. by recryst. from warm alcohol (Grossmann.)

Molybdenum sulphocyanide, $\text{Mo}(\text{SCN})_2(?)$.

Sol. in H_2O and ether. (Braun, Z. anal. 6. 86.)

Molybdenum potassium sulphocyanide,
 $\text{K}_2\text{Mo}(\text{SCN})_6 + 4\text{H}_2\text{O}$.

Cryst. from boiling H_2O and alcohol. (Chilesotti, Gazz. ch. it. 1904, 34. (2) 493.)

Molybdenum sodium sulphocyanide,
 $\text{Na}_2\text{Mo}(\text{SCN})_6 + 12\text{H}_2\text{O}$.

(Rosenheim, B. 1909, 42. 154.)

Molybdenum thalious sulphocyanide,
 $\text{MoTl}_3(\text{SCN})_6$.

(Rosenheim and Garfunkel, B. 1908, 41. 2388.)

Molybdenum sulphocyanide zinc amine,
 $2\text{Mo}(\text{SCN})_6, 3\text{Zn}(\text{NH}_3)_4$.

(Rosenheim and Garfunkel, B. 1908, 41. 2390.)

$2\text{Mo}(\text{SCN})_6(\text{OH}), \text{Zn}_3(\text{NH}_3)_{11}$. Can be cryst. from boiling $\text{NH}_4\text{OH} + \text{Aq}$. Air-dried salt probably has the composition $2\text{Mo}(\text{SCN})_6(\text{OH}), 3\text{Zn}(\text{NH}_3)_4 + 2\text{H}_2\text{O}$. (Maas and Sand, B. 1908, 41. 1510.)

$2\text{Mo}(\text{SCN})_6(\text{OH}), \text{Zn}_3(\text{NH}_3)_{11}$. (Maas and Sand.)

Nickel sulphocyanide, $\text{Ni}(\text{SCN})_2$.

Sol. in H_2O . (Grossmann, B. 1904, 37. 565.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O and alcohol. Insol. in acetone. (Krug and M'Elroy.)

+ $1\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O . (Rosenheim and Cohn, Z. anorg. 1901, 27. 292.)

Nickel potassium sulphocyanide,
 $\text{K}_2\text{Ni}(\text{SCN})_6 + 4\text{H}_2\text{O}$.

Sol. in H_2O with decomp.

Sl. sol. in cold, easily sol. in hot alcohol. (Rosenheim, Z. anorg. 1901, 27. 292.)

Nickel sodium sulphocyanide,
 $\text{NiNa}_2(\text{SCN})_6 + 8\text{H}_2\text{O}$.

Sol. in H_2O with decomp.

Sl. sol. cold, readily sol. hot alcohol. (Rosenheim, Z. anorg. 1901, 27. 292.)

Nickel sulphocyanide ammonia, $\text{Ni}(\text{SCN})_2, 3\text{NH}_3$.

(Peters, B. 1908, 41. 3178.)

$\text{Ni}(\text{SCN})_2, 4\text{NH}_3$. Decomp. by H_2O .

Platinous sulphocyanide, $\text{Pt}(\text{SCN})_2(?)$.

Insol. in H_2O .

See *Platinosulphocyanides, and Platinosulphocyanides*.

Potassium sulphocyanide, KSCN .

Deliquescent. Very sol. in H_2O . 100 pts. H_2O dissolve 177.2 pts. at 0° , and 217.0 pts. at 20° .

100 g. sat. $\text{KSCN} + \text{Aq}$ contain 70.5 g. KSCN at 25° . (Foote, Z. phys. Ch. 1903, 46. 81.)

150 pts. $\text{KSCN} + 100$ pts. H_2O at 10.8° lower the temp. 34.5° . (Rüdorff, B. 2. 68.)

Solubility of $\text{KSCN} + \text{AgSCN}$ at 25° .

% KSCN	% AgSCN	Solid phase
70.53	0.00	KSCN
66.55	9.32	$\text{KSCN} + 2\text{KSCN}, \text{AgSCN}$
64.47	10.62	$2\text{KSCN}, \text{AgSCN}$
61.25	11.76	"
58.34	13.55	"
53.21	17.53	"
50.68	20.43	$2\text{KSCN}, \text{AgSCN} + \text{KSCN}, \text{AgSCN}$
49.43	20.32	$\text{KSCN}, \text{AgSCN}$
32.51	18.34	"
24.68	16.41	"
23.86	16.07	$\text{KSCN}, \text{AgSCN} + \text{AgSCN}$

(Foote, Z. phys. Ch. 1903, 46. 81.)

See also AgSCN .

Sol. in alcohol, especially easily if boiling.

Sol. in acetone. (Krug and M'Elroy.)

Sol. in liquid SO_2 . (Walden, Z. anorg. 1902, 30. 160.)

100 g. acetone dissolve 20.75 g. KSCN at 22° , and 20.40 g. at 58° .

100 g. amyl alcohol dissolve 0.18 g. KSCN at 13° ; 1.34 g. at 65° ; 2.14 g. at 100° ; 3.15 g. at 133.5° .

100 g. ethyl acetate dissolve 0.44 g. KSCN at 0° ; 0.40 g. at 14° ; 0.20 g. at 79° .

100 g. pyridine dissolve 6.75 g. KSCN at 0° ; 6.15 g. at 20° ; 4.97 g. at 58° ; 3.88 g. at

97°; 3.21 g. at 115°. (Laszcynski, B. 1894, 27. 2285.)

100 g. acetonitrile dissolve 11.31 g. KSCN at 18°. (Naumann and Schier, B. 1914, 47. 249.)

Sl. sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3789.)

Potassium molybdenyl sulphocyanide,
3KSCN, Mo(OH)(SCN)₃+4H₂O.

Sol. in H₂O. (Sand and Maas, B. 1908, 41. 1506.)

Potassium silver sulphocyanide, KSCN, AgSCN.

Decomp. by H₂O.

See Donk under KSCN.

2KSCN, AgSCN. Stable in the air. (Wells, Am. Ch. J. 1902, 28. 265.)

See Donk under KSCN.

3KSCN, AgSCN. (Wells.)

Potassium stannic sulphocyanide,
K₂Sn(SCN)₆+4H₂O.

Very sol. in H₂O.

Sol. in alcohol and acetone. (Weinland and Barnes, Z. anorg. 1909, 62. 258.)

Potassium titanyl sulphocyanide,
2KSCN, TiO(SCN)₂+H₂O.

Sol. in cold H₂O without immediate decomp. but slowly decomp. (Rosenheim and Cohn, Z. anorg. 1901, 28. 169.)

Potassium vanadium sulphocyanide,
3KSCN, V(SCN)₃+4H₂O.

Sol. in H₂O. Sol. in alcohol with a green color. Sl. sol. in ether. (Ciocci, Z. anorg. 1898, 19. 309.)

Sol. in H₂O with decomp.; stable in aq. solution in the presence of an excess of KSCN; sol. in alcohol. (Locke, Am. Ch. J. 1898, 20. 604.)

Potassium vanadyl sulphocyanide,
K₂VO(SCN)₄+5H₂O.

Sol. in H₂O, alcohol, ether, amyl alcohol and ethyl acetate. (Koppel, Z. anorg. 1903, 36. 292.)

Potassium zinc sulphocyanide, 2KSCN, Zn(SCN)₂+3H₂O.

Easily sol. in alcohol. (Walden, Z. anorg. 1900, 23. 374.)

Potassium sulphocyanide mercuric bromide,
KSCN, HgBr₂.

Very sol. in H₂O.

Sol. in alcohol. (Grossmann, B. 1902, 35. 2945.)

2KSCN, HgBr₂. Very sol. in H₂O. Sol. in alcohol. (Grossman.)

Potassium sulphocyanide mercuric iodide,
2KSCN, HgI₂.

Undecomp. by solution in conc. alcohol, or in KSCN+Aq. (Grossmann, Z. anorg. 1903, 37. 421.)

+2H₂O. Decomp. by H₂O. (Philipp, Pogg, 1867, 131. 94.)

Silicon sulphocyanide, Si(SCN)₄.

Decomp. by H₂O and alcohol.

Sol. in CS₂, CHCl₃, and ligroin, (Reynolds, Proc. Chem. Soc. 1906, 22. 17.)

Silver sulphocyanide, AgSCN.

1 l. H₂O dissolves 1.08×10^{-4} g. mol AgSCN at 25°. (Küster and Thiel, Z. anorg. 1902, 33. 139.)

1 l. H₂O dissolves 1.25×10^{-4} gram-atoms of silver at 25°. (Abegg and Cox, Z. phys. Ch. 1903, 46. 11.)

Sl. sol. in H₂O. 1 liter of sat. solution at 19.96° contains 1.37×10^{-4} g. (Böttger, Z. phys. Ch. 1903, 46. 603.)

6.4 milligrams are dissolved in 1 liter of sat. solution at 100°. (Böttger, Z. phys. Ch. 1906, 56. 93.)

Solubility product of AgSCN is 0.49 and 1.16×10^{-12} mols. per l. at 18° and 25° respectively. (Kirschner, Z. phys. Ch. 1912, 79. 245.)

Solubility in H₂O = 1.2×10^{-4} g. mol per liter at 25°. (A. E. Hill, J. Am. Chem. Soc. 1908, 30. 74.)

1 l. H₂O dissolves 0.00025 g. AgSCN at 21°. (Whitby, Z. anorg. 1910, 67. 108.)

Insol. in acids, excepting conc. H₂SO₄, or HNO₃. Insol. in dil., sol. in conc. NH₄OH+Aq. Sol. in KSCN+Aq. Insol. in AgNO₃ or NH₄SCN+Aq. Sol. in Hg₂(NO₃)₂+Aq.

Solubility in KSCN at 25°.

Mol. KSCN in 1 litre	g. AgSCN in 1 litre
1.25	22.34
1.20	19.93
1.12	16.18
1.066	14.10
0.626	2.80
0.573	2.06

(Hellwig, Z. anorg. 1900, 25. 184.)

Solubility in N/10 KSCN+Aq at 18° = 2.5×10^{-4} . (Kirschner, Z. phys. Ch. 1912, 72. 247.)

See also KSCN.

1 l. of a 3-N solution of AgNO₃ dissolves 0.432 g. AgSCN at 25°. Nearly insol. in less dil. solution. (Hellwig, Z. anorg. 1900, 25. 179.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in ethyl acetate. (Hamers, Dissert. 1906; Naumann, B. 1910, 42. 314.)

Silver strontium sulphocyanide, 2AgSCN , $\text{Sr}(\text{SCN})_2 + 2\text{H}_2\text{O}$.

Stable in the air. (Wells, Am. Ch. J. 1902, 270.)

Silver zinc sulphocyanide, 2AgSCN , $\text{Zn}(\text{SCN})_2$.

Decomp. by hot H_2O . (Wells.)

Silver sulphocyanide ammonia, AgSCN , 2NH_3 .

Decomp. by H_2O .

Samarium sulphocyanide, $\text{Sm}(\text{SCN})_3 + 6\text{H}_2\text{O}$.

Very deliquescent. (Cleve.)

Sodium sulphocyanide, NaSCN .

Very deliquescent. Very sol. in H_2O and alcohol.

Sol. in benzonitrile. (Naumann, B. 1914, 47. 1369.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3789.)

Sodium stannic sulphocyanide, $\text{Na}_2\text{Sn}(\text{SCN})_6 + 6\text{H}_2\text{O}$.

Very sol. in H_2O . Sol. in alcohol and acetone. (Weinland and Bames, Z. anorg. 1909, 62. 257.)

Sodium vanadium sulphocyanide, 3NaSCN , $\text{V}(\text{SCN})_3 + 12\text{H}_2\text{O}$.

Very hygroscopic. Sol. in H_2O and alcohol. (Ciocci, Z. anorg. 1898, 19. 313.)

Strontium sulphocyanide, $\text{Sr}(\text{SCN})_2 + 3\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O and alcohol.

Strontium stannic sulphocyanide, $\text{SrSn}(\text{SCN})_6 + 12\text{H}_2\text{O}$.

Sol. in H_2O , alcohol and acetone. (Weinland and Bames, Z. anorg. 1909, 62. 259.)

Thallium sulphocyanide, TlSCN .

Sl. sol. in H_2O . 3.15 g. are contained in 1 liter of sat. solution at 20° ; 3.905 g. at 25° ; 7.269 g. at 39.75° . Insol. in alcohol. (Böttger, Z. phys. Ch. 1903, 46. 603.)

Titanyl sulphocyanide, $\text{TiO}(\text{SCN})_2 + 2\text{H}_2\text{O}$.

Sol. in cold H_2O .

Tin (stannous) sulphocyanide, $\text{Sn}(\text{SCN})_2$.

Sol. in H_2O and alcohol. (Classen, J. pr. 96. 349.)

Sol. in cold H_2O . (Rosenheim, Z. anorg. 1901, 28. 168.)

Yttrium sulphocyanide, $\text{Y}(\text{SCN})_3 + 6\text{H}_2\text{O}$.

Not deliquescent. Very sol. in H_2O , alcohol, or ether.

Zinc sulphocyanide, $\text{Zn}(\text{SCN})_2$.

Less sol. in H_2O and alcohol than most other cyanides.

Zinc sulphocyanide ammonia, $\text{Zn}(\text{SCN})_2$, 12NH_3 .

Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Sulphocyanoplatinic acid.

See Platinosulphocyanhydric acid.

Sulphocyanoplatinous acid.

See Platinosulphocyanhydric acid.

Sulphohypophosphoric acid.

Aluminum sulphohypophosphate, $\text{Al}_2(\text{PS}_2)_3$.

Unstable in the air. Sol. in H_2O with decomp. (Friedel, C. R. 1894, 119. 262.)

Cadmium —, $\text{Cd}_2\text{P}_2\text{S}_6$.

Partially decomp. in moist air. Decomp. by H_2O , cold HNO_3 or alkalis + Aq. (Ferrand, A. ch. 1899, (7) 17. 423; Bull. Soc. 1895, (3) 13. 116.)

Chromium —, $\text{Cr}_2\text{P}_2\text{S}_6$.

Insol. in HNO_3 . Very sl. attacked by aqua regia. (Ferrand.)

Cupric —, $\text{Cu}_2\text{P}_2\text{S}_6$.

Ppt. (Friedel, C. R. 1894, 119. 262.)

Iron (ferrous) —, $\text{Fe}_2\text{P}_2\text{S}_6$.

Sol. in HNO_3 and in a mixture of HNO_3 with KClO_3 . (Friedel.)

Lead —, $\text{Pb}_2\text{P}_2\text{S}_6$.

Not decomp. by boiling H_2O . (Friedel.)

Mercuric —, $\text{Hg}_2\text{P}_2\text{S}_6$.

Slowly decomp. by boiling H_2O , more rapidly by $\text{KOH} + \text{Aq}$. (Friedel.)

Nickel —, $\text{Ni}_2\text{P}_2\text{S}_6$.

Not attacked by boiling H_2O or hot or cold acids. Sl. attacked by aqua regia. (Ferrand, A. ch. 1899, (7) 17. 416.)

Silver —, $\text{Ag}_4\text{P}_2\text{S}_6$.

Ppt. (Friedel, C. R. 1894, 119. 263.)

Tin (stannous) hypophosphate —, SnPS_2 .

Decomp. by boiling H_2O . Sol. in dil. $\text{KOH} + \text{Aq}$. (Friedel, C. R. 1894, 119. 264.)

Tin (stannic) —, SnP_2S_6 .

Easily decomp. by boiling H_2O . Sol. in dil. $\text{KOH} + \text{Aq}$. (Friedel.)

Zinc sulphohypophosphate, $\text{Zn}_2\text{P}_2\text{S}_6$.

Decomp. in moist air. Insol. in H_2O . Partially decomp. by boiling H_2O . Violently attacked by HNO_3 . Sol. in aqua regia. Not attacked by HCl . (Ferrand, A. ch. 1899, (7) 17. 421.)

Zinc —, $\text{Zn}_2\text{P}_2\text{S}_6$.

Insol. in H_2O . Partly decomp. by boiling H_2O . Not decomp. by HCl or HNO_3 but by aqua regia. (Ferrand, Bull. Soc. 1895, (3) 13. 115.)

Sulphomolybdic acid.**Ammonium sulphomolybdate, $(\text{NH}_4)_2\text{MoS}_4$.**

Easily sol. in H_2O ; very sl. sol. in alcohol. (Berzelius, Pogg. 83. 261.)

Ammonium cupric sulphomolybdate.

Sl. sol. in H_2O . (Debray, C. R. 96. 1616.)

Barium sulphomolybdate, BaMoS_4 .

More sol. in H_2O than $\text{BaMo}_2\text{S}_{10}$. Known only in solution. (Berzelius.)

$\text{BaS}, 3\text{MoS}_3 = \text{BaMo}_2\text{S}_{10}$. Sl. sol. in cold, easily sol. in hot H_2O . Not decomp. by conc. cold $\text{HNO}_3 + \text{Aq}$, but more easily by dil. $\text{HNO}_3 + \text{Aq}$. (Berzelius.)

Cadmium sulphomolybdate.

Insol. in H_2O . (Berzelius.)

Cæsium sulphomolybdate, $\text{Cs}_2\text{S}, 3\text{MoS}_4 + 7\text{H}_2\text{O}$.

As Rb comp. (Herschfinkel, Dissert. 1907.) $3\text{Cs}_2\text{S}, 5\text{MoS}_4$. (Herschfinkel.)

Calcium sulphomolybdate, $\text{CaS}, 3\text{MoS}_3$.

Sol. in H_2O . (Berzelius.)

CaMoS_4 . More sol. in H_2O than $\text{CaS}, 3\text{MoS}_3$. Known only in solution. (Berzelius.)

Cerium sulphomolybdate.

Precipitate. (Berzelius.)

Cobalt sulphomolybdate, CoMoS_4 .

Sol. in $\text{K}_2\text{MoS}_4 + \text{Aq}$. (Berzelius.)

Cupric sulphomolybdate.

(Debray, C. R. 96. 1616.)

Ferrous sulphomolybdate, FeMoS_4 .

Sol. in H_2O . (Berzelius.)

Ferric sulphomolybdate, $\text{Fe}_2(\text{MoS}_4)_3$.

Sol. in $\text{K}_2\text{MoS}_4 + \text{Aq}$.

Lead sulphomolybdate.

Ppt. (Berzelius.)

Lithium sulphomolybdate.

Not deliquescent, but very easily sol. in H_2O . (Berzelius.)

Magnesium sulphomolybdate, MgMoS_4 .

Sol. in $\text{K}_2\text{MoS}_4 + \text{Aq}$. (Berzelius.)

Manganous sulphomolybdate, MnMoS_4 .

Sol. in H_2O . (Berzelius.)

Mercurous sulphomolybdate, Hg_2MoS_4 (?)

Ppt.

Mercuric sulphomolybdate, HgMoS_4 .

Insol. in $\text{K}_2\text{MoS}_4 + \text{Aq}$.

Nickel sulphomolybdate, NiMoS_4 .

Sol. in $\text{K}_2\text{MoO}_4 + \text{Aq}$. (Berzelius.)

Potassium sulphomolybdate, basic, K_2MoS_4 .

Easily sol. in H_2O . Insol. in alcohol and ether. (Kruss, B. 16. 2050.)

Potassium sulphomolybdate, K_2MoS_4 .

Sol. in H_2O , from which it is precipitated by alcohol. (Berzelius.)

Rubidium sulphomolybdate, $3\text{Rb}_2\text{S}, 8\text{MoS}_3 + 30\text{H}_2\text{O}$.

Very sl. sol. in H_2O . Sol. by addition of NH_3 . (Herschfinkel, Dissert. 1907.) $5\text{Rb}_2\text{S}, 6\text{MoS}_3$. (Herschfinkel.)

Silver sulphomolybdate, Ag_2MoS_4 .

Ppt.

Sodium sulphomolybdate, Na_2MoS_4 .

Sol. in H_2O , and not precipitated by alcohol from aqueous solution. (Berzelius.)

Strontium sulphomolybdates.

Exactly analogous to the Ba salts, which see. (Berzelius.)

Zinc sulphomolybdate.

Ppt. Insol. in H_2O . (Berzelius.)

Monosulphomolybdic acid.**Sodium monosulphomolybdate, $\text{Na}_2\text{MoO}_3\text{S}$.**

Rather hygroscopic. Sol. in H_2O ; forms deep blue solution with H_2SO_4 . Sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Kruss, A. 225. 1.)

Disulphomolybdic acid.**Ammonium disulphomolybdate, $(\text{NH}_4)_2\text{MoO}_3\text{S}_2$.**

Sl. sol. in cold, easily in hot H_2O . Insol. in sat. $\text{NH}_4\text{Cl} + \text{Aq}$ and absolute alcohol. Aqueous solution is decomp. by boiling. (Bodenstab, J. pr. 76. 183.)

am disulphomolybdate, $K_2MoO_7S_2$.

sol. in H_2O and alcohol. Sol. in O_2 + Aq. (Krüss, B. 16. 2046.)

homolybdic acid.

ium hydrogen trisulphopyromolyb-
e, $NH_4HMo_2O_7S_3$.

pitate. Insol. in alcohol or CS_2 .
B. 16. 2047.)

am hydrogen trisulphopyromolybdate,
 $HM_2O_7S_3$.

easily sol. in H_2O . (Krüss, B. 16.

hydrogen trisulphopyromolybdate,
 $HMo_2O_7S_3$.

pitate. Much more sol. in H_2O than
a compound. (Krüss, B. 16. 2047.)

um sulphomolybdate, $K_2Mo_4S_7O_7$.

a H_2O , $HC_2H_3O_2$, and H_2SO_4 . (Krüss,
771.)

alphomolybdic acid.

um pentasulphomolybdate, $KMoS_5$.

n warm H_2O . (Hofmann, Z. anorg.
l. 62.)

homolybdic acid, H_2MoS_5 .

pitate. Insol. in H_2O , alcohol, ether,
d acetic acid.

mp. slowly by hot H_2SO_4 . Sol. in
 ζOH + Aq, and cold K_2S + Aq. Not
d by cold KSH + Aq, but dissolves
ning. (Krüss, B. 17. 1773.)

ium persulphomolybdate,
 $(H_4)_2MoS_6$.

sl. sol. in cold, more easily in hot
Insol. in NH_4OH + Aq. (Berzelius.)

—, $BaMoS_5$.

in boiling H_2O or dil. HCl + Aq.
us.)

l —.

ultly sol. in H_2O . (Berzelius.)

—.

pitate. (Berzelius.)

—.

in Fe salts + Aq, but sol. in K_2MoS_5 +
erzelius.)

—.

Lithium persulphomolybdate.

Sl. sol. in cold, easily sol. in hot H_2O .
(Berzelius.)

Magnesium —.

Insol. precipitate. (Berzelius.)

Nickel —.

Ppt. Sol. in K_2MoS_5 + Aq, from which it
separates in 24 hours. (Berzelius.)

Potassium —, K_2MoS_5 .

Almost insol. in cold, more sol. in hot H_2O .
Insol. in cold KOH + Aq. (Berzelius.)

Potassium hydrogen —, $KHMoS_5$.

Sol. in H_2O . (Krüss.)

Sodium —, Na_2MoS_5 .

Sl. sol. in cold, easily in hot H_2O . (Ber-
zelius.)

Sodium hydrogen —, $NaHMoS_5$.

(Krüss.)

Persulphomolybdic acid, $HMoS_5$.

Sol. in H_2O . (Hofmann, Z. anorg. 1896,
12. 59.)

Ammonium —, $NH_4MoS_5 + H_2O$.

Sl. sol. in H_2O and in alcohol with decomp.
(Hofmann.)

Cæsium —, $CaMoS_5$.

Almost insol. in H_2O . (Hofmann.)

Potassium —, $KMoS_5$.

Sol. in H_2O . (Hofmann.)

Thallium —, $TlMoS_5$.

Insol. in H_2O . (Hofmann.)

Sulphonosmic acid.

Potassium sulphonosmate,

$7K_2O, 4OsO_3, 10SO_2$.

Sol. in H_2O . (Rosenheim, Z. anorg. 1899,
21. 127.)

+ $3H_2O$. Sol. in H_2O . (Rosenheim.)

+ $7H_2O$. Easily sol. in H_2O ; decomp. in
aq. solution at 70° . (Rosenheim.)

$11K_2O, 4OsO_3, 14SO_2 + 7H_2O$. Sol. in H_2O .
(Rosenheim.)

Sodium sulphonosmate,

$3Na_2O, OsO_3, 4SO_2 + 5H_2O$.

Easily sol. in H_2O ; decomp. in aq. solution.
(Rosenheim.)

Sulphopalladic acid.

Potassium palladious sulphopalladate, K_2S , Pd_2S , $PdS_2 = K_2Pd_2S_4$.

Insol. in H_2O . Moderately conc. $HCl + Aq$ dissolves out K without evolution of H_2S . (Schneider, Pogg. 141. 526.)

Silver sulphopalladate, Ag_2PdS_2 .
(Schneider.)

Silver palladious sulphopalladate, Ag_2S , Pd_2S , $PdS_2 = Ag_2Pd_2S_4$.
Extraordinarily stable. (Schneider.)

Sodium sulphopalladate, Na_2PdS_2 .
Slowly sol. in H_2O . Insol. in alcohol. (Schneider, Pogg. 141. 520.)

Sulphophosphide of M.
See *M* phosphosulphide.

Sulphophosphamic acid, $PS^{(OH)}_{NH_2}(?)$.
See Thiophosphamic acid.

Sulphophosphodiamic acid, $PS^{OH}_{(NH_2)_2}(?)$.
See Thiophosphodiamic acid.

Sulphophosphotriamide, $PS(NH_2)_3$.
See Thiophosphoryl triamide.

Sulphophosphoric acid, H_2PSO_3 .
See Thiophosphoric acid.
 H_2PS_4 . Known only in its salts.

Ammonium sulphophosphate, $(NH_4)_2PS_4$.
Stable in the air. (Ephraim, B. 1911, 44. 3408.)

Antimony sulphophosphate, $SbPS_4$.
Insol. in H_2O , alcohol, ether, CS_2 , $HCl + Aq$, dil. $H_2SO_4 + Aq$, C_2H_6 , or $HC_2H_3O_2$. Decomp. by boiling with conc. $HNO_3 + Aq$, H_2SO_4 , aqua regia, KOH , $NaOH$ or $NH_4OH + Aq$. (Glatzel, B. 24. 3886.)

Arsenic sulphophosphate, $AsPS_4$.
Insol. in H_2O , alcohol, $HCl + Aq$, etc. Decomp. by warm HNO_3 , aqua regia, dil. H_2SO_4 ; also sol. in KOH or $NH_4OH + Aq$. (Glatzel, Z. anorg. 4. 186.)

Barium sulphophosphate, $Ba_3(PS_4)_2 + xH_2O$.
(Ephraim, B. 1911, 44. 3409.)

Bismuth sulphophosphate, $BiPS_4$.
Insol. in H_2O , alcohol, ether, CS_2 , benzene, $HC_2H_3O_2$, or dil. $H_2SO_4 + Aq$. Decomp. by boiling $HCl + Aq$, conc. H_2SO_4 , HNO_3 , or aqua regia; also by $NaOH$, KOH , or $NH_4OH + Aq$. (Glatzel, Z. anorg. 4. 186.)

Cadmium sulphophosphate, $Cd_2(PS_4)_2$.
Insol. in H_2O , alcohol, ether, benzene, CS_2 , and $HC_2H_3O_2$. Decomp. by hot $HCl + Aq$. Very sl. attacked by dil. $H_2SO_4 + Aq$. Slowly sol. in hot HNO_3 , rapidly in aqua regia or hot conc. H_2SO_4 . (Glatzel, Z. anorg. 4. 186.)

Cuprous sulphophosphate, Cu_2PS_4 .
Insol. in H_2O , alcohol, etc.; also in HCl or dil. $H_2SO_4 + Aq$. Decomp. by HNO_3 , aqua regia, etc., not by KOH or $NaOH + Aq$. (Glatzel.)

Ferrous sulphophosphate, $Fe_2(PS_4)_2$.
Insol. in H_2O , alcohol, ether, etc.; insol. in HCl or hot dil. $H_2SO_4 + Aq$. Decomp. by HNO_3 , aqua regia, or conc. H_2SO_4 . Not attacked by KOH or $NH_4OH + Aq$. (Glatzel.)

Lead sulphophosphate, $Pb_2(PS_4)_2$.
Insol. in H_2O , alcohol, etc. Decomp. by warm $HCl + Aq$, conc. $HNO_3 + Aq$; not attacked by $NH_4OH + Aq$; sl. decomp. by $KOH + Aq$. (Glatzel.)

Manganous sulphophosphate, $Mn_2(PS_4)_2$.
Insol. in H_2O , alcohol, ether, benzene, CS_2 , or $HC_2H_3O_2$. Not attacked by $HCl + Aq$. Sol. in HNO_3 or aqua regia, with separation of S . Not attacked by dil. $H_2SO_4 + Aq$. (Glatzel, Z. anorg. 4. 186.)

Mercuric sulphophosphate, $Hg_2(PS_4)_2$.
Insol. in H_2O , alcohol, etc.; also in HCl , dil. HNO_3 , or $H_2SO_4 + Aq$. Not attacked by conc. HNO_3 or aqua regia; easily sol. in $HNO_3 + Br_2 + Aq$. (Glatzel.)

Nickel sulphophosphate, $Ni_2(PS_4)_2$.
As the ferrous salt. (Glatzel.)

Potassium sulphophosphate, $K_2PS_4 + H_2O$.
Easily sol. in H_2O . (Ephraim, B. 1911, 44. 3407.)

Silver sulphophosphate, Ag_2PS_4 .
Insol. in H_2O , alcohol, etc.; also in HCl , HNO_3 , or dil. $H_2SO_4 + Aq$. Decomp. by conc. H_2SO_4 , and aqua regia. (Glatzel.)

Sodium sulphophosphate, $Na_2PS_4 + 8H_2O$.
Decomp. by H_2O .
Sol. in $Na_2S + Aq$. (Glatzel, Z. anorg. 1905, 44. 65.)

Thallous sulphophosphate, Tl_2PS_4 .
Insol. in H_2O , alcohol, etc. Sol. in HCl , dil. $H_2SO_4 + Aq$, etc. Not attacked by $NH_4OH + Aq$; sl. decomp. by conc. $KOH + Aq$. (Glatzel.)

Stannous (stannous) sulphophosphate, $\text{Sn}_2(\text{PS}_2)_2$.
 Insol. in H_2O , alcohol, etc. Insol. in dil. H_2SO_4 or $\text{HCl} + \text{Aq}$. Decomp. by $\text{HNO}_3 + \text{Aq}$, aqua regia, NH_4OH , or $\text{KOH} + \text{Aq}$. (Glatzel.)

Zinc sulphophosphate, $\text{Zn}_2(\text{PS}_2)_2$.

Insol. in H_2O , alcohol, ether, etc. Sol. in $\text{HCl} + \text{Aq}$ or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Easily attacked by $\text{KOH} + \text{Aq}$; sl. decomp. by $\text{NH}_4\text{OH} + \text{Aq}$. (Glatzel.)

Sulphopyrophosphoric acid.

Aluminum sulphopyrophosphate, $\text{Al}_2\text{P}_2\text{S}_7$.

Decomp. in moist air.
 Violently decomp. by H_2O or acids. (Ferrand, A. ch. 1899, (7) 429.)

Cadmium —, $\text{Cd}_2\text{P}_2\text{S}_7$.

Decomp. in moist air.
 Not attacked by cold acids. (Ferrand.)

Chromium —, $\text{Cr}_2\text{P}_2\text{S}_7$.

Decomp. in moist air.
 Not readily attacked by acids. (Ferrand.)

Cuprous —, $\text{Cu}_2\text{P}_2\text{S}_7$.

Not attacked by cold H_2SO_4 or boiling HCl . (Ferrand.)

Sol. in hot conc. HNO_3 . (Ferrand.)
 Sol. in alkalies, and in all acids except HCl . (Ferrand, C. R. 1896, 122. 886.)

Ferrous —, $\text{Fe}_2\text{P}_2\text{S}_7$.

Insol. in cold acids.
 Sl. attacked by boiling HCl or hot $\text{KOH} + \text{Aq}$.
 Decomp. by fused KOH . (Ferrand, A. ch. 1899, (7) 17. 410.)

Lead —, $\text{Pb}_2\text{P}_2\text{S}_7$.

Not attacked by cold HNO_3 . (Ferrand.)

Mercurous —, $\text{Hg}_2\text{P}_2\text{S}_7$.

Decomp. by moist air or hot HNO_3 . (Ferrand.)
 Almost insol. in acids; decomp. by H_2O and moist air. (Ferrand, C. R. 1896, 122. 888.)

Nickel —, $\text{Ni}_2\text{P}_2\text{S}_7$.

Decomp. by H_2O and by conc. HNO_3 at 150° in a sealed tube. (Ferrand, A. ch. 1899, (7) 17. 418.)

Silver —, $\text{Ag}_2\text{P}_2\text{S}_7$.

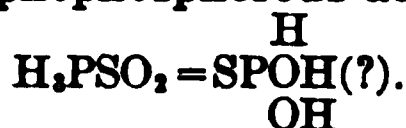
Not decomp. by H_2O .
 Decomp. by aqua regia.
 Not attacked by HNO_3 . (Ferrand.)

Zinc —, $\text{Zn}_2\text{P}_2\text{S}_7$.

Decomp. in moist air.
 Decomp. by H_2O .

Violently attacked by cold HNO_3 . (Ferrand.)

Sulphophosphorous acid,



See Thiophosphorous acid.

H_2PS_2 . Known only in its salts.

Aluminum sulphophosphite, $\text{Al}_2(\text{PS}_2)_2$.

Very unstable.
 Decomp. in the air. (Ferrand, C. R. 1896, 122. 622.)

Barium sulphophosphite, $\text{Ba}_2(\text{PS}_2)_2 + x\text{H}_2\text{O}$.

Sol. in dil. acids.
 Insol. in alcohol. (Ephraim, B. 1911, 44. 3412.)

Chromous sulphophosphite, $\text{Cr}_2(\text{PS}_2)_2$.

Easily attacked by hot conc. HNO_3 or aqua regia.

Decomp. by boiling $\text{NaOH} + \text{Aq}$. (Ferrand, A. ch. 1899, (7) 17. 419.)

Quite stable in moist air; very slowly attacked by acids. (Ferrand, C. R. 1896, 122. 622.)

Cuprous sulphophosphite, Cu_2PS_2 .

Not attacked by H_2O or hot conc. HCl .
 Sl. attacked by cold fuming HNO_3 .
 Violently attacked by HNO_3 , aqua regia and boiling conc. H_2SO_4 .
 Not attacked by boiling $\text{NaOH} + \text{Aq}$. (Ferrand, A. ch. 1899, (7) 17. 398.)
 Fairly stable decomp. by damp air. (Ferrand, C. R. 1896, 122. 621.)

Iron (ferrous) sulphophosphite, $\text{Fe}_2(\text{PS}_2)_2$.

Very stable and resists the action of alkalies and acids. (Ferrand, C. R. 1896, 122. 622.)
 Insol. in cold acids or hot NCl .
 Sol. in hot fuming HNO_3 .
 Insol. in hot 40% $\text{KOH} + \text{Aq}$. (Ferrand, A. ch. 1899, (7) 17. 412.)

Mercuric sulphophosphite, $\text{Hg}_2(\text{PS}_2)_2$.

Decomp. in moist air.
 Not attacked by cold HNO_3 . Decomp. by hot HNO_3 . (Ferrand.)
 Unstable in the air.
 Very slowly attacked by acids. (Ferrand, C. R. 1896, 122. 622.)

Nickel sulphophosphite, $\text{Ni}_2(\text{PS}_2)_2$.

Unstable in the air.
 Attacked slowly by HNO_3 . (Ferrand.)

Silver sulphophosphite, Ag_2PS_2 .

Insol. in most reagents. (Ferrand, C. R. 1896, 122. 622.)
 Not decomp. by H_2O .
 Not easily attacked by acids. (Ferrand, A. ch. 1899, (7) 17. 414.)

Sodium sulphophosphite, $\text{Na}_2\text{PS}_2 + x\text{H}_2\text{O}$.

Very sol. in H_2O , probably with decomp. (Ephraim, B. 1911, 44. 3410.)

Zinc sulphophosphite, $\text{Zn}_2(\text{PS}_2)_2$.

Decomp. in moist air.

Sl. attacked by H_2O .

Decomp. by HNO_3 . (Ferrand, A. ch. 1899, (7) 17. 422.)

Very unstable in the air, and attacked violently by acids. (Ferrand, C. R. 1896, 122. 622.)

Sulphoplatinic acid, $\text{H}_2\text{Pt}_4\text{S}_6$.

Insol. in H_2O , but decomp. on air. (Schneider, Pogg. 138. 604.)

$\text{H}_4\text{Pt}_4\text{S}_6$. Insol. in H_2O , but decomp. very rapidly on air. (Schneider.)

Copper sulphoplatinate, 2CuS , 2PtS , PtS_2 .

Insol. in H_2O . HCl , HNO_3 , or aqua regia dissolve out part of the Cu. (Schneider, Pogg. 139. 661.)

Lead sulphoplatinate, 2PbS , 2PtS , PtS_2 .

Insol. in hot or cold H_2O or $\text{HCl} + \text{Aq}$. $\text{HNO}_3 + \text{Aq}$ dissolves out Pb partly; aqua regia dissolves completely with difficulty. (Schneider, Pogg. 139. 662.)

Mercuric sulphoplatinate chloride, 2HgS , 2PtS , PtS_2 , 2HgCl_2 .

Insol. in H_2O ; not attacked by $\text{HCl} + \text{Aq}$, and only partially sol. in boiling aqua regia. (Schneider.)

Potassium sulphoplatinate, $\text{K}_2\text{Pt}_4\text{S}_6$.

Insol. in H_2O . $\text{HCl} + \text{Aq}$ dissolves out K without evolution of H_2S .

Composition its potassium platinous sulphoplatinate, K_2S , 3PtS , PtS_2 . (Schneider, Pogg. 138. 604.)

K_2PtS_2 .

Silver sulphoplatinate, $2\text{Ag}_2\text{S}$, 2PtS , PtS_2 .

Insol. in H_2O or $\text{HCl} + \text{Aq}$. $\text{HNO}_3 + \text{Aq}$ dissolves out Ag on warming. Aqua regia decomp. with formation of AgCl . (Schneider, Pogg. 138. 664.)

Sodium sulphoplatinate, $\text{Na}_4\text{Pt}_2\text{S}_6 = 2\text{Na}_2\text{S}$, 2PtS , PtS_2 .

Decomp. by hot H_2O , with residue of PtS_2 . (Schneider.)

$\text{Na}_2\text{Pt}_2\text{S}_6 = \text{Na}_2\text{S}$, PtS , 2PtS_2 . Insol. in H_2O . (Schneider, J. pr. (2) 48. 418.)

Thallium sulphoplatinate, $2\text{Tl}_2\text{S}$, 2PtS , PtS_2 .

Insol. in cold H_2O . Dil. acids dissolve out all the thallium. (Schneider, Pogg. 138. 626.)

Sulphoplatinous acid, H_2PtS_2 .

Known only in solution in H_2O , which soon decomposes. (Schneider, J. pr. (2) 48. 424.)

Sodium sulphoplatinite, Na_2PtS_2 .

Sol. in H_2O with decomp. (Schneider, J. pr. (2) 48. 420.)

$\text{H}_4\text{Na}_2(\text{PtS}_2)_2$. Sol. in H_2O , from which it is pptd. by alcohol. (Schneider.)

Sulphoselenantimonous acid.

See Selenosulphantimonous acid.

Sulphoselenarsenic acid.

See Selenosulpharsenic acid.

Sulphoselenostannic acid.

See Selenosulphostannic acid.

Sulphoselenoxyarsenic acid.

See Selenosulphoxyarsenic acid.

Sulphoselenyl chloride, SSeO_2Cl_2 .

Deliquescent; decomposed by H_2O . (Clemmenger, B. 11. 2007.)

Metasulphosilicic acid.

Sodium metasulphosilicate, Na_2SiS_2 .

Decomp. by H_2O . (Hempel, Z. anorg. 1900, 23. 41.)

Sulphostannic acid, H_2SnS_3 .

Ppt. (Kühn, A. 84. 110.)

Does not exist. (Storch, W. A. B. 2b. 236.)

Ammonium sulphostannate, $(\text{NH}_4)_2\text{S}$, $3\text{SnS}_2 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O , and easily decomp. (Ditte, C. R. 95. 641.)

$(\text{NH}_4)_2\text{SnS}_3 + 3\text{H}_2\text{O}$, and $+7\text{H}_2\text{O}$. Decomp. by acid. (Stanek, Z. anorg. 1896, 11. 124.)

Barium sulphostannate, $\text{BaSnS}_3 + 8\text{H}_2\text{O}$.

Sol. in cold H_2O . (Ditte, C. R. 95. 641.)

Calcium sulphostannate, 2CaS , $\text{SnS}_2 + 14\text{H}_2\text{O}$.

Sol. in H_2O . (Ditte, C. R. 95. 641.)

Tetraplatinous sulphostannate, 4PtS , SnS_2 .

Not decomp. by acids. (Schneider, J. pr. (2) 7. 214.)

Platinum potassium sulphostannate, 3PtS , K_2S , SnS_2 .

Insol. in cold H_2O . Dil. HCl or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ dissolves out all the potassium. (Schneider, Pogg. 138. 109.)

Platinum sodium sulphostannate, $3\text{PtS}_2, \text{SnS}_2$.

in cold H_2O . (Schneider, Pogg. 136.)

Potassium sulphostannate, K_2SnS_3 .

insol. in H_2O . (Kühn, A. 84. 110.)

in H_2O . (Ditte, C. R. 95. 641.)

$\text{SnS}_4 + 4\text{H}_2\text{O}$. Sol. in H_2O : pptd. by H_2S . (Weinland, Z. anorg. 1898, 17. 419.)

Sodium sulphostannate, $\text{Na}_2\text{SnS}_3 + 2\text{H}_2\text{O}$.

insol. in H_2O . (Kühn, A. 84. 110.)

in H_2O . (Ditte, C. R. 95. 641.)

in H_2O . Sol. in H_2O . (Höring, Zeitsch. 1851. 120.)

$\text{SnS}_4 + 12\text{H}_2\text{O}$. Melts in crystal H_2O on heating. Very sol. in H_2O . (Kühn.)

Strontium sulphostannate, $\text{SrSnS}_3 + 12\text{H}_2\text{O}$.

insol. in H_2O . (Ditte, C. R. 95. 641.)

Thallium sulphostannate, Tl_2SnS_4 .

Practically insol. in H_2O . (Hawley, Chem. Soc. 1907, 29. 1011.)

Tetrapersulphuric acid.

Sodium disulphopersulphate, $\text{Na}_2\text{S}_4\text{O}_8$.

insol. in H_2O . Cryst. in cold with $2\text{H}_2\text{O}$. (Berzelius, C. R. 106. 851, 1354.)

contains 4H more and is sodium tetraperphosphate, $\text{NaS}_4\text{O}_8, 2\text{H}_2\text{O}$. (Villiers, C. R. 1872.)

Telluric acid.

Mercurous sulphotellurate, $3\text{Hg}_2\text{S}, \text{TeS}_2$.

Mercurous —, $3\text{HgS}, \text{TeS}_2$.

(Berzelius.)

Potassium —, K_2TeS_4 .

insol. in H_2O . (Oppenheim, J. pr. 71. 279.)

Sodium —.

insol. in H_2O . (Oppenheim.)

Tellurous acid.

Ammonium sulphotellurite, $3(\text{NH}_4)_2\text{S}, \text{TeS}_2$.

decomp. on air. Sol. in H_2O .

Sodium —.

very slowly sol. in H_2O .

Mercurous —.

slightly sol. in H_2O .

Sodium —.

insol. ppt.

Cobalt sulphotellurite, Co_2TeS_4 .

Ppt.

Copper —, Cu_2TeS_4 .

Ppt.

Ferrous —.

Ppt.

Ferric —.

Ppt.

Lead —.

Ppt.

Lithium —.

Sol. in H_2O .

Magnesium —.

Sol. in H_2O and alcohol.

Manganous —.

Ppt.

Potassium —, $3\text{K}_2\text{S}, \text{TeS}_2$.

Sol. in H_2O .

Silver —, $3\text{Ag}_2\text{S}, \text{TeS}_2$.

(Berzelius.)

Sodium —.

Sol. in H_2O .

Strontium —.

Sol. in H_2O .

Zinc —, $3\text{ZnS}, \text{TeS}_2$.

Ppt. (Berzelius.)

Sulphotungstic acid.

Ammonium sulphotungstate, $(\text{NH}_4)_2\text{WS}_4$.

Very deliquescent. Easily sol. in H_2O , and still more easily in $\text{NH}_4\text{OH} + \text{Aq}$. (Corleis, A. 232. 244.)

More sol. in pure H_2O than in H_2O acidified with HCl . Decomp. slowly on air. (Berzelius.)

Barium —.

Sol. in $\text{BaS} + \text{Aq}$.

Cadmium —, CdWS_4 .

Ppt. (Berzelius.)

Calcium —.

Sol. in H_2O and alcohol. (Berzelius.)

Cobalt —, CoWS_4 .

Sl. sol. in H_2O .

Copper sulphotungstate, CuWS_4 .

Ppt.

Glucinum —, GlWS_4 .Sol. in H_2O (?).**Ferrous —, FeWS_4 .**Sol. in H_2O .**Ferric —.**

Ppt.

Lead —, PbWS_4 .

Ppt. (Berzelius.)

Magnesium —, MgWS_4 .Easily sol. in H_2O or alcohol.**Manganous —, MnWS_4 .**Sol. in H_2O . (Berzelius.)**Mercurous —.**

Ppt. (Berzelius.)

Mercuric —, HgWS_4 .

Ppt. (Berzelius.)

Nickel —, NiWS_4 .

Ppt. (Berzelius.)

Potassium —, K_2WS_4 .Sol. in H_2O . Alcohol precipitates from aqueous solutions, but is not entirely insol. in alcohol. (Berzelius.)Very sol. in H_2O . (Corleis, A. 232. 264.)**Potassium — nitrate, K_2WS_4 , KNO_3 .**Very sol. in cold or hot H_2O , from which it is precipitated by alcohol. (Berzelius.)**Potassium — tungstate, $\text{K}_2\text{WO}_3\text{S}_2 = \text{K}_2\text{WS}_4$, K_2WO_4 .**Easily sol. in H_2O . Not precipitated by alcohol. (Berzelius.)Is potassium trisulphotungstate, K_2WOS_3 , which see. (Corleis, A. 232. 244.)**Silver —, Ag_2WS_4 .**

Ppt. (Berzelius.)

Sodium —, Na_2WS_4 .Very sol. in H_2O ; less sol. in alcohol. (Berzelius.)

Very deliquescent. (Corleis, A. 232. 264.)

Strontium —.Sol. in H_2O , and in $\text{SrS} + \text{Aq}$.**Stannous —, SnWS_4 .**

Ppt. (Berzelius.)

Stannic —, SnWS_4 .

Ppt. (Berzelius.)

Zinc sulphotungstate, ZnWS_4 .Sol. in H_2O with subsequent pptn. (Berzelius.)**Monosulphotungstic acid.****Potassium monosulphotungstate,** $\text{K}_2\text{WOS} + \text{H}_2\text{O}$.Deliquescent in moist air. Very sol. in H_2O . (Corleis, A. 232. 244.)**Disulphotungstic acid.****Ammonium disulphotungstate, $(\text{NH}_4)_2\text{WOS}_2$.**Sol. in H_2O and alcohol. (Berzelius.)

Decomp. easily when moist. (Corleis, A. 232. 264.)

Trisulphotungstic acid.**Potassium trisulphotungstate, $\text{K}_2\text{WOS}_3 + \text{H}_2\text{O}$.**Hygroscopic. Effloresces on dry air and easily decomposed. Easily sol. in H_2O . (Corleis, A. 232. 244.)**Sulphovanadic acid, V_2O_5 , $3\text{SO}_3 + 3\text{H}_2\text{O}$.**

See Vanadiousulphuric acid, and Sulphate, vanadium.

Sulphovanadates.Alkali sulphovanadates are sol. in H_2O . Ca, Sr, and Ba sulphovanadates are sl. sol. in H_2O , and all other sulphovanadates are insol. in H_2O . (Berzelius.)**Ammonium sulphovanadate, $(\text{NH}_4)_2\text{VS}_4$.**Easily sol. in H_2O . Very sl. sol. in conc. $\text{NH}_4\text{SH} + \text{Aq}$. Insol. in ether, CS_2 , or CHCl_3 . (Krüss and Ohnmais, A. 263. 46.)

See also Sulphoxyvanadic acid.

Sodium pentasulphopyrovanadate, $\text{Na}_5\text{V}_2\text{O}_{15}$.Hydroscopic; sol. in H_2O with rapid decomp. (Locke, Am. Ch. J. 1898, 20. 375.)**Sulphoxyantimonic acid.****Potassium sulphonyantimonate, $\text{K}_2\text{HSbO}_5 + 2\text{H}_2\text{O}$.**Sol. in hot, less sol. in cold H_2O . Decomp. by cold H_2O . (Weinland and Gutmann, Z. anorg. 1898, 17. 414.)**Sulphoxyarsenic acid, H_3AsO_5 .**

Known only in aqueous solution. (McCay, Am. Ch. J. 10. 459.)

Ammonium monosulphonyarsenate, $(\text{NH}_4)_2\text{AsSO}_5 + 3\text{H}_2\text{O}$.Decomp. in the air; sol. in H_2O , decomp. on boiling. (Weinland, B. 1896, 29. 1009.)

Very sol. in H_2O ; insol. in alcohol; decomp. in aq. solution and also in the air. (Weinland, Z. anorg. 1897, 14. 53.)
Decomp. in the air. (McLaughlan, B. 1901, 34. 2166.)

Ammonium hydrogen monosulphoxyarsenate, $(NH_4)_2HAsSO_4$.
Ppt. (McLaughlan, B. 1901, 34. 2168.)

Barium monosulphoxyarsenate, $BaHAsO_4 + 10H_2O$.
 (Preis, A. 257. 184.)
 $Ba_3(AsSO_4)_2 + 6H_2O$. **Ppt.** (Weinland, Z. anorg. 1897, 14. 54.)

Barium disulphoxyarsenate, $Ba_3(AsS_2O_7)_2 + 4H_2O$.
Ppt. (Preis, A. 257. 185.)
 $+ 6H_2O$. (Weinland and Rumpf, Z. anorg. 1897, 14. 64.)

Barium potassium trisulphoxyarsenate, $KBaAsS_3O + 7H_2O$.
Ppt. (McCay, Z. anorg. 1904, 41. 469.)

Barium sodium monosulphoxyarsenate, $BaNaAsSO_4 + 9H_2O$.
Ppt. (Weinland, Z. anorg. 1897, 14. 55.)

Barium sodium sulphoxyarsenate, $Ba_7Na_2As_8O_7S_{14} + 12H_2O$.
 (McCay and Foster, Z. anorg. 1904, 41. 467.)

Calcium trisulphoxyarsenate, $Ca_3(AsS_2O_7)_2 + 20H_2O$.
Ppt. (McCay and Foster, Z. anorg. 1904, 41. 463.)

Potassium monosulphoxyarsenate, K_2AsSO_4 .
Hygroscopic. (Weinland, B. 1896, 29. 109.)
Sol. in conc. $KOH + Aq$, free from carbonate; very hygroscopic. (Weinland, Z. anorg. 1897, 14. 51.)

Potassium hydrogen monosulphoxyarsenate, $K_2HAsSO_4 + 2\frac{1}{2}H_2O$.
Very hygroscopic. (Weinland and Rumpf, Z. anorg. 1897, 14. 59.)
 KH_2AsSO_4 . **Sol. in H_2O ; solution slowly decomp. on standing.** (McCay, Am. Ch. J. 10. 459.)
Formula given by Bouquet and Cloez (A. ch. (3) 13. 44) is $K_2H_4As_2S_2O_6$.

Potassium disulphoxyarsenate, $K_2AsS_2O_7 + 10H_2O$.
Very hygroscopic; decomp. by H_2O . (Weinland, Z. anorg. 1897, 14. 63.)

Potassium trisulphoxyarsenate, $K_3AsS_3O + 7H_2O$.
Yellow oil which cryst. at -20° . (McCay and Foster, Z. anorg. 1904, 41. 468.)

Sodium monosulphoxyarsenate, $Na_2AsSO_4 + 12H_2O$.

Easily sol. in H_2O . (Preis, A. 257. 180.)
 (McLaughlan, B. 1901, 34. 2170.)

Sol. in H_2O . (Weinland, B. 1896, 29. 1009.)
Sl. efflorescent. Insol. in alcohol. (McCay, Z. anorg. 1902, 29. 42.)

Sol. in $NaOH + Aq$; decomp. by boiling with conc. $NaOH$. (Weinland, Z. anorg. 1897, 14. 49.)

Sodium hydrogen monosulphoxyarsenate, NaH_2AsSO_4 .

Decomp. by H_2O ; insol. in alcohol. (Weinland, Z. anorg. 1897, 14. 58.)

$Na_2HAsSO_4 + 8H_2O$. **Easily sol. in H_2O .** (Preis.)

Sodium disulphoxyarsenate, $Na_3AsS_2O_7 + 10H_2O$.

Easily sol. in H_2O . (Preis.)
Sol. in H_2O ; pptd. by alcohol. (McCay, B. 1899, 32. 2472.)

Not decomp. by boiling $NaOH + Aq$. (Weinland, Z. anorg. 1897, 14. 62.)

Insol. in alcohol. (McCay, Z. anorg. 1900, 25. 461.)

$+ 11H_2O$. (McLaughlan, B. 1901, 34. 2170.)

Insol. in alcohol. (McCay, Z. anorg. 1902, 29. 46.)

Sodium trisulphoxyarsenate, $Na_3AsS_3O + 11H_2O$.

Decomp. by H_2O . (McCay and Foster, Z. anorg. 1904, 41. 454.)

Sodium trisulphoxydiarsenate, $As_2O_3S_3, 3Na_2O + 24H_2O$.

Easily sol. in H_2O . (Geuther, A. 240. 208.)
 $2As_2O_3S_3, Na_2O + 7H_2O$. **Sol. in H_2O .** (Nilson, J. pr. (2) 14. 14.)

Correct composition is $Na_3As_{18}S_{24}O_7 + 30H_2O$. (Preis.)

Sodium sulphoxyarsenate, $Na_3As_{18}S_{24}O_7 + 30H_2O = 4Na_2O, 6As_2S_2, 3As_2S_4O + 30H_2O$.

Decomp. by H_2O . Sol. in NH_4OH or $KOH + Aq$. (Preis, A. 257. 187.)

= Sodium oxytrisulpharsenate of Nilson.

Sodium pentasulphoxytetraarsenate, $Na_{12}As_4S_5O_{11} + 48H_2O$.

Less sol. in H_2O than other sulphoxyarsenates. (Preis.)

Sodium strontium trisulphoxyarsenate, $NaSrAsS_3O + 10H_2O$.

Unstable. (McCay and Foster, Z. anorg. 1904, 41. 462.)

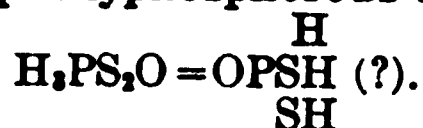
Trisulphoxyazotic acid, $\text{ON}(\text{SO}_3\text{H})_3$.

Known only in its salts. (Claus, A., 158. 52 and 194.)

Has the formula $(\text{SO}_3\text{H})_3\text{N} < \overset{\text{O}}{\underset{\text{O}}{\text{O}}} > \text{N}(\text{SO}_3\text{H})_3$.
(Raschig, A. 241. 161.)

Potassium trisulphoxyazotate, $\text{ON}(\text{SO}_3\text{K})_3 + \text{H}_2\text{O} = (\text{SO}_3\text{K})_3\text{N} < \overset{\text{O}}{\underset{\text{O}}{\text{O}}} > \text{N}(\text{SO}_3\text{K})_3$.

Easily sol. in H_2O without decomp., even on boiling. (Claus, A. 157. 210.)

Sulphoxyphosphorous acid,

See Thiophosphorous acid.

Sulphoxyvanadic acid.

Ammonium pyrohexasulphoxyvanadate, $(\text{NH}_4)_4\text{V}_2\text{S}_6\text{O}$.

Sol. in H_2O . (Krüss and Ohnmais, A. 263. 53.)

Potassium pyrohexasulphoxyvanadate, $\text{K}_4\text{V}_2\text{S}_6\text{O} + 3\text{H}_2\text{O}$.

Melts in crystal H_2O . (Krüss and Ohnmais.)

$\text{K}_2\text{V}_4\text{S}_{12}\text{O}_2 + 3\text{H}_2\text{O}$. More sol. in H_2O than preceding comp. (K. and O.)

Sodium orthotrisulphoxyvanadate, $\text{Na}_3\text{VS}_3\text{O} + 5\text{H}_2\text{O}$.

Very deliquescent, and easily sol. in H_2O . Somewhat sol. in alcohol. (Krüss and Ohnmais.)

Sodium orthomonosulphoxyvanadate, $\text{Na}_3\text{VSO}_3 + 10\text{H}_2\text{O}$.

Less sol. in H_2O than other sulphoxyvanadates. (K. and O.)

Sulphur, S.

The various modifications of sulphur have been classified in many different ways, and there is a difference of opinion as to whether certain forms are true allotropic modifications or not.

The data, as far as concerns the solubility, may be arranged as follows:—

A. Sol. in CS_2 . 1. Rhombic, octahedral, or alpha sulphur, ordinary sulphur. Easily sol. in CS_2 , etc. See below for solubility in various solvents.

2. Prismatic, monoclinic, or beta sulphur. Sol. in CS_2 , but is converted into A, 1. Prismatic sulphur obtained by melting brimstone is not wholly sol. in CS_2 on account of admixture of gamma sulphur.

Monoclinic modification is more sol. than rhombic in CHCl_3 , ether and benzene. (Meyer C. C. 1903, II. 481.)

3. Soft sulphur, milk of sulphur.

4. Amorphous sol. sulphur is also a separate modification, according to Berthelot.

B. Soft sulphur, obtained by strong heating and quickly cooling, is sol. in CS_2 , but becomes insol. therein by repeatedly dissolving and evaporating. More easily sol. in CS_2 than A, 1.

C. Insol. in CS_2 . 1. By action of strong light on S in CS_2 .

2. By heating to b.-pt., cooling suddenly, and allowing to stand until hard. Has been called gamma sulphur, but is a mixture of $\frac{2}{3}$ A, 4 and $\frac{1}{3}$ insol. S.

3. Insol. S in flowers of sulphur. Converted into A. 1 by standing 3 days with alcohol.

According to Berthelot (A. ch. (3) 49. 430) there are only two varieties of S. I. "Octahedral," II. "Amorphous."

I. *Octahedral*. Sol. in CS_2 . Scarcely acted upon by $\text{KHSO}_3 + \text{Aq}$. Converted by oxidising agents into II.

II. *Amorphous*. Insol. in neutral solvents, viz. H_2O , alcohol, ether, CS_2 , etc.

Sol. with tolerable rapidity in $\text{KHSO}_3 + \text{Aq}$. By long action of $\text{Na}_2\text{S} + \text{Aq}$, a portion is dissolved, and the remainder converted into I. Less easily oxidised by $\text{HNO}_3 + \text{Aq}$ than I. Some varieties of this modification are sol. to a certain extent in alcohol and ether, and by boiling the rest of the sulphur is converted into I; also by long-continued contact with cold alcohol. Berthelot holds that the modification is changed before dissolving. Solutions of the alkalis, alkali salts, and alkali sulphides change insol. into sol. sulphur (Berthelot.)

Elastic sulphur obtained by pouring molten sulphur at a temp. of over 260° into H_2O contains 35% or more of a modification of S which is insol. in CS_2 , hot or cold, but sol. in absolute alcohol; this modification can be converted back into ord. sulphur by heating to 100° . (Pelouze and Fremy.) (See C. 2.)

This modification can be obtained also by action of HCl on thiosulphates. (Förde and Gélis.)

The soft pasty sulphur obtained by decomposition of H_2S by SO_2 forms an almost clear emulsion (pseudo-solution) with H_2O , from which it is pptd. by various salts and substances which have no chemical affinity for it. 23 pts. S combine in this way with 100 pts. H_2O . When pptd. by saline solutions, some of the S remains in solution. When solution is exposed to the light, S gradually separates out; also on boiling the same takes place. The above pseudo-solution is pptd. by mineral acids, and the pptd. S may still be dissolved in fresh water, if not left in contact for some time with the acid. Also pptd. by K salts, with loss of power of forming pseudo-solutions. Pptd. by NH_4 and Na salts without losing that power.

Alkali hydrates, carbonates, or sulphides convert it into insol. S.

The solution may be mixed with alcohol without change. Decomp. by long shaking with naphtha or oil of turpentine. The pseudo-solution combines with CS_2 , forming an emulsion which subsequently decomposes. The S itself is only partially sol. in CS_2 . (Selmi, J. pr. 57. 49.)

By treatment of amorphous "insoluble" S with CS_2 or CCl_4 , a small part goes into solution, the amount being dependent on the time of contact with the temp., and nature of the solvent, but independent of the amount of the solvent. It is assumed that this is due to a partial change of the "insoluble" into soluble S. (Wigand, Z. phys. Ch. 1910, 75. 235.)

"Delta" sulphur. Partly sol. in H_2O . (Debus, Chem. Soc. 53. 18.)

A colloidal form wholly sol. in H_2O exists, which, however, decomposes very easily. (Engel, C. R. 112. 866.)

Black sulphur. Insol. in alcohol, ether, CS_2 , fatty oils even at 200° , cold alkali hydroxides + Aq, H_2SO_4 , HNO_3 , or aqua regia. (Knapp, J. pr. (2) 43. 305.)

Green modification. Five times more sol. than ordinary sublimed sulphur in a mixture of salicylaldehyde and benzene. (Orloff, C. C. 1902, I. 1264.)

The following data relate to octahedral or ordinary sulphur (A. 1):—

Sol. in warm liquid H_2S (Niemann); warm P_2S_5 , SBr_2 , SCl_2 , Br_2 , NCl_3 , BaS + Aq (Dumas); in alcoholic solution of K_2S_2 , but is reprecipitated by addition of H_2O to sat. solution.

Sol. in liquid SO_2 .

Sol. in aqueous solution of alkali sulphates, especially when hot. Sl. sol. in boiling conc. HSCN + Aq, from which it mostly separates on cooling.

Na_2CO_3 + Aq (5.6% Na_2CO_3) dissolves no S at 20° ; 0.06775% at 100° . (Pohl, Dingl. 197. 508.)

The solubility of S in Na_2S + Aq between 0° and 50° diminishes slightly with increase in temp., but increases with dilution of the solution, having its largest value in a N/16 solution of Na_2S + Aq when the relation of Na_2S to dissolved S equals about 1:4. (Küster, Z. anorg. 1905, 43. 56.)

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913, 84. 27.)

Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54. 674.)

Sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 822.)

Sol. in liquid NH_3 . 1 gr. S is sol. in 3–4 ccm. liquid NH_3 . (Hugot, A. ch. 1900, (7) 21. 32.)

The solubility of S in liquid NH_3 is constant from -23° to -84° and equals 39%. (Ruff, Z. angew. Ch. 1910, 23. 1830.)

Solubility in liquid NH_3 .
(g. S in 100 g. solution.)

t°	S	t°	S
—78	38.6	16.4	25.65
—20.5	38.1	30	21.0
0	32.34	40	18.5

(Ruff and Hecht, Z. anorg. 1911, 70. 62.)

Sl. sol. in liquid NO_2 . (Frankland, Chem. Soc. 1901, 79. 1361.)

S_2Cl_2 dissolves 66.74% S at ord. temp. to form a liquid of 1.7 sp. gr. (Rose.)

Solubility of S in S_2Cl_2 varies according to the variety of sulphur used. Aten has published an extended investigation on the subject, which see for details. (Z. phys. Ch. 1905–14, 54. 86, 124; 81. 268; 83. 443; 86. 1; 88. 321.)

Solubility in SnCl_4 .

100 g. SnCl_4 dissolve at:

99° 101° 110° 110°
5.8 6.2 8.7 9.1 pts. solid S,

112° 112° 121°
9.4 9.9 17.0 pts. liquid S.

(Gerardin.)

Sol. in alkalis + Aq with decomp.

Sol. in 1926.7 pts. absolute alcohol at 15° . (Pohl, W. A. B. 6. 600.)

Sol. in 20 pts. hot nearly absolute alcohol, less sol. in weaker alcohol. (Lauragais.)

Sol. in 600 pts. boiling alcohol of 40° B. (Chevallier, J. ch. méd. 2. 587); in 500 pts. alcohol (Meissner); 200 pts. alcohol (Pelouze and Fremy).

100 pts. absolute alcohol dissolve 0.42 pt. at b.-pt., and 0.12 pt. S at 16° ; 100 pts. ether dissolve 0.54 pt. at b.-pt., and 0.19 pt. S at 16° ; 100 pts. benzene dissolve 17.04 pts. at b.-pt., and 1.79 pts. S at 16° ; 100 pts. oil of turpentine dissolve 16.16 pts. at b.-pt., and 1.35 pts. S at 16° ; 100 pts. CS_2 dissolve 73.46 pts. at b.-pt., and 38.70 pts. S at 16° ; 100 pts. naphtha dissolve 10.56 pts. at b.-pt., and 2.77 pts. S at 16° ; 100 pts. tar-oil dissolve 26.98 pts. at b.-pt., and 1.51 pts. S at 16° . (Payen, C. R. 34. 456.)

100 pts. absolute methyl alcohol dissolve 0.028 pt. at 18.5° ; 100 pts. absolute ethyl alcohol dissolve 0.053 pt. at 18.5° . (de Bruyn, Z. phys. Ch. 10. 781.)

Solubility in amyl alcohol.

95° 110° 110°
1.5 2.1 2.2 pts. solid S,

112° 112° 120° 131°
2.6 2.7 3.0 5.3 pts. liquid S.

(Gerardin, A. ch. (4) 5. 134.)

Quickly sol. in 12.5 pts. ether. (Favre.)

100 pts. benzene dissolve 0.965 pt. S at 26° ; 100 pts. benzene dissolve 4.377 pts. S at 71° ; 100 pts. toluene dissolve 1.479 pts. S at 23° ; 100 pts. ethyl ether dissolve 0.972 pt. S at

23.5°; 100 pts. chloroform dissolve 1.205 pts. S at 22°; 100 pts. phenol dissolve 16.35 pts. S at 174°; 100 pts. aniline dissolve 85.27 pts. S at 130°. (Cossa, B. 1. 139.)

Solubility in benzene at t°.

t°	g. S in 10 g. of solution
15 17	0.1480
19.29	0.1692

(Brönsted, Z. phys. Ch. 1906, 55. 377.)

A mixture of S and toluene separates into two layers, containing 33 and 92.5% S respectively. (Haywood, J. phys. Ch. 1897, 1. 232.)

CS₂ dissolves 0.35 pt. ordinary sulphur; some varieties of S, however, are not entirely sol. in CS₂, thus—

Variety of Sulphur	Pts. sol. in 1 pt. CS ₂	Fraction of original wt. insol. in CS ₂
Octahedral, from Sicily	0.335	0.000
Crystallised in dry way, recently prepared	0.415	0.029
Do., prepared 8 years	0.33	0.004
Do., prepared 9 years		0.020
Do., prepared 15 years		0.051
Red needles, recently prepared	0.382	0.023
Soft yellow, do		0.353
Do., prepared 2 years	0.316	0.157
Soft red, recently prepared	0.374	0.157
Do., prepared 5 years		0.181
Flowers of sulphur	0.351	0.113
Do., another sample		0.234
Roll brimstone, outside		0.029
Do., inside		0.073

(Deville, A. ch. (3) 47. 99.)

The pt. insol. in CS₂ is sol. in hot absolute alcohol, crystallising on cooling; less sol. in chloroform or ether. (Deville.)

100 pts. pure CS₂ dissolve pts. S at t°.

t°	Pts. S	t°	Pts. S
11	16.54	22	46.05
- 6	18.75	38	94.57
0	23.99	48.5	146.21
+15	37.15	55	181.34
18.5	41.65		

(Cossa, B. 1. 138.)

Neither ordinary stick S nor flowers of S is completely sol. in CS₂. Pptd. S is completely sol. in 5 pts. CS₂. (Tittenger, C. C. 1894, II. 267.)

2.99 g. S are sol. in 100 grams CS₂ at -117°. (Arcetowski, C. R. 1895, 121. 124.)

Solubility in CS₂.

100 g. of the sat. solution contain at:

-77°	-84.5°	-89°	-116°
4.84	4.46	4.29	2.90 g. S.

(Arcetowski, Z. anorg. 1896, 11. 274.)

When 20 pts. S dissolve in 50 pts. CS₂ at 22° the temp. is lowered 5°. (Cossa.)

Sat. solution of S in CS₂ boils at 57°. (Cossa.)

Sp. gr. of S dissolved in CS₂ at 15°.

(Pts. S per 100 pts. CS₂.)

Sp. gr.	Pts. S	Sp. gr.	Pts. S	Sp. gr.	Pts. S
1.271	0.0	1.312	9.9	1.352	19.6
1.272	0.2	1.313	10.2	1.353	19.9
1.273	0.4	1.314	10.4	1.354	20.1
1.274	0.6	1.315	10.6	1.355	20.4
1.275	0.9	1.316	10.9	1.356	20.6
1.276	1.2	1.317	11.1	1.357	21.0
1.277	1.4	1.318	11.3	1.358	21.2
1.278	1.6	1.319	11.6	1.359	21.5
1.279	1.9	1.320	11.8	1.360	21.8
1.280	2.1	1.321	12.1	1.361	22.1
1.281	2.4	1.322	12.3	1.362	22.3
1.282	2.6	1.323	12.6	1.363	22.7
1.283	2.9	1.324	12.8	1.364	23.0
1.284	3.1	1.325	13.1	1.365	23.2
1.285	3.4	1.326	13.3	1.366	23.6
1.286	3.6	1.327	13.5	1.367	24.0
1.287	3.9	1.328	13.8	1.368	24.3
1.288	4.1	1.329	14.0	1.369	24.8
1.289	4.4	1.330	14.2	1.370	25.1
1.290	4.6	1.331	14.5	1.371	25.6
1.291	4.8	1.332	14.7	1.372	26.0
1.292	5.1	1.333	15.0	1.373	26.5
1.293	5.3	1.334	15.2	1.374	26.9
1.294	5.6	1.335	15.4	1.375	27.4
1.295	5.8	1.336	15.6	1.376	28.1
1.296	6.0	1.337	15.9	1.377	28.5
1.297	6.3	1.338	16.1	1.378	29.0
1.298	6.5	1.339	16.4	1.379	29.7
1.299	6.7	1.340	16.6	1.380	30.2
1.300	7.0	1.341	16.9	1.381	30.6
1.301	7.2	1.342	17.1	1.382	31.4
1.302	7.5	1.343	17.4	1.383	31.9
1.303	7.8	1.344	17.6	1.384	32.6
1.304	8.0	1.345	17.9	1.385	33.2
1.305	8.2	1.346	18.1	1.386	33.8
1.306	8.5	1.347	18.4	1.387	34.5
1.307	8.7	1.348	18.6	1.388	35.2
1.308	8.9	1.349	18.9	1.389	36.1
1.309	9.2	1.350	19.0	1.390	36.7
1.310	9.4	1.351	19.3	1.391	37.2
1.311	9.7				

(Mascagno, C. N. 42. 192.)

dissolved in CS_2 at 15° . Water at $4^\circ = 1$.

S	Sp. gr.	% S	Sp. gr.	% S
0	1.2736	0.6	1.2764	1.2
2	1.2745	0.8	1.2774	1.4
4	1.2755	1.0	1.2783	1.6
8	1.3096	8.0	1.3409	14.2
0	1.3105	8.2	1.3419	14.4
2	1.3115	8.4	1.3430	14.6
4	1.3125	8.6	1.3440	14.8
6	1.3135	8.8	1.3450	15.0
8	1.3145	9.0	1.3460	15.2
0	1.3155	9.2	1.3471	15.4
2	1.3165	9.4	1.3481	15.6
4	1.3175	9.6	1.3491	15.8
6	1.3185	9.8	1.3502	16.0
8	1.3195	10.0	1.3512	16.2
0	1.3205	10.2	1.3522	16.4
2	1.3215	10.4	1.3532	16.6
4	1.3226	10.6	1.3543	16.8
6	1.3236	10.8	1.3553	17.0
8	1.3246	11.0	1.3563	17.2
0	1.3256	11.2	1.3573	17.4
2	1.3266	11.4	1.3584	17.6
4	1.3277	11.6	1.3594	17.8
6	1.3287	11.8	1.3604	18.0
8	1.3297	12.0	1.3615	18.2
0	1.3307	12.2	1.3625	18.4
2	1.3317	12.4	1.3635	18.6
4	1.3328	12.6	1.3646	18.8
6	1.3338	12.8	1.3656	19.0
8	1.3348	13.0	1.3667	19.2
0	1.3358	13.2	1.3677	19.4
2	1.3368	13.4	1.3688	19.6
4	1.3379	13.6	1.3698	19.8
6	1.3389	13.8	1.3709	20.0
8	1.3399	14.0		

er, Z. anorg. 1897, 15. 200.)

tone. (Eidmann, C. C. 1899, II.

y of S in acetone + Aq at 25° .

als. g. S in 100 cc. of the solution.
one in 100 g. acetone + Aq.

S	Sp. gr.
65.0	0.78540
45.0	0.79114
33.0	0.81654
25.3	0.82958

(noch, Z. anorg. 1905, 45. 263.)

Solubility in organic solvents.

Solvent	t°	Sat. solution contains % S
CS_2	-61	3.6
	-55	4.4
	-19	10.6
	-18	10.8
	-17	11.5
	-13	12.4
	-11	13.3
	-11	13.5
	-2	17.2
	+3	19.5
	9	23.1
	11	23.7
	14	25.9
	17	27.2
	19	28.9
	20	28.5
	21	29.7
	26	33.4
	27	34.6
	29	37.8
	30 5	39.7
	33	42.2
	40	48.7
	44	53.2
	46	56.2
	48	57.5
	53	60.0
	54	60.6
	65	67.9
	77.5	76.4
	81.0	79.4
	92.0	87.8
	98.0	90.1
Ethylene dibromide	9	1.7
	22	2.4
	40	4.4
	50	6.4
	72	12.4
	95	30.2
	108	60.0
Benzene	8	1.2
	10	1.3
	21	1.8
	30	2.6
	39	3.3
	47	4.0
	54	4.9
	65	6.8
	72	8.6
	100	17.5
	123	31.9
	127	34.0
	150	36.8
Hexane	-20	0.07
	0	0.16
	+26	0.41
	+68	1.2
	+130	5.2
	+142	6.2
	+184	8.3

(Étard, A. ch. 1894, (7) 2. 571.)

Solubility in CHCl_3 at t° .

t°	g. S in 10 g. of solution
12.25	0.0744
19.29	0.0918

(Brönsted, Z. phys. Ch. 1906, 55. 377.)

Solubility of octahedral and prismatic S in organic solvents at t° .

Solvent	t°	% prismatic S	% octahedral S
Benzene	18.6	2.004	1.512
	25.3	2.335	1.835
Chloroform	0	1.101	0.788
	15.5	1.658	1.253
	40	2.9	2.4
Ethyl ether	0	0.113	0.080
	25.3	0.253	0.200
Ethyl bromide	0	0.852	0.611
	25.3	1.676	1.307
Ethyl formate	0	0.028	0.019
Ethyl alcohol	25.3	0.066	0.052

(Brönsted, Z. phys. Ch. 1906, 55. 377.)

Solubility in organic solvents at 25° . (G. S dissolved in 1 g. mol. of solvent.)

Solvent	g. S
Ethylene chloride	0.831
Tetrachlorethane	2.063
Dichlorethylene	1.237
Pentachlorethane	2.421
Trichlorethylene	2.43
Perchlorethylene	2.537
Carbon tetrachloride	1.354

(Hoffmann et al. 1910, B. 43. 188.)

100 g. trichlorethylene dissolve 1.19 g. S at 15° . (Wester and Bruins, Pharm. Weekbl. 1914, 51. 1443.)

Solubility in benzyl chloride at t° .

t°	g. S per 100 g. of solution	
	in upper layer	in lower layer
0	0.99	
17	1.78	
35	2.57	
46.1	3.64	
63.3	6.15	
78.0	9.88	
99.1	19.89	
109.6	...	90.62
114.6	...	87.99
118.8	37.29	...
121.4	40.04	85.02
130.0	49.71	80.07
134.2	56.20	72.23

Above 134.2° sulphur is miscible with

benzyl chloride in all proportions; at temp. two layers are formed.

(Bogusky, J. Russ. Phys. Chem. S 37. 92-99; C. C. 1905, I. 120)

Easily sol. in boiling acetic a (Rosenfeld, B. 13. 1475.)

Sol. in considerable amount in w $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, but very sl. sol. if (bermann, B. 10. 866.)

Sol. in stearic acid + Aq. (Vulp Pharm. (3) 13. 38.)

Acetic ether dissolves 6% S. (F. Difficultly sol. in methyl acetat mann, B. 1909, 42. 3790.)

Sol. in ethyl acetate. (Naumann 37. 3601.)

Sl. sol. in benzonitrile at ord. tem more sol. at higher temp. (Nau 1914, 47. 1369.)

Sol. in 12 pts. hot petroleum froe but nearly insol. in cold. (de Saum 100 pts. nicotine at 100° dissolve S, but this separates out as the solu (Klever, C. C. 1872. 434.)

Sol. in warm aniline. (Barral, 20. 352.)

Easily sol. in hot, less sol. in co (Fritzsche.)

Very sol. in aniline and quinoline, when warm. (Hofmann.)

Sol. in quinoline but reacts with t with evolution of H. (Beckmann a Z. anorg. 1906, 51. 236.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.03 6 days. (Gates, J. phys. Chem. 143.)

Sol. in 2.6 pts. of boiling, sl. a creosote.

Sol. by digestion in 2 pts. oil of t Sol. in hot oil of copaiba, crysta cooling.

Sol. in hot oil of mandarin, cr. on cooling.

Sol. in hot oil of caraway, crysta cooling.

Somewhat sol. in hot, less in a spirit.

Sl. sol. in lignone, bromoform, zene, but easily in hot benzene. (J Chem. Soc. 1. 262.)

Sol. in ethyl sulphide, and car ride. (Rathke, A. 152. 187.)

Sol. in mercuric methyl.

Sol. in 20 pts. ethyl nitrate, from is not pptd. by H_2O .

Sol. in naphtha, aldehyde, iodol chloroform, warm chloral, sinka ethyl chloride, warm benzoyl chlori 100 pts. methylene iodide dissolv S at 10° . Melted sulphur is misc hot methylene iodide. (Retgers, 3. 343.)

S dissolves in 2000 pts. glycerin and Garot, J. Pharm. (3) 26. 81.)

Glycerine dissolves 0.10% S. (Kle 1872. 434.)

100 g. glycerine dissolve 0.14 g. at 15.5°. (Ossendowski, Pharm. J. 1907, 79. 575.)

Sol. in butyl sulphhydrate, and warm retinole.

Sol. in ethyl sulphhydrate.

Very sol. in coniine, hexyl alcohol, warm allyl sulphocyanide, cacodyl oxide. Somewhat sol. in hot styrene, separating out on cooling.

Readily sol. in warm, less readily in cold toluene or resin-oil.

Sol. in olive oil at 115°, from which it mostly separates on cooling.

Sol. in hot oil of amber, crystallising upon cooling. Sol. in 2 pts. hot, sl. sol. in cold mouthchin.

Insol. in valerianic acid, amyl valerate, amyl hydride.

Linseed oil dissolves % S at t°.

t°	% S	t°	% S	t°	% S
25	0.630	95	2.587	160	9.129
60	1.852	130	4.935

(Pohl.)

Solubility in olive oil (sp. gr. = 0.885).
100 pts. dissolve pts. S at t°.

t°	Pts. S	t°	Pts. S	t°	Pts. S
15	2.3	65	20.6	110	30.3
40	5.6	100	25.0	130	43.2

(Pelouze, C. R. 68. 1179.)

Solubility in 100 pts. coal-tar oil at t°.

t°	Pts. S in		
	Oil of 0.870 sp. gr. B.-pt. 80-100°	Oil of 0.880 sp. gr. B.-pt. 85-120°	Oil of 0.882 sp. gr. B.-pt. 120-200°
15	2.1	2.3	2.5
30	3.0	4.0	5.3
50	5.2	6.1	8.3
80	11.8	13.7	15.2
100	15.2	18.7	23.0
110	...	23.0	26.2
120	...	27.0	32.0
130	38.7

t°	Pts. S in		
	Oil of 0.885 sp. gr. B.-pt. 150-200°	Oil of 1.010 sp. gr. B.-pt. 210-300°	Oil of 1.020 sp. gr. B.-pt. 220-300°
15	2.6	6.0	7.0
30	5.8	8.5	8.5
50	8.7	10.0	12.0
80	21.0	37.0	41.0
100	26.4	52.5	54.0
110	31.0	105.0	115.0
120	38.0	∞	∞
130	43.8	∞	∞

(Pelouze, C. R. 69. 56.)

Sulphur bromide, S₂Br₂.

Decomp. gradually with H₂O. Dissolves S on warming, which crystallises out on cooling. Sol. in CS₂.

Decomp. by current of dry air into S and Br. (Hannay, Chem. Soc. 35. 16.)

Decomp. slowly by cold H₂O, rapidly by hot H₂O. Decomp. by dil. KOH+Aq or NaHCO₃+Aq. (Korndörfer, Arch. Pharm. 1904, 242. 156.)

A study of the mpt. curve of a series of mixtures of sulphur and bromine gave no evidence for the existence of the compounds SBr₂ and SBr₄. (Ruff, B. 1903, 36. 2446.)

Sulphur monochloride, S₂Cl₂.

Slowly decomp. by H₂O. Miscible with CS₂ and C₆H₆. Sol. in alcohol and ether with subsequent decomposition. Sol. in oil of turpentine.

Moderately sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 830.)

Sol. in CCl₄, and C₆H₆. (Oddo, Gazz. ch. it. 1899, 29. (2) 318.)

Sulphur dichloride, SCl₂.

Decomp. slowly with H₂O, immediately by alcohol or ether.

Sulphur tetrachloride, SCl₄.

Violently decomp. by H₂O. Decomp. at temperatures above -22°. (Michaelis, A. 170. 1.)

Sulphur stannic chloride, 2SCl₄, SnCl₄.

Decomp. by H₂O. Sol. in dil. HNO₃+Aq. Forms a mass with fuming HNO₃, which is sol. in HNO₃+Aq. Sol. in POCl₃. (Casselmann.)

Very hygroscopic. Fumes in moist air. Very easily sol. in dry abs. ether and in benzene. Sol. in CHCl₃, SO₂Cl₂, CS₂, POCl₃, ligroin and petroleum ether. (Ruff, B. 1904, 37. 4517.)

Sulphur titanium chloride, SCl₄, 2TiCl₄.

Very deliquescent. Easily sol. in dil. HNO₃+Aq. (Weber, Pogg. 132. 454.)

SCl₄, TiCl₄. Sol. in SO₂Cl₂, CHCl₃, CS₂, and petroleum ether. (Ruff, B. 1904, 37. 4516.)

Sulphur chloride ammonia, S₂Cl₂, 4NH₃.

Insol. in H₂O, but gradually decomp. thereby; sol. without decomp. in absolute alcohol, from which it is pptd. by H₂O. (Mertens.)

Does not exist. (Fordos and Gélis, C. R. 31. 702.)

SCl₂, 2NH₃. Decomp. by H₂O. Sol. in alcohol or ether. (Soubeiran, A. ch. 67. 71.) Not a true chemical compound, but a mixture. (Fordos and Gélis, C. R. 31. 702.)

SCl2, 4NH3. Decomp. by H2O. Sl. sol. in absolute alcohol and ether (Soubeiran, A. ch. 67. 71); mixture (Fordos and Gélis).

Sulphur chloride nitrogen sulphide.
See Nitrogen sulphochloride.

Sulphur perfluoride, SF6.
Very sl. sol. in H2O; sl. sol. in alcohol. (Moissan, C. R. 1900, 130. 868.)

Sulphur monoiodide, S2I2.
Insol. in H2O. Decomp. by alcohol, which dissolves out I2. Sl. sol. in cold caoutchin, the solution decomposing when boiled. Freely sol. in glycerine. Sol. in 60 pts. glycerine, and 82 pts. olive oil. (Cap and Garot, J. Pharm. (3) 26. 81.)
Very sol. in liquid NH3. (Franklin, Am. Ch. J. 1898, 20. 830.)
Sol. in CS2. (Linebarger, Am. Ch. J. 1895, 17. 58.)

Sulphur hexiodide, SI6.
Decomp. on air. Alcohol or alkalis dissolve out iodine. (vom Rath, Pogg. 110. 116.)
Does not exist. (M'Leod, Rep. Brit. Assn. Advn. Sci. 1892. 690.)

Sulphur stannic iodide.
See Tin sulphur iodide.

Sulphur sesquioxide, S2O3.
Deliquescent. Violently decomp. by H2O at ordinary temp. Sol. in fuming H2SO4. Insol. in SO2. Decomp. by alcohol or other. (Weber, Pogg. 156. 531.)

Sulphur dioxide, SO2.
Liquid. Insol. in H2O if brought in contact therewith below the b.-pt. of SO2.
Sol. in 3 vols. CS2 on warming, separating out on cooling. Dissolves some P, little S, and no sulphuric or phosphoric acids.
Dissolves ether, chloroform, P, Br, S, I, CS2, colophonium, and other gums; also benzene when warmed. (Sestini, Bull. Soc. (2) 10. 226.)
Miscible with liquid SO2, but not with H2SO4.
Gas.
1 vol. H2O absorbs 30 vols. SO2 gas at 18° (Davy); 20 vols. at ord. temp. (Dalton); 43.78 vols. at ord. temp. (de Saussure); 50 vols. at 20° and 760 mm. (Pelouze and Fremy); 33 vols. at ord. temp. (Thomson).
1 pt. SO2 (by weight) is sol. in 0.1429 pt. H2O at 5°, and the solution has 1.020 sp. gr.
1 pt. SO2 is sol. in 0.0400 pt. H2O at ord. temp. (Priestley); in 0.0009 pt. H2O at 16°, and sp. gr. of the solution = 1.0513 (Thomson).
Sol. in 2 pts. H2O at 10°. (Pierre, A. ch. (3) 23. 421.)
100 vols. H2O at 18° and 760 mm. absorb 4378 vols. SO2 gas; 100 vols. alcohol of 0.84 sp. gr. at 760 mm. absorb 11,577 vols. (de Saussure, 1814.)

Solubility of SO2 gas in H2O. t° = temp.; V = vols. SO2 reduced to 0° and 760 mm. contained in 1 vol. sat. SO2 + Aq; V_1 = vols. SO2 gas reduced to 0° and 760 mm. dissolved by 1 vol. H2O under 760 mm. pressure.

t°	V	V_1	t°	V	V_1
0	68.861	79.789	21	34.986	37.970
1	67.003	77.210	22	33.910	36.617
2	65.169	74.691	23	32.847	35.302
3	63.360	72.230	24	31.800	34.026
4	61.576	69.828	25	30.766	32.786
5	59.816	67.485	26	29.748	31.584
6	58.080	65.200	27	28.744	30.422
7	56.369	62.973	28	27.754	29.314
8	54.683	60.805	29	26.788	28.250
9	53.021	58.697	30	25.819	27.161
10	51.383	56.647	31	24.873	26.151
11	49.770	54.655	32	23.942	25.178
12	48.182	52.723	33	23.025	24.244
13	46.618	50.849	34	22.122	23.347
14	45.079	49.033	35	21.234	22.489
15	43.564	47.276	36	20.361	21.666
16	42.073	45.578	37	19.502	20.866
17	40.608	43.939	38	18.658	20.141
18	39.165	42.360	39	17.827	19.435
19	37.749	40.838	40	17.013	18.768
20	36.206	39.374

(Schönfeld, A. 95. 5.)

This table may be formulated as follows:
1 vol. H2O absorbs $79.789 - 2.6077t + 0.029349t^2$ vols. SO2 at temp. between 0° and 20°, or 1 vol. sat. solution contains $68.861 - 1.87025t + 0.01225t^2$ vols. SO2. Coefficient of absorption between 21° and 40° = $75.192 - 2.1716t + 0.01903t^2$ vols. SO2 or 1 vol. sat. solution between 21° and 40° contains $60.952 - 1.38898t + 0.00726t^2$ vols. SO2.

Solubility of SO2 in H2O at various temps. and 760 mm. t° = temp.; G = grammes SO2 dissolved in 1 g. H2O; V = vols. SO2 dissolved in 1 g. H2O.

t°	G	V	t°	G	V
8	0.168	58.7	30	0.078	27.3
10	0.154	53.9	32	0.073	25.7
12	0.142	49.6	34	0.069	24.3
14	0.130	45.6	36	0.065	22.8
16	0.121	42.2	38	0.062	21.6
18	0.112	39.3	40	0.058	20.4
20	0.104	36.4	42	0.055	19.3
22	0.098	34.2	44	0.053	18.4
24	0.092	32.3	46	0.050	17.4
26	0.087	30.5	48	0.047	16.4
28	0.083	28.9	50	0.045	15.6

(Sims, A. 113. 340.)

— solubility of SO_2 in H_2O at various pressures.
 — P = "partial pressure," i. e. the total pressure minus the tension of aqueous vapour at given temp.; G at P = weight SO_2 in grammes, which is dissolved in 1 g. H_2O at pressure P ; G at 760 = calculated weight SO_2 that would be contained in 1 g. H_2O at 760 mm. if the absorption were proportional to the pressure; V = the volume of G grammes of SO_2 at 0° and 760 mm.

P	7°			
	G at P	G at 760	V at P	V at 760
30	0.010	0.263	3.634	92.06
40	0.013	0.242	4.451	84.55
50	0.015	0.223	5.129	77.95
60	0.017	0.818	6.024	76.28
70	0.020	0.213	6.868	74.55
80	0.022	0.210	7.743	73.55
90	0.025	0.208	8.598	72.62
100	0.027	0.205	9.421	71.60
120	0.032	0.201	11.09	70.20
140	0.036	0.197	12.71	69.00
160	0.041	0.195	14.34	68.15
180	0.046	0.193	15.97	67.40
200	0.050	0.191	17.59	66.83
220	0.055	0.190	19.19	66.30
240	0.059	0.188	20.79	65.84
260	0.064	0.187	22.40	65.44
280	0.069	0.186	23.99	65.10
300	0.073	0.185	25.59	64.81
350	0.085	0.184	29.55	64.16
400	0.096	0.182	33.51	63.65
450	0.107	0.181	37.44	63.25
500	0.118	0.180	41.42	62.94
550	0.130	0.179	45.31	62.60
600	0.141	0.178	49.20	62.32
650	0.152	0.178	53.10	62.09
700	0.163	0.177	56.98	61.86
750	0.174	0.176	60.88	61.69
760	0.176	0.176	61.65	61.65
800	0.185	0.176	64.74	61.50
850	0.196	0.175	68.57	61.30
900	0.207	0.175	72.41	61.15
950	0.218	0.175	76.25	61.00
1000	0.229	0.174	80.01	60.88
1050	0.240	0.174	83.97	60.77
1100	0.251	0.174	87.80	60.65
1200	0.273	0.173	95.45	60.45
1300	0.295	0.172	103.00	60.25

P	20°			
	G at P	G at 760	V at P	V at 760
40	0.007	0.143	2.637	50.09
50	0.009	0.138	3.171	48.20
60	0.011	0.135	3.718	47.10
70	0.012	0.131	4.205	45.64
80	0.013	0.127	4.663	44.30
90	0.015	0.125	5.169	43.65
100	0.016	0.124	5.692	43.25
120	0.019	0.121	6.683	42.33
140	0.022	0.119	7.690	41.75
160	0.025	0.118	8.711	41.17
180	0.028	0.117	9.652	40.75
200	0.030	0.116	10.62	40.35
220	0.033	0.115	11.59	40.03
240	0.036	0.114	12.54	39.70
260	0.038	0.112	13.45	39.30
280	0.041	0.112	14.41	39.10
300	0.044	0.111	15.34	38.87
350	0.050	0.110	17.66	38.35
400	0.059	0.109	20.56	38.10
450	0.064	0.108	22.37	37.77
500	0.071	0.107	24.67	37.50
550	0.077	0.106	26.93	37.20
600	0.083	0.105	29.14	36.90
650	0.090	0.105	31.39	36.70
700	0.096	0.105	33.62	36.50
750	0.103	0.104	35.94	36.43
760	0.104	0.104	36.43	36.43
800	0.110	0.104	38.32	36.40
1000	0.137	0.104	47.85	36.37
1200	0.178	0.104	62.10	36.31
1600	0.218	0.104	76.35	36.27
1900	0.259	0.104	90.53	36.21

P	30.8°			
	G at P	G at 760	V at P	V at 760
200	0.016	0.062	5.675	21.57
300	0.024	0.061	8.368	21.20
400	0.031	0.060	11.03	20.95
500	0.039	0.059	13.67	20.77
600	0.047	0.059	16.29	20.64
760	0.059	0.059	20.50	20.50
800	0.062	0.059	21.58	20.50
1000	0.077	0.058	26.84	20.40
1500	0.113	0.057	39.65	20.09
2000	0.149	0.057	52.11	19.80

P	50°			
	G at P	G at 760	V at P	V at 760
400	0.012	0.045	4.156	15.97
400	0.024	0.045	8.275	15.72
600	0.035	0.045	12.36	15.65
700	0.045	0.045	15.62	15.62
800	0.047	0.045	16.43	15.60
1000	0.059	0.045	20.51	15.59
1500	0.088	0.044	30.73	15.57
2000	0.012	0.044	39.07	15.55

(Sims, A. 118. 340.)

1 g. H₂O dissolves 0.0909 g. SO₂ = 34.73 cc. (at 25°) at 25° and 748 mm. pressure. (Walden and Centnerszwer, Z. phys. Ch. 1901, 42. 462.)

Solubility of SO₂ in H₂O at t° and 760 mm. pressure.

t°	G. SO ₂ per 1 g. H ₂ O	t°	G. SO ₂ per 1 g. H ₂ O
0	0.236	7	0.176
2	0.218	8	0.168
4	0.201	10	0.154
6	0.184	12	0.142

(Roozeboom, R. t. c. 1884, 3. 29.)

From a gas containing 10% by vol. of SO₂ at 10°, 1.63% by wt. is dissolved by 1 litre of H₂O; if the pressure is increased to 5 atmospheres, 8.14% by wt. is dissolved. (Harpf, Chem. Zeitschr., 1905, 4. 136.)

Solubility of SO₂ in H₂O at t°.

C = g. SO₂ in 1 cc. of the solution.

P = Pressure in mm. of Hg.

t°	C	P	$\frac{C}{P} \times 10^4$
0	0.000537	0.4	13.4
"	0.00237	3.5	6.78
"	0.01227	29.4	4.17
"	0.03804	109.4	3.48
25	0.000534	1.4	3.81
"	0.00234	11.75	2.00
"	0.01212	87.9	1.379
"	0.03750	313.0	1.198
50	0.000525	4.9	1.07
"	0.002276	30.5	0.746
"	0.01181	204.5	0.577
"	0.03628	696.0	0.521

(Lindner, M. 1912, 33. 645.)

Sp. gr. of sat. solution at—

0° 10° 20° 40°

1.06091 1.05472 1.02386 0.95548

(Bunsen and Schönfeld, A. 95. 2.)

Sat. SO₂ + Aq has sp. gr. = 1.0040. (Berthollet.)

Sp. gr. of sat. SO₂ + Aq at t°.

t°	Sp. gr.	t°	Sp. gr.	t°	Sp. gr.
0	1.0609	9	1.0548	17	1.0358
1	1.0596	10	1.0547	18	1.0321
2	1.0585	11	1.0528	19	1.0281
3	1.0576	12	1.0505	20	1.0239
4	1.0569	13	1.0481	21	1.0195
5	1.0562	14	1.0454	22	1.0147
6	1.0557	15	1.0424	23	1.0099
7	1.0552	16	1.0392	24	0.9991
8	1.0549				

(Schiff, A. 107. 312.)

Sp. gr. of SO₂ + Aq at 4°.

% SO ₂	Sp. gr.	% SO ₂	Sp. gr.	% SO ₂	Sp. gr.
1	1.0024	8	1.0217	15	1.0445
2	1.0049	9	1.0247	16	1.0480
3	1.0075	10	1.0278	17	1.0517
4	1.0102	11	1.0311	18	1.0553
5	1.0130	12	1.0343	19	1.0591
6	1.0158	13	1.0376	20	1.0629
7	1.0187	14	1.0410	21	1.0667

(Schiff, calculated by Gerlach, Z. anal. & 292

Sp. gr. of SO₂ + Aq.

% SO ₂	Temp.	Sp. gr.
0.99	15.5°	1.0051
2.05	"	1.0102
2.87	"	1.0148
4.04	"	1.0204
4.99	"	1.0252
5.89	"	1.0297
7.01	"	1.0353
8.08	"	1.0399
8.68	"	1.0439
9.80	"	1.0492
10.75	"	1.0541
11.65	12.5°	1.0597
13.09	11.0°	1.0668

(Giles and Schearer, Jour. Soc. Ch. Ind. 4 303.)

Sp. gr. of SO₂ + Aq.

% SO ₂	Sp. gr.	% SO ₂	Sp. gr.	% SO ₂	Sp. gr.
1	1.0052	4	1.0167	7	1.0283
2	1.0094	5	1.0208	8	1.0329
3	1.0134	6	1.0242	9	1.0402

(Anthon.)

Sp. gr. of SO₂ + Aq.

% SO ₂	Sp. gr.	% SO ₂	Sp. gr.	% SO ₂	Sp. gr.
1	1.0042	5	1.0210	8	1.0348
2	1.0083	6	1.0252	9	1.0392
3	1.0125	7	1.0295	10	1.0438
4	1.0167

(Hager, Adjumenta varia, Leipzig. 1871 146.)

Sp. gr. of SO₂ + Aq at 15°.

% SO ₂	Sp. gr.	% SO ₂	Sp. gr.	% SO ₂	Sp. gr.
0.5	1.0028	4.0	1.0221	7.5	1.0401
1.0	1.0056	4.5	1.0248	8.0	1.0426
1.5	1.0085	5.0	1.0275	8.5	1.0450
2.0	1.0113	5.5	1.0302	9.0	1.0474
2.5	1.0141	6.0	1.0328	9.5	1.0497
3.0	1.0168	6.5	1.0353	10.0	1.0520
3.5	1.0194	7.0	1.0377

(Scott, Polyt. Centralbl. 1873. 826.)

Conc. H₂SO₄ absorbs 0.009 pt. by weight (58 vols.), and SO₂ is more soluble in dilute H₂SO₄ + Aq, the more H₂O there is present. (Kolb, Dingl. 209. 270.)

Solubility in H₂SO₄.

Sp. gr. of H ₂ SO ₄	Absorbs SO ₂ per kg.	Absorbs SO ₂ per litre
1.841	0.009	5.8
1.839	0.014	8.9
1.540	0.021	11.2
1.407	0.032	15.9
1.227	0.068	29.7
1.020	0.135	49.0

(Kolb, Bull. Soc. Ind. Mullhouse, 1872. 224.)

Coefficient of absorption for H₂SO₄ (1.841 sp. gr. at 15° and 760 mm.) is 28.14 at 17°, and 28.86 at 16°. (Dunn, C. N. 43. 121.)

Solubility of SO₂ in H₂SO₄ of 1.84 sp. gr.

t°	Sp. gr. of sat. solution	Coeff. of absorption (760 mm.)	t°	Sp. gr. of sat. solution	Coeff. of absorption (760 mm.)
0	...	53	50	1.8186	9.5
10	1.8232	35.0	60	1.8165	7.0
20	1.8225	25.0	70	1.8140	5.5
25	1.8221	21.0	80	1.8112	4.5
30	1.8216	18.0	90	1.8080	4.0
40	1.8205	13.0

(Dunn, C. N. 1882, 45. 272. Calc. by Seidell, Solubilities, 1st Ed.)

Solubility of SO₂ in H₂SO₄ + Aq.

t°	Sp. gr. of H ₂ SO ₄ solution	Approx. % H ₂ SO ₄	Coeff. of absorption	t°	Sp. gr. of H ₂ SO ₄ solution	Approx. % H ₂ SO ₄	Coeff. of absorption
6.9	1.139	20	48.67	15.2	1.173	25	31.82
6.9	1.300	40	45.38	16.8	1.151	21	31.56
8.6	1.482	58	39.91	14.8	1.277	36	30.41
9.8	1.703	78	29.03	15.1	1.458	56	29.87
5.5	..067	10	36.78	15.6	1.609	70	25.17
6.6	1.102	15	34.08	15.0	1.739	81	20.83

(Dunn, C. N. 1882, 45. 272; Seidell, Solubilities, 1st Ed.)

Coefficient of absorption in H₂SO₄ (sp. gr. = 1.841) = 5.8; (sp. gr. = 1.839) = 8.9. (Lunge.)

Solubility in salts + Aq at 35°.

l = coefficient of absorption of SO₂ in the given salt solution at 35°.
lo = coefficient of absorption of SO₂ in water at 35° = 22.43.

Salt		3-normal	2.5-normal	2-normal	1.5-normal	1-normal	0.5-normal
KI	l	45.43	41.87	38.04	34.64	30.25	26.30
	l-lo	23.00	19.44	15.61	12.21	7.82	3.87
KBr	l	36.14	34.12	31.93	29.64	27.49	24.83
	l-lo	13.71	11.69	9.50	7.21	5.01	2.40
KCl	l	30.02	28.93	27.94	26.54	25.15	23.74
	l-lo	7.59	6.50	5.31	4.11	2.72	1.81
KCNS	l	42.94	38.13	35.05	32.03	28.79	25.63
	l-lo	18.51	15.70	12.62	9.60	6.36	3.20
NH ₄ NO ₃	l	27.43	26.66	25.57	24.78	24.23	23.35
	l-lo	5.00	4.23	3.14	2.35	1.80	0.92
KNO ₃	l	27.33	26.54	25.72	24.79	24.03	23.27
	l-lo	4.90	4.11	3.29	2.36	1.60	0.84

Solubility in salts + Aq at 35°.—Continued.

l = coefficient of absorption of SO₂ in the given salt solution at 35°.
 lo = coefficient of absorption of SO₂ in water at 35° = 22.43.

Salt		3-normal	2.5-normal	2-normal	1.5-normal	1-normal	0.5-normal
$\frac{1}{2}(\text{NH}_4)_2\text{SO}_4$	l	24.60	24.23	23.93	23.49	23.14	22.91
	l-lo	2.17	1.80	1.50	1.06	0.71	0.48
$\frac{1}{2}\text{CdI}_2$	l	24.30	23.99	23.71	23.38	23.06	22.75
	l-lo	1.87	1.56	1.28	0.95	0.63	0.32
$\frac{1}{2}\text{Na}_2\text{SO}_4$	l	19.27	19.79	20.20	20.81	21.35	21.88
	l-lo	-3.16	-2.64	-2.23	-1.62	-1.08	-0.55
$\frac{1}{2}\text{CdBr}_2$	l	19.17	19.70	20.60	20.81	21.46	21.88
	l-lo	-3.26	-2.73	-1.83	-1.62	-0.97	-0.55
$\frac{1}{2}\text{CdCl}_2$	l	18.68	19.23	20.02	20.55	21.23	21.73
	l-lo	-3.75	-3.20	-2.41	-1.88	-1.20	-0.70
$\frac{1}{2}\text{CdSO}_4$	l	16.25	17.41	18.31	19.42	20.43	21.45
	l-lo	-6.81	-5.02	-4.12	-3.01	-2.00	-0.98

Solubility in salts + Aq at 25°.

l = coefficient of absorption of SO₂ in the given solution at 25°.
 lo = coefficient of absorption of SO₂ in water at 25° = 32.76.

Salt		3-normal	2.5-normal	2-normal	1.5-normal	1-normal	0.5-normal
KI	l	68.36	62.63	56.75	50.58	44.76	38.66
	l-lo	35.60	29.87	23.99	17.82	12.00	5.90
$\frac{1}{2}\text{CdI}_2$	l	35.77	34.98	34.74	34.16	33.76	33.27
	l-lo	3.01	2.22	1.98	1.40	1.00	0.51
NH ₄ Br	l	52.25	49.17	46.06	42.78	39.46	36.28
	l-lo	19.49	16.41	13.30	10.02	6.70	3.52
KBr	l	52.26	48.87	44.96	42.41	39.11	35.94
	l-lo	19.00	15.71	12.70	9.15	6.35	3.18
NaBr	l	37.74	36.84	36.26	35.27	34.54	33.76
	l-lo	4.98	4.08	3.50	2.51	1.78	1.00
$\frac{1}{2}\text{CdBr}_2$	l	27.46	28.15	29.27	30.17	31.01	31.91
	l-lo	-5.30	-4.61	-3.49	-2.59	-1.75	-0.85
NH ₄ Cl	l	42.78	41.37	39.76	38.06	36.37	34.58
	l-lo	10.02	8.61	7.00	5.30	3.61	1.80
KCl	l	42.27	40.96	39.32	37.76	36.05	34.42
	l-lo	9.51	8.20	6.56	5.00	3.29	1.66

Solubility in salts + Aq at 25°—Continued.

coefficient of absorption of SO₂ in the given solution at 25°.
coefficient of absorption of SO₂ in water at 25°= 32.76.

Salt		3-normal	2.5-normal	2-normal	1.5-normal	1-normal	0.5-normal
NaCl	1	31.36	31.51	31.76	31.96	32.25	32.46
	1-lo	—1.40	—1.25	—1.00	—0.80	—0.51	—0.30
CdCl ₂	1	26.06	27.09	28.16	29.46	30.55	31.66
	1-lo	—6.70	—5.67	—4.60	—3.30	—2.21	—1.10
H ₄ CNS	1	61.46	57.01	52.26	47.26	42.74	37.78
	1-lo	28.70	24.25	19.50	14.50	9.98	5.02
KCNS	1	61.26	55.87	51.86	47.02	42.38	37.57
	1-lo	28.50	23.11	19.10	14.26	9.62	4.81
CaCNS	1	48.34	45.86	43.37	40.78	38.24	35.44
	1-lo	15.58	13.10	10.61	8.02	5.48	2.68
H ₄ NO ₃	1	39.14	38.01	37.27	36.28	35.07	33.96
	1-lo	6.38	5.25	4.51	3.52	2.31	1.20
KNO ₃	1	38.52	37.57	36.66	35.77	34.79	33.80
	1-lo	5.76	4.81	3.90	3.01	2.03	1.04
(NH ₄) ₂ SO ₄	1	35.96	35.47	34.95	34.34	33.82	33.35
	1-lo	3.20	2.71	2.19	1.58	1.06	0.59
K ₂ SO ₄	1	33.61	33.20
	1-lo	0.85	0.48
Na ₂ SO ₄	1	28.44	28.66	29.51	30.45	31.14	31.96
	1-lo	—4.32	—4.10	—3.25	—2.31	—1.62	—0.80
CdSO ₄	1	23.76	25.14	26.58	28.24	29.71	31.11
	1-lo	—9.00	—7.62	—6.18	—4.52	—3.05	—1.85

(Fox, Z. phys. Ch. 1902, 41. 462.)

in Cl₂ + Aq. Sol. in Br₂. Solidification curves determined. (Van der Goot, Z. phys. 13, 84. 419.)

Solubility of SO₂ in alcohol. 1 vol. alcohol at t° and 760 mm. dissolves V vols. SO₂ gas at 0° and 760 mm.

t°	V	t°	V	t°	V
0	328.62	9	201.33	17	130.61
1	311.98	10	190.31	18	124.58
2	295.97	11	179.91	19	119.17
3	280.58	12	170.13	20	114.48
4	265.81	13	160.98	21	110.22
5	251.67	14	152.45	22	106.68
6	238.16	15	144.55	23	103.77
7	225.26	16	137.27	24	101.47
8	212.98

(Bunsen's Gasometry.)

100 pts. absolute methyl alcohol dissolve 247 pts. SO₂ at 0° and 760 mm.; 47 pts. at 26° and 760 mm.; 100 pts. absolute ethyl alcohol dissolve 115 pts. SO₂ at 0° and 760 mm.; 32.3 pts. at 26° and 760 mm. (de Bruyn, Z. phys. Ch. 10. 783.)

Sol. in ether.
Absorbed by oil of turpentine.
Rapidly absorbed by anhydrous aldehyde in the cold, 11 pts. aldehyde absorbing 19 pts. SO₂.

Absorption coefficient of aldehyde for SO₂ is 1.4 times greater than that of alcohol, and 7 times greater than that of H₂O. (Geuther and Cartmell, Proc. Roy. Soc. 10. 111.)

1 pt. camphor dissolves 0.880 pt. by weight (=308 vols.) SO₂ at 0° and 725 mm.; 1 pt. glacial HC₂H₃O₂ dissolves 0.961 pt. by weight (=318 vols.) SO₂ at 0° and 725 mm.; 1 pt. formic acid dissolves 0.821 pt. by weight (=351 vols.) SO₂ at 0° and 725 mm.; 1 pt. acetone dissolves 2.07 pts. by weight (=589 vols.) SO₂ at 0° and 725 mm.; 1 pt. sulphuryl chloride dissolves 0.323 pt. by weight (=187 vols.) SO₂ at 0° and 725 mm. (Schulze, J. pr. (2) 24. 168.)

Solubility of SO₂ in CHCl₃.

C = g. SO₂ in 1 cc. of the solution.

P = Pressure in mm. Hg.

t°	C	P	$\frac{C}{P} \times 10^4$
0	0.000701	2.7	2.6
"	0.001790	5.6	3.14
"	0.006982	22.0	3.17
"	0.03097	90.2	3.43
"	0.08217	219.6	3.74
25	0.000669	5.7	1.17
"	0.001712	12.9	1.37
"	0.006723	48.0	1.40
"	0.02954	206.2	1.47
"	0.07839	488.8	1.60

(Lindner, M. 1912, 33. 645.)

Distribution of SO₂ between H₂O and CHCl₃ at 20°.

c₁ = g. SO₂ per l. of H₂O solution.

c₂ = g. SO₂ per l. of CHCl₃ solution.

C ₁	C ₂	$\frac{C_1}{C_2}$
1.738	1.123	1.55
1.753	1.122	1.56
2.326	1.704	1.37
2.346	1.703	1.38
2.628	1.897	1.38
3.039	2.395	1.27
3.058	2.385	1.28
3.686	3.063	1.21
3.735	3.062	1.22
4.226	3.626	1.17
5.269	4.798	1.10
5.372	4.813	1.12
6.588	6.183	1.07
31.92	33.84	0.94
33.26	37.25	0.89

(McCrae, Z. anorg. 1903, 35. 12.)

Distribution of SO₂ between HCl + Aq and CHCl₃ at 20°.

c₁ = g. SO₂ per l. of HCl + Aq solution

c₂ = g. SO₂ per l. of CHCl₃ solution.

HCl = normality of HCl + Aq used.

HCl	C ₁	C ₂	$\frac{C_1}{C_2}$
0.05-N	1.86	1.46	1
	3.076	2.830	1
	4.277	4.07	1
	5.340	5.42	0
0.1 -N	1.25	1.41	0
	1.324	1.416	0
	2.78	3.08	0
	3.86	4.08	0
0.2 -N	5.161	5.715	0
	1.268	1.509	0
	1.914	2.274	0
	2.464	3.040	0
0.4 -N	3.967	4.898	0
	1.202	1.614	0
	1.894	2.263	0

(McCrae, Z. anorg. 1903, 35. 14.)

Sulphur dioxide ammonia, SO₂, NH₃.

Very hygroscopic. Easily sol. in H₂O decomp. (Schumann, Z. anorg. 1900 49.)

SO₂, 2NH₃. Somewhat hygroscopic Sol. in H₂O with evolution of NH₃. (Schumann, Z. anorg. 1900, 23. 50.)

5SO₂, 4NH₃. Very deliquescent. Very sol. in H₂O. (Divers and O. Chem. Soc. 1901, 79. 1103.)

Sulphur trioxide, SO₃.

Fumes on air. Miscible with H₂O, evolution of much heat. Sol. in H₂SO₄ comp. by alcohol and ether.

is in two modifications, one of which is and miscible with H_2SO_4 , while the other is only slowly sol. therein. Miscible with CS_2 at 30° , but at 15° CS_2 is only $\frac{1}{2}$ pt. SO_2 , and SO_2 , $\frac{1}{2}$ pt. (Schults-Sellack, Pogg. 139. 480.) There is only one modification, the liquid, absorbs H_2O and becomes solid. (Rebs, 356.)

Miscible with liquid SO_2 . (Schults-)

Iso Sulphuric acid.

Heptoxide, S_7O_{17} .

Exposes on air. Slowly decomp. at 0° , increasingly on warming. Sol. in conc. (Berthelot, J. pr. (2) 17. 48.)

is compound S_2O_7 , $2\text{H}_2\text{O}$.

Formula is SO_4 , according to Traube (B. 4), and S_2O_7 is $\text{SO}_3 + \text{SO}_4$.

Iso Marshall (Chem. Soc. 59. 771).

He (B. 26. 148) denies the existence of

oxybromide, SOBr_2 .

Thionyl bromide.

oxychloride, SOCl_2 .

Thionyl chloride.

Cl_2 . See Sulphuryl chloride.

Cl_2 . See Pyrosulphuryl chloride.

Cl . See Sulphuryl hydroxyl chloride.

Cl . Decomp. by H_2O and alcohol. (C. R. 94. 446.)

ure of about $178\text{Cl}_2 + 28\text{SOCl}_2$ and (Knoll, B. 1898, 31. 2183.)

oxytetrachloride, $\text{S}_2\text{O}_4\text{Cl}_4$.

ntly decomp. by H_2O , dil. acids, or (Millon, A. ch. (3) 29. 327.)

in warm S_2Cl_2 . (Carius, A. 106. 295.) mp. violently with CS_2 .

oxyfluoride, SO_2F_2 .

Sulphuryl fluoride.

. See Thionyl fluoride.

diphosphide, P_2S .

Phosphorus monosulphide.

tetraphosphide, P_4S .

Phosphorus semisulphide.

retted hydrogen, H_2S

hydrogen sulphide.

ric acid, H_2SO_4 .

ble with H_2O in all proportions.

Sp. gr. of $\text{H}_2\text{SO}_4 + \text{Aq.}$

Baume degrees	Sp. gr.	% H_2SO_4	Baume degrees	Sp. gr.	% H_2SO_4
66	1.842	100	66	1.844	100
60	1.725	84.22	60	1.717	82.34
55	1.618	74.32	55	1.618	74.32
50	1.524	66.45	54	1.603	72.70
45	1.466	58.02	53	1.588	71.17
40	1.375	50.41	52	1.566	69.30
35	1.315	43.21	51	1.550	68.03
30	1.260	36.52	50	1.532	66.45
25	1.210	30.12	49	1.515	64.37
20	1.162	24.01	48	1.500	62.80
15	1.114	17.89	47	1.482	61.32
10	1.076	11.73	46	1.466	59.85
5	1.023	5.60	45	1.454	58.02

(Vauquelin, A. ch. 76. 260.)

(Darcet, A. ch. (2) 1. 198.)

Sp. gr. of $\text{H}_2\text{SO}_4 + \text{Aq.}$

% H_2SO_4	Sp. gr. at 15°	Sp. gr. at 25°	% H_2SO_4	Sp. gr. at 15°	Sp. gr. at 25°
0	0.9986	0.9955	50	1.3866	1.3780
2.5		1.0115	55	1.4347	
5	1.0284	1.0272	60	1.4860	1.4767
10	1.0659	1.0604	65	1.5402	
15	1.0998		70	1.5946	1.5863
20	1.1378	1.3311	75	1.6534	
25	1.1787		80	1.7092	1.6996
30	1.2154	1.2078	85	1.7602	
35	1.2502		90	1.8050	1.7940
40	1.2876	1.2858	95	1.8318	
45	1.3409		100	1.8406	1.8286

(Delezenne, 1888.)

Sp. gr. at 15.56° , and b.-pt. of $\text{H}_2\text{SO}_4 + \text{Aq.}$

Sp. gr.	% SO_2	B.-pt.	Sp. gr.	% SO_2	B.-pt.
1.850	81	326°	1.769	67	217°
1.849	80	318	1.757	66	210
1.848	79	310	1.744	65	205
1.847	78	301	1.730	64	200
1.845	77	293	1.715	63	195
1.842	76	285	1.699	62	190
1.838	75	277	1.684	61	186
1.833	74	268	1.670	60	182
1.827	73	260	1.650	58.6	177
1.819	72	253	1.620	50	143
1.810	71	245	1.408	40	127
1.801	70	238	1.300	30	116
1.791	69	230	1.200	20	107
1.780	68	224	1.100	10	103

(Dalton, N. Syst. 2. 210.)

Sp. gr. of $\text{H}_2\text{SO}_4 + \text{Aq}$ at 15° .

Sp. gr.	% SO_2	% H_2SO_4	Sp. gr.	% SO_2	% H_2SO_4
1.8485	81.54	100	1.5975	57.08	70
1.8460	79.90	98	1.5760	55.45	68
1.8410	78.28	96	1.5503	53.82	66
1.8336	76.65	94	1.5280	52.18	64
1.8233	75.02	92	1.5060	50.55	62
1.8115	73.39	90	1.4860	48.92	60
1.7962	71.75	88	1.4660	47.29	58
1.7774	70.12	86	1.4460	45.66	56
1.7570	68.49	84	1.4265	44.03	54
1.7360	66.86	82	1.4073	42.40	52
1.7120	65.23	80	1.3884	40.77	50
1.6870	63.60	78	1.3697	39.14	48
1.6630	61.97	76	1.3530	37.51	46
1.6415	60.34	74	1.3345	35.88	44
1.6204	58.71	72	1.3165	34.25	42

Sp. gr. of H_2SO_4 + Aq at 15° —Continued.

Sp. gr.	% SO_3	% H_2SO_4	Sp. gr.	% SO_3	% H_2SO_4
1.2999	32.61	40	1.1410	16.31	20
1.2925	30.94	38	1.1249	14.08	18
1.2854	29.35	36	1.1090	13.05	16
1.2499	27.72	34	1.0953	11.41	14
1.2334	25.09	32	1.0809	9.78	12
1.2184	24.46	30	1.0682	8.15	10
1.2032	22.83	28	1.0544	6.52	8
1.1876	21.20	26	1.0405	4.89	6
1.1706	19.57	24	1.0268	3.26	4
1.1549	17.94	22	1.0140	1.63	2

(U're. Schw. J. 35. 444.)

Sp. gr. of H_2SO_4 + Aq

Degrees Baume	Sp. gr.	At 0°		At 15°	
		% SO_3	% H_2SO_4	% SO_3	% H_2SO_4
5	1.038	5.1	4.2	5.4	4.5
10	1.075	10.3	8.4	10.9	8.9
15	1.116	15.5	12.7	16.3	13.3
20	1.161	21.2	17.3	22.4	18.3
25	1.209	27.2	22.2	28.3	23.1
30	1.242	33.6	27.4	34.8	28.4
33	1.290	37.6	30.7	38.9	31.8
35	1.320	40.4	33.0	41.6	34.0
36	1.332	41.7	34.1	43.0	35.1
37	1.345	43.1	35.2	44.3	36.2
38	1.357	44.5	36.3	45.5	37.2
39	1.370	45.9	37.5	46.9	38.3
40	1.383	47.3	38.6	48.4	39.5
41	1.397	48.7	39.7	49.9	40.7
42	1.410	50.0	40.8	51.2	41.8
43	1.424	51.4	41.9	52.5	42.9
44	1.438	52.8	43.1	54.0	44.1
45	1.453	54.3	44.3	55.4	45.2
46	1.468	55.7	45.5	56.9	46.4
47	1.484	57.1	46.6	58.2	47.5
48	1.498	58.5	47.8	59.6	48.7
49	1.514	60.0	49.0	61.1	50.0
50	1.530	61.4	50.1	62.6	51.1
51	1.546	62.9	51.3	63.9	52.2
52	1.563	64.4	52.6	65.4	53.4
53	1.581	65.9	53.8	66.9	54.6
54	1.597	67.4	55.0	68.4	55.8
55	1.615	68.9	56.2	70.0	57.1
56	1.634	70.5	57.5	71.6	58.4
57	1.652	72.1	58.8	73.2	59.7
58	1.671	73.6	60.1	74.7	61.0
59	1.691	75.2	61.4	76.3	62.3
60	1.711	76.9	62.8	78.0	63.6
61	1.732	78.5	64.2	79.8	65.1
62	1.753	81.4	65.7	81.7	66.7
63	1.774	82.4	67.2	83.9	68.5
64	1.796	83.6	69.0	85.3	70.4
65	1.819	87.4	71.3	89.5	73.0
66.5	1.840	89.1	72.2	91.8	74.9
66.8	1.847	90.4	73.8	94.5	77
66	1.842	91.3	74.5	100.0	81.6
66.2	1.846	92.5	75.5		
66.4	1.852	93.0	77.5		
66.6	1.857	100.0	81.6		

(Bineau. A. ch. (3) 34. 121.)

The sp. gr. found at 1° can be reduced to sp. gr. at 0° by multiplying by $\frac{144.38}{144.38-t}$, or by using the following table (Bineau.)

Correction of sp. gr. for temperature, to be added for lowering of the temp. of 10°, or subtracted for corresponding increase.

Sp. gr. of acid at 0°	Corr.	Sp. gr. of acid at 0°	Corr.	Sp. gr. of acid at 0°	Corr.
1.04	0.002	1.15	0.005	1.45	0.008
1.07	0.003	1.20	0.006	1.70	0.008
1.10	0.004	1.30	0.007	1.95	0.008

(Bineau.)

Sp. gr. of H_2SO_4 + Aq at 15°. a = % SO_3 ; b = sp. gr. if % is SO_3 ; c = sp. gr. if % is H_2SO_4 .

a	b	c	a	b	c
1	1.009	1.0064	51	1.530	1.405
2	1.017	1.013	52	1.545	1.415
3	1.025	1.019	53	1.556	1.425
4	1.034	1.0256	54	1.573	1.435
5	1.041	1.032	55	1.585	1.445
6	1.049	1.039	56	1.600	1.456
7	1.058	1.0464	57	1.615	1.466
8	1.067	1.0536	58	1.627	1.476
9	1.076	1.061	59	1.642	1.490
10	1.085	1.068	60	1.656	1.510
11	1.095	1.0756	61	1.675	1.512
12	1.104	1.083	62		1.523
13	1.114	1.091		1.701	1.534
14	1.123	1.098	64	1.716	1.545
15	1.133	1.106	65	1.730	1.557
16	1.142	1.1136	66	1.742	1.575
17	1.150	1.121	67	1.765	1.590
18	1.160	1.129	68	1.770	1.592
19	1.170	1.136	69	1.781	1.604
20	1.180	1.144	70	1.792	1.615
21	1.190	1.1516	71	1.802	1.627
22	1.200	1.159	72	1.810	1.639
23	1.210	1.167	73	1.819	1.651
24	1.220	1.174	74	1.825	1.663
25	1.229	1.182	75	1.830	1.675
26	1.239	1.190	76	1.834	1.686
27	1.248	1.198	77	1.837	1.696
28	1.258	1.2066	78	1.839	1.710
29	1.268	1.215	79	1.841	1.722
30	1.278	1.223	80	1.842	1.734
31	1.288	1.231	81		1.745
32	1.300	1.239	82		1.756
33	1.310	1.2476	83		1.767
34	1.320	1.256	84		1.777
35	1.332	1.264	85		1.786
36	1.344	1.272	86		1.794
37	1.354	1.281	87		1.802
38	1.367	1.289	88		1.809
39	1.378	1.2976	89		1.816
40	1.390	1.306	90		1.822
41	1.401	1.315	91		1.827
42	1.415	1.324	92		1.831
43	1.427	1.333	93		1.834
44	1.440	1.342	94		1.836
45	1.451	1.351	95		1.8376
46	1.465	1.361	96		1.8384
47	1.478	1.370	97		1.840
48	1.490	1.379	98		1.8406
49	1.501	1.3886	99		1.842
50	1.517	1.398	100		1.8426

(Bineau, calculated by Gerlach, Z. anal. 3. 202.)

Sp. gr. of H₂SO₄+Aq at 15°; H₂O at 0°=1.

Sp. gr. of H₂SO₄, etc.—Continued.

$\% \text{H}_2\text{SO}_4$	Sp. gr.	$\% \text{H}_2\text{SO}_4$	Sp. gr.	$\% \text{H}_2\text{SO}_4$	Sp. gr.	Sp. gr.	$\% \text{SO}_3$	$\% \text{H}_2\text{SO}_4$	Sp. gr.	$\% \text{SO}_3$	$\% \text{H}_2\text{SO}_4$
1	1.006	35	1.264	68	1.592	1.150	17.07	20.91	1.455	45.31	55.50
2	1.012	36	1.272	69	1.604	1.155	17.59	21.55	1.460	45.69	55.97
3	1.018	37	1.281	70	1.615	1.160	18.11	22.19	1.465	46.07	56.43
4	1.025	38	1.290	71	1.626	1.165	18.64	22.83	1.470	46.45	56.90
5	1.032	39	1.298	72	1.638	1.170	19.06	23.47	1.475	46.83	57.37
6	1.039	40	1.307	73	1.650	1.175	19.69	24.12	1.480	47.21	57.83
7	1.046	41	1.316	74	1.662	1.180	20.21	24.76	1.485	47.57	58.28
8	1.053	42	1.324	75	1.674	1.185	20.73	25.40	1.490	47.95	58.74
9	1.061	43	1.333	76	1.684	1.190	21.26	26.04	1.495	48.34	59.22
10	1.069	44	1.342	77	1.697	1.195	21.78	26.68	1.500	48.73	59.70
11	1.076	45	1.352	78	1.710	1.200	22.30	27.32	1.505	49.12	60.18
12	1.084	46	1.361	79	1.721	1.205	22.82	27.95	1.510	49.51	60.65
13	1.091	47	1.370	80	1.732	1.210	23.33	28.58	1.515	49.89	61.12
14	1.099	48	1.379	81	1.743	1.215	23.84	29.21	1.520	50.28	61.59
15	1.106	49	1.389	82	1.753	1.220	24.36	29.84	1.525	50.66	62.06
16	1.114	50	1.399	83	1.763	1.225	24.88	30.48	1.530	51.04	62.53
17	1.122	51	1.409	84	1.773	1.230	25.39	31.11	1.535	51.43	63.00
18	1.129	52	1.418	85	1.783	1.235	25.88	31.70	1.540	51.78	63.43
19	1.137	53	1.428	86	1.792	1.240	26.35	32.28	1.545	52.12	63.85
20	1.145	54	1.438	87	1.800	1.245	26.83	32.86	1.550	52.46	64.26
21	1.153	55	1.448	88	1.807	1.250	27.29	33.40	1.555	52.79	64.67
22	1.161	56	1.459	89	1.814	1.255	27.76	34.00	1.560	53.12	65.08
23	1.168	57	1.469	90	1.820	1.260	28.22	34.57	1.565	53.46	65.49
24	1.176	58	1.480	91	1.825	1.265	28.69	35.14	1.570	53.80	65.90
25	1.184	59	1.491	92	1.8294	1.270	29.15	35.71	1.575	54.13	66.30
26	1.191	60	1.501	93	1.8339	1.275	29.62	36.29	1.580	54.46	66.71
27	1.199	61	1.512	94	1.8372	1.280	30.10	36.87	1.585	54.80	67.13
28	1.207	62	1.523	95	1.8390	1.285	30.57	31.45	1.590	55.18	67.59
29	1.215	63	1.535	96	1.8406	1.290	31.04	38.03	1.595	55.55	68.05
30	1.223	64	1.546	97	1.8410	1.295	31.52	38.61	1.600	55.93	68.51
31	1.231	65	1.558	98	1.8412	1.300	31.99	39.19	1.605	56.30	68.97
32	1.239	66	1.569	99	1.8403	1.305	32.46	39.77	1.610	56.68	69.43
33	1.247	67	1.580	100	1.8384	1.310	32.94	40.35	1.615	57.05	69.89
34	1.256	1.315	33.41	40.93	1.620	57.40	70.32

(From 1–91 % according to Kolb, calculated by Gerlach; from 92–100% according to Lunge and Naef, calculated by Gerlach, Z. anal. 27. 316.)

Sp. gr. of H₂SO₄ at 15° compared with H₂O at 4° and 0 mm. pressure.

Sp. gr.	$\% \text{SO}_3$	$\% \text{H}_2\text{SO}_4$	Sp. gr.	$\% \text{SO}_3$	$\% \text{H}_2\text{SO}_4$	Sp. gr.	$\% \text{SO}_3$	$\% \text{H}_2\text{SO}_4$	Sp. gr.	$\% \text{SO}_3$	$\% \text{H}_2\text{SO}_4$
1.000	0.07	0.09	1.075	8.90	10.90	1.370	38.32	46.94	1.675	61.20	74.97
1.005	0.68	0.83	1.080	9.47	11.60	1.375	38.75	47.47	1.680	61.57	75.42
1.010	1.28	1.57	1.085	10.04	12.30	1.380	39.18	48.00	1.685	61.93	75.86
1.015	1.88	2.30	1.090	10.60	12.99	1.385	39.62	48.53	1.690	62.29	76.30
1.020	2.47	3.03	1.095	11.16	13.67	1.390	40.05	49.06	1.695	62.64	76.73
1.025	3.07	3.76	1.100	11.71	14.35	1.395	40.48	49.59	1.700	63.00	77.17
1.030	3.67	4.49	1.105	12.27	15.07	1.400	40.91	50.11	1.705	63.35	77.60
1.035	4.27	5.23	1.110	12.82	15.71	1.405	41.33	50.63	1.710	63.70	78.04
1.040	4.87	5.96	1.115	13.36	16.36	1.410	41.76	51.15	1.715	64.07	78.48
1.045	5.45	6.67	1.120	13.89	17.01	1.415	42.17	51.66	1.720	64.43	78.92
1.050	6.02	7.37	1.125	14.42	17.66	1.420	42.57	52.15	1.725	64.78	79.36
1.055	6.59	8.07	1.130	14.95	18.31	1.425	42.96	52.63	1.730	65.14	79.80
1.060	7.16	8.77	1.135	15.48	18.96	1.430	43.36	53.11	1.735	65.50	80.24
1.065	7.73	9.47	1.140	16.01	19.61	1.435	43.75	53.59	1.740	65.86	80.68
1.070	8.32	10.19	1.145	16.54	20.26	1.440	44.14	54.07	1.745	66.22	81.12
						1.445	44.53	54.55	1.750	66.58	81.56
						1.450	44.92	55.03	1.755	66.94	82.00

Sp. gr. of H_2SO_4 , etc.—Continued.

Sp. gr.	% SO_3	% H_2SO_4	Sp. gr.	% SO_3	% H_2SO_4
1.760	67.30	82.44	1.829	75.03	91.90
1.765	67.65	82.88	1.830	75.19	92.10
1.770	68.02	83.32	1.831	75.35	92.30
1.775	68.49	83.90	1.832	75.53	92.52
1.780	68.98	84.50	1.833	75.72	92.75
1.785	69.74	85.10	1.834	75.96	93.05
1.790	69.96	85.70	1.835	76.27	93.43
1.795	70.45	86.30	1.836	76.57	93.80
1.800	70.94	86.90	1.837	76.90	94.20
1.805	71.50	87.60	1.838	77.23	94.60
1.810	72.08	88.30	1.839	77.55	95.00
1.815	72.69	89.05	1.840	78.04	95.60
1.820	73.51	90.05	1.8405	78.33	95.95
1.821	73.63	90.20	1.8415	79.19	97.00
1.822	73.80	90.40	1.8410	79.76	97.70
1.823	73.96	90.60	1.8415	80.16	98.20
1.824	74.12	90.80	1.8400	80.57	98.70
1.825	74.29	91.00	1.8400	80.98	99.20
1.826	74.49	91.25	1.8395	81.18	99.45
1.827	74.69	91.50	1.8390	81.39	99.70
1.828	74.86	91.70	1.8385	81.59	99.95

(Lunge and Isler, Zeit. angew. Ch. 9. 129.)

Sp. gr. of conc H_2SO_4 + Aq at 15°.

% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.
100	1.8384	99.02	1.8417
99.98	1.8385	98.98	1.8418
99.96	1.8386	98.94	1.8419
99.94	1.8387	98.84	1.8420
99.92	1.8388	98.84	1.8421
99.90	1.8389	98.78	1.8422
99.88	1.8390	98.71	1.8423
99.86	1.8391	98.63	1.8424
99.84	1.8392	98.56	1.8425
99.81	1.8393	98.48	1.8426
99.78	1.8394	98.40	1.8427
99.76	1.8395	98.32	1.8428
99.73	1.8396	98.22	1.8429
99.70	1.8397	98.08	1.8430
99.67	1.8398	97.85	1.8431
99.64	1.8399	97.50	1.8432
99.61	1.8400	97.10	1.8431
99.58	1.8401	96.93	1.8430
99.55	1.8402	96.76	1.8429
99.52	1.8403	96.65	1.8428
99.49	1.8404	96.55	1.8427
99.46	1.8405	96.46	1.8426
99.43	1.8406	96.39	1.8425
99.40	1.8407	96.31	1.8424
99.37	1.8408	96.24	1.8423
99.33	1.8409	96.16	1.8422
99.29	1.8410	96.09	1.8421
99.25	1.8411	96.02	1.8420
99.22	1.8412	95.95	1.8419
99.19	1.8413	95.88	1.8418
99.16	1.8414	95.81	1.8417
99.11	1.8415	95.74	1.8416
99.06	1.8416	95.67	1.8415

Sp. gr. of conc H_2SO_4 , etc.—Continued.

% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.
95.61	1.8414	93.32	1.8352
95.55	1.8413	93.29	1.8351
95.50	1.8412	93.26	1.8350
95.45	1.8411	93.23	1.8349
95.40	1.8410	93.20	1.8348
95.35	1.8409	93.17	1.8347
95.30	1.8408	93.14	1.8346
95.25	1.8407	93.12	1.8345
95.21	1.8406	93.09	1.8344
95.16	1.8405	93.06	1.8343
95.12	1.8404	93.00	1.8342
95.08	1.8403	92.98	1.8341
95.04	1.8402	92.95	1.8340
95.00	1.8401	92.93	1.8339
94.96	1.8400	92.90	1.8337
94.92	1.8399	92.87	1.8336
94.88	1.8398	92.84	1.8335
94.84	1.8397	92.82	1.8334
94.81	1.8396	92.79	1.8333
94.77	1.8395	92.77	1.8332
94.73	1.8394	92.73	1.8331
94.69	1.8393	92.71	1.8330
94.65	1.8392	92.69	1.8329
94.61	1.8391	92.66	1.8328
94.57	1.8390	92.63	1.8327
94.53	1.8389	92.61	1.8326
94.49	1.8388	92.59	1.8325
94.46	1.8387	92.56	1.8324
94.42	1.8386	92.54	1.8323
94.38	1.8385	92.52	1.8322
94.34	1.8384	92.49	1.8321
94.31	1.8383	92.46	1.8320
94.27	1.8382	92.44	1.8319
94.24	1.8381	92.41	1.8318
94.20	1.8380	92.39	1.8317
94.17	1.8379	92.37	1.8316
94.13	1.8378	92.34	1.8315
94.10	1.8377	92.32	1.8314
94.07	1.8376	92.29	1.8313
94.03	1.8375	92.27	1.8312
94.00	1.8374	92.24	1.8311
93.97	1.8373	92.22	1.8310
93.93	1.8372	92.19	1.8309
93.90	1.8371	92.17	1.8308
93.87	1.8370	92.15	1.8307
93.83	1.8369	92.12	1.8306
93.80	1.8368	92.10	1.8305
93.77	1.8367	92.07	1.8304
93.74	1.8366	92.05	1.8303
93.71	1.8365	92.02	1.8302
93.68	1.8364	92.00	1.8301
93.65	1.8363	91.98	1.8300
93.62	1.8362	91.95	1.8299
93.59	1.8361	91.93	1.8298
93.56	1.8360	91.91	1.8297
93.53	1.8359	91.88	1.8296
93.51	1.8358	91.86	1.8295
93.47	1.8357	91.84	1.8294
93.44	1.8356	91.81	1.8293
93.41	1.8355	91.78	1.8292
93.38	1.8354	91.76	1.8291
93.35	1.8353	91.74	1.8290

Sp. gr. of conc. H_2SO_4 , etc.—Continued.

% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.
91.72	1.8298	90.78	1.8244
91.70	1.8288	90.76	1.8243
91.68	1.8287	90.74	1.8242
91.65	1.8286	90.72	1.8241
91.63	1.8285	90.70	1.8240
91.61	1.8284	90.68	1.8239
91.59	1.8283	90.66	1.8238
91.56	1.8282	90.64	1.8237
91.54	1.8281	90.62	1.8236
91.52	1.8280	90.60	1.8235
91.50	1.8279	90.59	1.8234
91.47	1.8278	90.57	1.8233
91.45	1.8277	90.55	1.8232
91.43	1.8276	90.53	1.8231
91.41	1.8275	90.51	1.8230
91.39	1.8274	90.49	1.8229
91.37	1.8273	90.47	1.8228
91.35	1.8272	90.46	1.8227
91.32	1.8271	90.44	1.8226
91.30	1.8270	90.42	1.8225
91.28	1.8269	90.40	1.8224
91.26	1.8268	90.38	1.8223
91.24	1.8267	90.37	1.8222
91.22	1.8266	90.35	1.8221
91.20	1.8265	90.33	1.8220
91.18	1.8264	90.31	1.8219
91.16	1.8263	90.29	1.8218
91.14	1.8262	90.28	1.8217
91.12	1.8261	90.26	1.8216
91.10	1.8260	90.24	1.8215
91.08	1.8259	90.23	1.8214
91.06	1.8258	90.20	1.8213
91.04	1.8257	90.18	1.8212
91.02	1.8256	90.17	1.8211
91.00	1.8255	90.15	1.8210
90.98	1.8254	90.13	1.8209
90.96	1.8253	90.11	1.8208
90.94	1.8252	90.10	1.8207
90.92	1.8251	90.08	1.8206
90.90	1.8250	90.06	1.8205
90.88	1.8249	90.04	1.8204
90.86	1.8248	90.02	1.8203
90.84	1.8247	90.01	1.8202
90.82	1.8246	89.99	1.8201
90.80	1.8245	89.97	1.8200

(Richmond [calculated from Pickering, Chem. Soc. 57. 64], Jour Soc. Ch. Ind. 9. 479.)

Sp. gr. of conc. H_2SO_4 + Aq at 15°.

% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.
90	1.8185	96	1.8406
*90.20	1.8195	97	1.8410
91	1.8241	*97.70	1.8413
*91.48	1.8271	98	1.8412
92	1.8294	*98.39	1.8406
*92.83	1.8334	*98.66	1.8409
93	1.8339	99	1.8403
94	1.8372	*99.47	1.8395
*94.84	1.8387	100	1.8384
95	1.8390	*100.35	1.8411
*95.97	1.8406		

*Determined by experiment.

(Lunge and Naef, Dingl 248. 91.)

Sp. gr. of H_2SO_4 + Aq at room temp. containing:

7.875 15.503 23.429% H_2SO_4
1.0651 1.1305 1.2003

(Wagner, W. Ann. 1883, 18. 265).

Sp. gr. of H_2SO_4 + Aq at 25°.

Concentration of H_2SO_4 + Aq	Sp. gr.
1—normal	1.0303
$\frac{1}{2}$ —"	1.0154
$\frac{1}{3}$ —"	1.0074
$\frac{1}{4}$ —"	1.0035

(Wagner, Z. phys. Ch. 1890, 5. 40.)

Sp. gr. of dil. H_2SO_4 + Aq

G-equivalents H_2SO_4 per liter	t°	Sp. gr. t°/t°
0.005049	17.343	1.0002082
0.01009	17.360	1.0004020
0.01512	17.382	1.0005879
0.02014	17.398	1.000769
0.03014	17.419	1.001125
0.002526	18.039	1.0001065
0.005050	18.040	1.0002084
0.01006	18.040	1.0004009
0.02005	18.040	1.0007068
0.03001	18.039	1.0011208
0.04980	18.040	1.0018096
0.00864	18.048	1.003460
0.146560	18.070	1.005045
0.19354	18.060	1.006580
0.28042	18.052	1.009686
0.47466	18.055	1.015616
0.4980	17.73	1.01634
4.980	17.95	1.15234
0.005176	12.997	1.0002106
0.01035	13.020	1.000411
0.01551	13.005	1.000603
0.12648	13.031	1.004438
0.25151	13.011	1.008565
0.37672	13.007	1.012639
0.50503	12.998	1.016758

(Kohlrausch, W. Ann. 1894, 53. 22.)

Sp. gr. of H_2SO_4 + Aq.			
% H_2SO_4	79.68	60.98	35.77
Sp. gr. 20°/20°	1.7383	1.5181	1.2719
% H_2SO_4	10.10	4.78	
Sp. gr. 20°/20°	1.0685	1.0317	
(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 268.)			

Sp. gr. of $N-H_2SO_4$ + Aq at 18°/4° = 1.0306.
(Loomis, W. Ann. 1896, 60. 550.)

Sp. gr. of H_2SO_4 + Aq at 19.4°, when p = percent strength of solution; d = observed density; w = volume conc. in grams per cc. ($\frac{pd}{100} = w$).

p	d	w
94 10	1.8380	1.7295
84 59	1.7998	1.5223
73 08	1.6743	1.2235
61 35	1.5341	0.9412
40 72	1.3220	0.5383
31 94	1.2430	0.3970
23 77	1.1747	0.2792
14 72	1.1023	0.1623
9 802	1.0670	0.1046
4 826	1.0320	0.0498

(Barnes, J. Phys. Chem. 1898, 2. 546.)

Sp. gr. of H_2SO_4 + Aq at 20°.

Normality of H_2SO_4 + Aq	% H_2SO_4	Sp. gr.
11 53	70 07	1.6129
9 01	59 26	1.4901
6 95	49 10	1.3672
4 77	36 68	1.2756
3 008	25 00	1.1791
1 002	9 25	1.0612

(Forebheimer, Z. phys. Ch. 1900, 24. 27.)

Sp. gr. of conc. and fuming H_2SO_4 at 15° and 45°

% H_2SO_4	Total SO_3 %	Free SO_3 %	Sp. gr. at 15°	Sp. gr. at 45°
95 98	78 35		1.8418	
96 68	78 92		1.8429	
96 99	79 18		1.8431	
97 66	79 72		1.8434 max.	
98 65	80 53		1.8403	
99 40	81 14		1.8388 min.	
99 76	81 44		1.8418	
100 00	81 63	0 01	1.8500	1.822
	83 46	10 01	1.888	1.858
	85 30	20 01	1.920	1.887
	87 14	30 01	1.957	1.920
	88 97	40 01	1.979	1.945
	90 81	50 02	1.999	1.964 max
	92 65	60 02	2.020 max.	1.959
	94 48	70 02	2.018	1.942
	96 32	80 02	2.008	1.890
	98 16	90 01	1.990	1.864
	100 00	100 01	1.984	1.814

(Knietzsch, B. 1901, 24. 4102.)

Sp. gr. of fuming H_2SO_4 at 35°.

Total SO_3 %	Free SO_3 %	Sp. gr.	Total SO_3 %	Free SO_3 %	Sp. gr.
81.63	0	1.8186	91.18	52	1.929
81.99	2	1.8270	91.55	54	1.939
82.36	4	1.8360	91.91	56	1.947
82.73	6	1.8425	92.28	58	1.954
83.09	8	1.8498	92.65	60	1.958
83.46	10	1.8566	93.02	62	1.969
83.82	12	1.8627	93.38	64	1.967
84.20	14	1.8692	93.75	66	1.968
84.56	16	1.8756	94.11	68	1.968
84.92	18	1.8830	94.48	70	1.964
85.30	20	1.8919	94.85	72	1.962
85.66	22	1.9020	95.21	74	1.962
86.03	24	1.9092	95.58	76	1.959
86.40	26	1.9158	95.95	78	1.955
86.76	28	1.9220	96.32	80	1.951
87.14	30	1.9287	96.69	82	1.948
87.50	32	1.9333	97.05	84	1.945
87.87	34	1.9405	97.42	86	1.946
88.24	36	1.9474	97.78	88	1.949
88.60	38	1.9534	98.16	90	1.956
88.97	40	1.9584	98.53	92	1.960
89.33	42	1.9612	98.90	94	1.972
89.70	44	1.9643	99.26	96	1.986
90.07	46	1.9672	99.63	98	1.996
90.44	48	1.9702	100.00	100	1.970
90.81	50	1.9733	...		

(Knietzsch, B. 1901, 24. 4101.)

Sp. gr. of H_2SO_4 + Aq at 15°/15° in air

Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4
1.000	0.00	1.025	4.12	1.056	8.19
1.001	0.15	1.026	4.26	1.057	8.33
1.002	0.31	1.030	4.41	1.058	8.47
1.003	0.46	1.031	4.56	1.059	8.62
1.004	0.60	1.032	4.70	1.060	8.76
1.005	0.73	1.033	4.85	1.061	8.90
1.006	0.87	1.034	5.00	1.062	9.04
1.007	1.01	1.035	5.14	1.063	9.18
1.008	1.15	1.036	5.29	1.064	9.33
1.009	1.30	1.037	5.44	1.065	9.47
1.010	1.45	1.038	5.58	1.066	9.61
1.011	1.60	1.039	5.73	1.067	9.75
1.012	1.75	1.040	5.88	1.068	9.89
1.013	1.89	1.041	6.03	1.069	10.04
1.014	2.04	1.042	6.17	1.070	10.18
1.015	2.19	1.043	6.32	1.071	10.31
1.016	2.34	1.044	6.46	1.072	10.45
1.017	2.49	1.045	6.60	1.073	10.59
1.018	2.64	1.046	6.75	1.074	10.73
1.019	2.79	1.047	6.89	1.075	10.87
1.020	2.93	1.048	7.04	1.076	11.00
1.021	3.08	1.049	7.18	1.077	11.14
1.022	3.23	1.050	7.32	1.078	11.28
1.023	3.38	1.051	7.47	1.079	11.42
1.024	3.53	1.052	7.61	1.080	11.56
1.025	3.67	1.053	7.76	1.081	11.69
1.026	3.82	1.054	7.90	1.082	11.83
1.027	3.97	1.055	8.04	1.083	11.97

gr. of H_2SO_4 + Aq at 15°/15° in air —
Continued.Sp. gr. of H_2SO_4 + Aq at 15°/15° in air.—
Continued.

% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4
12.11	1.145	20.25	1.206	27.95	1.267	35.33	1.328	42.35	1.389	48.92		
12.24	1.146	20.38	1.207	28.08	1.268	35.45	1.329	42.46	1.390	49.02		
12.38	1.147	20.51	1.208	28.20	1.269	35.57	1.330	42.57	1.391	49.13		
12.52	1.148	20.64	1.209	28.32	1.270	35.68	1.331	42.68	1.392	49.23		
12.66	1.149	20.77	1.210	28.45	1.271	35.80	1.332	42.79	1.393	49.34		
12.79	1.150	20.90	1.211	28.57	1.272	35.92	1.333	42.90	1.394	49.44		
12.93	1.151	21.03	1.212	28.69	1.273	36.04	1.334	43.01	1.395	49.54		
13.07	1.152	21.16	1.213	28.82	1.274	36.15	1.335	43.12	1.396	49.65		
13.20	1.153	21.28	1.214	28.94	1.275	36.27	1.336	43.23	1.397	49.75		
13.34	1.154	21.41	1.215	29.06	1.276	36.39	1.337	43.35	1.398	49.86		
13.48	1.155	21.54	1.216	29.18	1.277	36.51	1.338	43.46	1.399	49.96		
13.61	1.156	21.67	1.217	29.31	1.278	36.62	1.339	43.57	1.400	50.06		
13.75	1.157	21.80	1.218	29.43	1.279	36.70	1.340	43.68	1.401	50.16		
13.89	1.158	21.93	1.219	29.55	1.280	36.86	1.341	43.79	1.402	50.26		
14.02	1.159	22.05	1.220	29.69	1.281	36.97	1.342	43.90	1.403	50.37		
14.16	1.160	22.18	1.221	29.80	1.282	37.09	1.343	44.01	1.404	50.47		
14.29	1.161	22.31	1.222	29.92	1.283	37.21	1.344	44.12	1.405	50.57		
14.43	1.162	22.44	1.223	30.04	1.284	37.32	1.345	44.23	1.406	50.67		
14.56	1.163	22.56	1.224	30.17	1.285	37.44	1.346	44.34	1.407	50.77		
14.70	1.164	22.69	1.225	30.29	1.286	37.56	1.347	44.45	1.408	50.88		
14.83	1.165	22.82	1.226	30.41	1.287	37.68	1.348	44.56	1.409	50.98		
14.97	1.166	22.94	1.227	30.53	1.288	37.79	1.349	44.67	1.410	51.08		
15.10	1.167	23.07	1.228	30.65	1.289	37.91	1.350	44.77	1.411	51.18		
15.24	1.168	23.20	1.229	30.78	1.290	38.03	1.351	44.88	1.412	51.28		
15.37	1.169	23.32	1.230	30.90	1.291	38.14	1.352	44.99	1.413	51.38		
15.51	1.170	23.45	1.231	31.02	1.292	38.26	1.353	45.10	1.414	51.48		
15.64	1.171	23.57	1.232	31.14	1.293	38.37	1.354	45.21	1.415	51.58		
15.78	1.172	23.71	1.233	31.26	1.294	38.49	1.355	45.32	1.416	51.68		
15.91	1.173	23.83	1.234	31.38	1.295	38.60	1.356	45.43	1.417	51.78		
16.05	1.174	23.96	1.235	31.50	1.296	38.72	1.357	45.53	1.418	51.89		
16.18	1.175	24.08	1.236	31.62	1.297	38.83	1.358	45.64	1.419	51.99		
16.31	1.176	24.21	1.237	31.75	1.298	38.95	1.359	45.75	1.420	52.09		
16.45	1.177	24.34	1.238	31.87	1.299	39.06	1.360	45.86	1.421	52.19		
16.58	1.178	24.46	1.239	31.99	1.300	39.18	1.361	45.97	1.422	52.29		
16.71	1.179	24.59	1.240	32.11	1.301	39.29	1.362	46.07	1.423	52.39		
16.84	1.180	24.71	1.241	32.23	1.302	39.41	1.363	46.18	1.424	52.49		
16.98	1.181	24.84	1.242	32.35	1.303	39.52	1.364	46.29	1.425	52.59		
17.11	1.182	24.97	1.243	32.47	1.304	39.64	1.365	46.39	1.426	52.69		
17.24	1.183	25.09	1.244	32.59	1.305	39.75	1.366	46.50	1.427	52.79		
17.37	1.184	25.22	1.245	32.71	1.306	39.86	1.367	46.61	1.428	52.89		
17.51	1.185	25.34	1.246	32.83	1.307	39.98	1.368	46.71	1.429	52.98		
17.64	1.186	25.47	1.247	32.95	1.308	40.09	1.369	46.82	1.430	53.08		
17.77	1.187	25.59	1.248	33.07	1.309	40.20	1.370	46.92	1.431	53.18		
17.90	1.188	25.72	1.249	33.19	1.310	40.32	1.371	47.03	1.432	53.28		
18.03	1.189	25.84	1.250	33.31	1.311	40.43	1.372	47.14	1.433	53.38		
18.16	1.190	25.97	1.251	33.43	1.312	40.54	1.373	47.24	1.434	53.48		
18.30	1.191	26.09	1.252	33.55	1.313	40.66	1.374	47.35	1.435	53.58		
18.43	1.192	26.22	1.253	33.67	1.314	40.77	1.375	47.45	1.436	53.68		
18.56	1.193	26.34	1.254	33.79	1.315	40.88	1.376	47.56	1.437	53.78		
18.69	1.194	26.47	1.255	33.91	1.316	40.99	1.377	47.67	1.438	53.88		
18.82	1.195	26.59	1.256	34.02	1.317	41.11	1.378	47.77	1.439	53.97		
18.95	1.196	26.71	1.257	34.14	1.318	41.22	1.379	47.88	1.440	54.07		
19.08	1.197	26.84	1.258	34.26	1.319	41.33	1.380	47.98	1.441	54.17		
19.22	1.198	26.96	1.259	34.38	1.320	41.45	1.381	48.09	1.442	54.27		
19.34	1.199	27.09	1.260	34.50	1.321	41.56	1.382	48.10	1.443	54.36		
19.47	1.200	27.21	1.261	34.62	1.322	41.67	1.383	48.30	1.444	54.46		
19.60	1.201	27.33	1.262	34.74	1.323	41.79	1.384	48.40	1.445	54.56		
19.73	1.202	27.46	1.263	34.86	1.324	41.90	1.385	48.50	1.446	54.65		
19.86	1.203	27.58	1.264	34.98	1.325	42.01	1.386	48.61	1.447	54.75		
19.99	1.204	27.71	1.265	35.09	1.326	42.12	1.387	48.71	1.448	54.85		
20.12	1.206	27.83	1.266	35.21	1.327	42.23	1.388	48.82	1.449	54.94		

Sp. gr. of H ₂ SO ₄ +Aq at 15°/15° in air.— Continued.						Sp. gr. of H ₂ SO ₄ +Aq at 15° 15° in air.— Continued.					
Sp. gr.	% H ₂ SO ₄	Sp. gr.	% H ₂ SO ₄	Sp. gr.	% H ₂ SO ₄	Sp. gr.	% H ₂ SO ₄	Sp. gr.	% H ₂ SO ₄	Sp. gr.	% H ₂ SO ₄
1.450	55.04	1.511	60.78	1.572	66.23	1.633	71.48	1.694	76.65	1.755	82.01
1.451	55.14	1.512	60.87	1.573	66.31	1.634	71.57	1.695	76.74	1.756	82.11
1.452	55.24	1.513	60.96	1.574	66.40	1.635	71.65	1.696	76.82	1.757	82.21
1.453	55.33	1.514	61.05	1.575	66.49	1.636	71.74	1.697	76.91	1.758	82.31
1.454	55.43	1.515	61.14	1.576	66.57	1.637	71.82	1.698	76.99	1.759	82.41
1.455	55.53	1.516	61.24	1.577	66.66	1.638	71.91	1.699	77.08	1.760	82.51
1.456	55.62	1.517	61.33	1.578	66.75	1.639	71.99	1.700	77.17	1.761	82.61
1.457	55.72	1.518	61.42	1.579	66.83	1.640	72.07	1.701	77.25	1.762	82.71
1.458	55.82	1.519	61.51	1.580	66.92	1.641	72.16	1.702	77.34	1.763	82.80
1.459	55.91	1.520	61.60	1.581	67.01	1.642	72.25	1.703	77.42	1.764	82.90
1.460	56.01	1.521	61.69	1.582	67.10	1.643	72.33	1.704	77.51	1.765	83.00
1.461	56.11	1.522	61.78	1.583	67.18	1.644	72.42	1.705	77.60	1.766	83.10
1.462	56.20	1.523	61.87	1.584	67.27	1.645	72.50	1.706	77.68	1.767	83.20
1.463	56.30	1.524	61.96	1.585	67.36	1.646	72.59	1.707	77.77	1.768	83.30
1.464	56.39	1.525	62.05	1.586	67.44	1.647	72.67	1.708	77.85	1.769	83.40
1.465	56.49	1.526	62.14	1.587	67.53	1.648	72.76	1.709	77.94	1.770	83.50
1.466	56.59	1.527	62.23	1.588	67.62	1.649	72.84	1.710	78.03	1.771	83.60
1.467	56.68	1.528	62.32	1.589	67.70	1.650	72.93	1.711	78.11	1.772	83.70
1.468	56.78	1.529	62.41	1.590	67.79	1.651	73.01	1.712	78.20	1.773	83.80
1.469	56.87	1.530	62.50	1.591	67.88	1.652	73.10	1.713	78.28	1.774	83.90
1.470	56.97	1.531	62.59	1.592	67.97	1.653	73.18	1.714	78.37	1.775	84.00
1.471	57.06	1.532	62.68	1.593	68.05	1.654	73.27	1.715	78.46	1.776	84.10
1.472	57.16	1.533	62.77	1.594	68.14	1.655	73.35	1.716	78.54	1.777	84.20
1.473	57.25	1.534	62.86	1.595	68.23	1.656	73.43	1.717	78.63	1.778	84.30
1.474	57.35	1.535	62.95	1.596	68.31	1.657	73.52	1.718	78.72	1.779	84.40
1.475	57.44	1.536	63.04	1.597	68.40	1.658	73.52	1.719	78.80	1.780	84.50
1.476	57.54	1.537	63.13	1.598	68.49	1.659	73.69	1.720	78.89	1.781	84.60
1.477	57.63	1.538	63.22	1.599	68.57	1.660	73.77	1.721	78.97	1.782	84.70
1.478	57.73	1.539	63.31	1.600	68.66	1.661	73.86	1.722	79.06	1.783	84.80
1.479	57.82	1.540	63.40	1.601	68.74	1.662	73.94	1.723	79.15	1.784	84.90
1.480	57.92	1.541	63.49	1.602	68.83	1.663	74.02	1.724	79.23	1.785	85.00
1.481	58.01	1.542	63.58	1.603	68.92	1.664	74.11	1.725	79.32	1.786	85.10
1.482	58.10	1.543	63.67	1.604	69.00	1.665	74.19	1.726	79.41	1.787	85.20
1.483	58.20	1.544	63.76	1.605	69.09	1.666	74.27	1.727	79.49	1.788	85.30
1.484	58.29	1.545	63.85	1.606	69.17	1.667	74.36	1.728	79.58	1.789	85.40
1.485	58.38	1.546	63.94	1.607	69.26	1.668	74.44	1.729	79.67	1.790	85.50
1.486	58.48	1.547	64.03	1.608	69.35	1.669	74.53	1.730	79.75	1.791	85.60
1.487	58.57	1.548	64.12	1.609	69.43	1.670	74.61	1.731	79.84	1.792	85.70
1.488	58.66	1.549	64.20	1.610	69.52	1.671	74.69	1.732	79.93	1.793	85.80
1.489	58.75	1.550	64.29	1.611	69.60	1.672	74.78	1.733	80.02	1.794	85.90
1.490	58.85	1.551	64.38	1.612	69.69	1.673	74.86	1.734	80.11	1.795	86.00
1.491	58.94	1.552	64.47	1.613	69.78	1.674	74.95	1.735	80.20	1.796	86.10
1.492	59.03	1.553	64.55	1.614	69.86	1.675	75.03	1.736	80.29	1.797	86.20
1.493	59.12	1.554	64.64	1.615	69.95	1.676	75.12	1.737	80.38	1.798	86.30
1.494	59.22	1.555	64.73	1.616	70.03	1.677	75.20	1.738	80.47	1.799	86.40
1.495	59.31	1.556	64.82	1.617	70.12	1.678	75.29	1.739	80.56	1.800	86.50
1.496	59.41	1.557	64.91	1.618	70.20	1.679	75.37	1.740	80.65	1.801	86.60
1.497	59.50	1.558	65.00	1.619	70.29	1.680	75.46	1.741	80.74	1.802	86.70
1.498	59.59	1.559	65.08	1.620	70.38	1.681	75.54	1.742	80.84	1.803	86.80
1.499	59.68	1.560	65.17	1.621	70.46	1.682	75.63	1.743	80.92	1.804	86.90
1.500	59.78	1.561	65.26	1.622	70.55	1.683	75.71	1.744	81.01	1.805	87.00
1.501	59.87	1.562	65.35	1.623	70.63	1.684	75.80	1.745	81.10	1.806	87.10
1.502	59.96	1.563	65.44	1.624	70.72	1.685	75.88	1.746	81.19	1.807	87.20
1.503	60.05	1.564	65.52	1.625	70.80	1.686	75.97	1.747	81.28	1.808	87.30
1.504	60.14	1.565	65.61	1.626	70.89	1.687	76.05	1.748	81.37	1.809	87.40
1.505	60.23	1.566	65.70	1.627	70.97	1.688	76.14	1.749	81.46	1.810	87.50
1.506	60.33	1.567	65.79	1.628	71.06	1.689	76.22	1.750	81.55	1.811	87.60
1.507	60.42	1.568	65.88	1.629	71.14	1.690	76.31	1.751	81.64	1.812	87.70
1.508	60.51	1.569	65.96	1.630	71.23	1.691	76.39	1.752	81.73	1.813	87.80
1.509	60.60	1.570	66.05	1.631	71.31	1.692	76.48	1.753	81.82	1.814	87.90
1.510	60.69	1.571	66.14	1.632	71.40	1.693	76.56	1.754	81.92	1.815	88.00

Sp. gr. of H₂SO₄+Aq at t°. Sp. gr. of H₂O at 15° = 1.

	0°	10°	15°	20°	25°	30°	40°	50°	60°
0	1.00074	1.00060	1.00000	0.99910	0.99794	0.99654	0.99311	0.98895	0.98418
1	1.00833	1.00773	1.00698	1.00594	1.00465	1.00312	0.99950	0.99522	0.99034
2	1.01563	1.01466	1.01381	1.01266	1.01126	1.00963	1.00585	1.00143	0.99644
3	1.02281	1.02153	1.02055	1.01928	1.01777	1.01607	1.01216	1.00761	1.00252
4	1.03001	1.02841	1.02728	1.02590	1.02428	1.02251	1.01848	1.01383	1.00865
5	1.03728	1.03533	1.03406	1.03258	1.03086	1.02902	1.02487	1.02013	1.01484
6	1.04461	1.04232	1.04092	1.03934	1.03756	1.03565	1.03138	1.02653	1.02114
7	1.05199	1.04939	1.04786	1.04618	1.04434	1.04235	1.03796	1.03302	1.02752
8	1.05942	1.05652	1.05486	1.05308	1.05116	1.04910	1.04458	1.03952	1.03393
9	1.06689	1.06370	1.06192	1.06002	1.05799	1.05585	1.05119	1.04605	1.04041
0	1.07439	1.07093	1.06903	1.06702	1.06490	1.06267	1.05787	1.05264	1.04696
1	1.08194	1.07821	1.07619	1.07408	1.07186	1.06955	1.06462	1.05930	1.05357
2	1.08954	1.08555	1.08342	1.08120	1.07890	1.07650	1.07145	1.06604	1.06027
3	1.09718	1.09294	1.09071	1.08839	1.08600	1.08352	1.07834	1.07284	1.06703
4	1.10488	1.10040	1.09805	1.09564	1.09316	1.09061	1.08530	1.07971	1.07385
5	1.11261	1.10790	1.10546	1.10295	1.10039	1.09776	1.09233	1.08666	1.08075
6	1.12040	1.11547	1.11292	1.11033	1.10768	1.10498	1.09944	1.09368	1.08772
7	1.12823	1.12309	1.12045	1.11777	1.11505	1.11228	1.10661	1.10077	1.09476
8	1.13610	1.13076	1.12803	1.12526	1.12246	1.11963	1.11385	1.10792	1.10186
9	1.14402	1.13848	1.13566	1.13282	1.12995	1.12704	1.12115	1.11514	1.10902
0	1.15199	1.14625	1.14335	1.14043	1.13748	1.13451	1.12851	1.12242	1.11625
1	1.15998	1.15407	1.15109	1.14809	1.14508	1.14205	1.13594	1.12977	1.12353
2	1.16803	1.16194	1.15888	1.15581	1.15273	1.14964	1.14343	1.13718	1.13089
3	1.17611	1.16986	1.16673	1.16359	1.16045	1.15731	1.15100	1.14467	1.13832
4	1.18424	1.17784	1.17464	1.17143	1.16823	1.16503	1.15862	1.15221	1.14579
5	1.19240	1.18586	1.18260	1.17933	1.17607	1.17282	1.16631	1.15982	1.15335
6	1.20061	1.19393	1.19060	1.18728	1.18396	1.18066	1.17406	1.16749	1.16096
7	1.20885	1.20204	1.19865	1.19527	1.19190	1.18854	1.18186	1.17522	1.16862
8	1.21710	1.21019	1.20675	1.20332	1.19990	1.19650	1.18973	1.18302	1.17635
9	1.22539	1.21838	1.21489	1.21142	1.20796	1.20452	1.19767	1.19087	1.18414
0	1.23370	1.22661	1.22308	1.21957	1.21607	1.21259	1.20566	1.19879	1.19198
1	1.24204	1.23487	1.23131	1.22776	1.22423	1.22071	1.21371	1.20677	1.19989
2	1.25038	1.24316	1.23957	1.23600	1.23244	1.22887	1.22179	1.21476	1.20779
3	1.25878	1.25151	1.24789	1.24429	1.24069	1.23712	1.22999	1.22292	1.21589
4	1.26723	1.25990	1.25626	1.25263	1.24901	1.24540	1.23822	1.23109	1.22400
5	1.27571	1.26834	1.26468	1.26102	1.25738	1.25375	1.24652	1.23933	1.23219
6	1.28424	1.27683	1.27314	1.26947	1.26580	1.26214	1.25487	1.24763	1.24045
7	1.29283	1.28538	1.28167	1.27797	1.27429	1.27061	1.26329	1.25601	1.24878
8	1.30149	1.29400	1.29027	1.28655	1.28284	1.27915	1.27179	1.26448	1.25721
9	1.31022	1.30268	1.29894	1.29520	1.29148	1.28776	1.28038	1.27304	1.26575
0	1.31901	1.31144	1.30767	1.30392	1.30018	1.29646	1.28905	1.28169	1.27440
1	1.32788	1.32027	1.31648	1.31271	1.30896	1.30522	1.29779	1.29042	1.28311
2	1.33683	1.32917	1.32537	1.32158	1.31782	1.31407	1.30662	1.29924	1.29193
3	1.34587	1.33817	1.33435	1.33054	1.32676	1.32300	1.31553	1.30813	1.30081
4	1.35501	1.34727	1.34342	1.33960	1.33580	1.33202	1.32452	1.31710	1.30976
5	1.36425	1.35647	1.35261	1.34877	1.34496	1.34116	1.33363	1.32618	1.31881
6	1.37361	1.36579	1.36191	1.35805	1.35422	1.35040	1.34284	1.33536	1.32797
7	1.38308	1.37522	1.37132	1.36744	1.36359	1.35975	1.35215	1.34464	1.33721
8	1.39267	1.38476	1.38084	1.37694	1.37306	1.36921	1.36157	1.35401	1.34655
9	1.40238	1.39441	1.39047	1.38654	1.38264	1.37877	1.37108	1.36349	1.35600
0	1.41219	1.40418	1.40021	1.39627	1.39235	1.38845	1.38073	1.37310	1.36556
1	1.42214	1.41407	1.41007	1.40610	1.40215	1.39823	1.39047	1.38280	1.37524
2	1.43220	1.42408	1.42005	1.41605	1.41208	1.40814	1.40033	1.39262	1.38502

Sp. gr. of H₂SO₄ + Aq. at t°. Sp. gr. of H₂O at 15° = 1 *Continued*

% H ₂ SO ₄	0°	10°	15°	20°	25°	30°	40°	50°	60°
53	1.44239	1.43420	1.43014	1.42611	1.42211	1.41814	1.41028	1.40254	1.39
54	1.45269	1.44443	1.44034	1.43628	1.43225	1.42825	1.42034	1.41255	1.40
55	1.46311	1.45477	1.45065	1.44656	1.44250	1.43847	1.43051	1.42268	1.41
56	1.47364	1.46523	1.46107	1.45695	1.45285	1.44880	1.44078	1.43290	1.41
57	1.48427	1.47578	1.47159	1.46743	1.46331	1.45922	1.45115	1.44322	1.41
58	1.49499	1.48643	1.48221	1.47802	1.47387	1.46975	1.46162	1.45364	1.41
59	1.50583	1.49719	1.49292	1.48870	1.48452	1.48037	1.47218	1.46415	1.41
60	1.51676	1.50804	1.50374	1.49949	1.49527	1.49109	1.48285	1.47476	1.41
61	1.52778	1.51899	1.51465	1.51036	1.50611	1.50190	1.49360	1.48546	1.41
62	1.53889	1.53002	1.52564	1.52132	1.51703	1.51278	1.50442	1.49622	1.41
63	1.55008	1.54113	1.53672	1.53236	1.52804	1.52376	1.51533	1.50708	1.41
64	1.56135	1.55233	1.54788	1.54348	1.53913	1.53481	1.52632	1.51801	1.51
65	1.57270	1.56360	1.55912	1.55469	1.55030	1.54595	1.53740	1.52903	1.51
66	1.58414	1.57496	1.57044	1.56597	1.56154	1.55716	1.54854	1.54011	1.51
67	1.59565	1.58640	1.58184	1.57733	1.57287	1.56846	1.55978	1.55128	1.51
68	1.60724	1.59792	1.59332	1.58878	1.58427	1.57981	1.57104	1.56246	1.51
69	1.61892	1.60951	1.60488	1.60030	1.59577	1.59129	1.58247	1.57384	1.51
70	1.63068	1.62118	1.61651	1.61189	1.60732	1.60280	1.59391	1.58521	1.51
71	1.64251	1.63293	1.62821	1.62355	1.61894	1.61437	1.60540	1.59663	1.51
72	1.65439	1.64473	1.63997	1.63527	1.63062	1.62601	1.61696	1.60811	1.51
73	1.66633	1.65658	1.65178	1.64704	1.64234	1.63769	1.62855	1.61961	1.61
74	1.67831	1.66847	1.66362	1.65883	1.65408	1.64939	1.64015	1.63111	1.61
75	1.69030	1.68037	1.67547	1.67063	1.66584	1.66109	1.65175	1.64260	1.61
76	1.70228	1.69225	1.68731	1.68242	1.67757	1.67278	1.66332	1.65405	1.61
77	1.71424	1.70411	1.69911	1.69416	1.68926	1.68439	1.67481	1.66540	1.61
78	1.72615	1.71589	1.71083	1.70582	1.70085	1.69591	1.68616	1.67658	1.61
79	1.73798	1.72758	1.72243	1.71735	1.71231	1.70731	1.69741	1.68767	1.61
80	1.74970	1.73909	1.73386	1.72868	1.72356	1.71847	1.70842	1.69854	1.61
81	1.76120	1.75038	1.74504	1.73979	1.73458	1.72942	1.71921	1.70916	1.61
82	1.77244	1.76140	1.75595	1.75057	1.74524	1.73998	1.72962	1.71945	1.71
83	1.78312	1.77193	1.76642	1.76097	1.75557	1.75022	1.73972	1.72943	1.71
84	1.79316	1.78191	1.77636	1.77087	1.76543	1.76006	1.74943	1.73902	1.71
85	1.80250	1.79123	1.78567	1.78016	1.77470	1.76929	1.75863	1.74816	1.71
86	1.81108	1.79982	1.79428	1.78878	1.78331	1.77789	1.76721	1.75674	1.71
87	1.81887	1.80767	1.80214	1.79666	1.79123	1.78584	1.77519	1.76473	1.71
88	1.82589	1.81476	1.80926	1.80381	1.79839	1.79302	1.78242	1.77199	1.71
89	1.83216	1.82111	1.81564	1.81022	1.80484	1.79950	1.78895	1.77856	1.71
90	1.83771	1.82677	1.82135	1.81597	1.81063	1.80532	1.79483	1.78448	1.71
91	1.84263	1.83179	1.82642	1.82109	1.81580	1.81054	1.80013	1.78985	1.71
92	1.84691	1.83619	1.83088	1.82561	1.82037	1.81516	1.80487	1.79471	1.71
93	1.85059	1.83997	1.83471	1.82950	1.82432	1.81918	1.80902	1.79900	1.71
94	1.85363	1.84311	1.83790	1.83275	1.82763	1.82255	1.81253	1.80266	1.71
95	1.85598	1.84555	1.84040	1.83526	1.83022	1.82520	1.81528	1.80553	1.71
96	1.85765	1.84729	1.84217	1.83709	1.83207	1.82708	1.81724	1.80758	1.71
97	1.85854	1.84816	1.84305	1.83798	1.83297	1.82800	1.81822	1.80863	1.71
98	1.85836	1.84789	1.84275	1.83766	1.83264	1.82767	1.81792	1.80840	1.71
99	1.85671	1.84612	1.84093	1.83581	1.83076	1.82578	1.81604	1.80658	1.71
100	(1.85330)	(1.84255)	(1.83729)	(1.83213)	(1.82705)	(1.82205)	(1.81231)	(1.80288)	(1.79

Auszug aus Band 5 der wissenschaftlichen Abhandlungen der Normaleichungskommis
Berlin 1904, P. 257. Springer's publication.

(Domke, Z. anorg. 1905, 43. 176.)

Sp. gr. of $\text{H}_2\text{SO}_4 + \text{Aq}$ at $15^\circ/15^\circ$ in air.—
Continued from page 915.

Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4	Sp. gr.	% H_2SO_4
1.816	89.11	1.828	91.30	1.840	94.57
1.817	89.27	1.829	91.52	1.841	94.96
1.818	89.44	1.830	91.74	1.842	95.40
1.819	89.61	1.831	91.98	1.843	96.02
1.820	89.79	1.832	92.22	1.844	96.93
1.821	89.97	1.833	92.46	1.8442	97.50
1.822	90.15	1.834	92.71	1.844	99.08
1.823	90.33	1.835	92.98	1.843	99.84
1.824	90.51	1.836	93.26	1.842	99.29
1.825	90.70	1.837	93.56	1.841	98.61
1.826	90.90	1.838	93.87	1.840	98.88
1.827	91.10	1.839	94.20	1.8394	100.00

Lunge, calculated by Marshall, J. Soc. Chem Ind. 1902, **21**, 1509.

Sp. gr. at 20° of $\text{H}_2\text{SO}_4 + \text{Aq}$ containing
M g. mols. H_2SO_4 per liter.

M.	0.01	0.025	0.05
Sp. gr.	1.000719	1.001907	1.003551
M.	0.075	0.10	0.25
Sp. gr.	1.005152	1.00677	1.01618
M.	0.50	0.75	1.0
Sp. gr.	1.03218	1.04760	1.06307
M.	1.5	2.0	
Sp. gr.	1.09345	1.12316	

(Jones and Pearce, Am. Ch. J. 1907, **38**, 733.)

Boiling-point of $\text{H}_2\text{SO}_4 + \text{Aq}$.

% H_2SO_4	B. pt.	% H_2SO_4	B. pt.
5	101.0°	70	170.0°
10	102.0	72	174.5
15	105.5	74	180.5
20	105.0	76	189.0
25	106.5	78	199.0
30	108.0	80	207.0
35	110.0	82	218.5
40	114.0	84	227.0
45	118.5	86	238.5
50	124.0	88	251.5
53	129.5	90	262.5
56	133.0	91	268.0
60	141.5	92	274.5
62.5	147.0	93	281.5
65	153.5	94	288.5
67.5	161.0	95	295.0

(Lunge, B. **11**, 370.)

Freezing- and melting-points of $\text{H}_2\text{SO}_4 + \text{Aq}$.

Sp. gr. at 15°	F. pt.	M. pt.
1.671	liq. at 20°	
1.691	"	
1.712	"	
1.727	-7.5	-7.5
1.732	-8.5	-8.5
1.749	-0.2	+4.5
1.767	+1.6	+6.5
1.790	+4.5	+8.0
1.807	-9.0	-6.0
1.822	liq. at -20°	
1.842	"	

(Lunge, B. **15**, 2644.)

Effect of impurities on sp. gr. of $\text{H}_2\text{SO}_4 + \text{Aq}$.

The figures show the increase in sp. gr. of $\text{H}_2\text{SO}_4 + \text{Aq}$ caused by adding 0.1% of an impurity to acid of different strengths.

Salt	100 %	98 %	94 %	80 %	70 %
Na_2SO_4	0.0011	0.0010	0.0007	0.0008	0.0007
CaSO_4	0.0012	0.0011	0.0009	0.0007	0.0006
$\text{Al}_2(\text{SO}_4)_3$	insol	insol	insol	0.0012	0.0011
$\text{Fe}_2(\text{SO}_4)_3$	"	"	0.0003	0.0008	0.0007
PbSO_4	0.0017	0.0014	0.0015	insol	insol.
MgSO_4	0.0011	0.0010	0.0012	0.0009	0.0009
As_2O_3	0.0013	0.0013		0.0010	
HSO_4	0.0002	0.00027		0.00023	

(Marshall, J. Soc. Chem. Ind. 1902, **21**, 1508.)

Sp. gr. of mixtures of H_2SO_4 (96.5%) and
 HNO_3 (94%) at $18^\circ/18^\circ$ in air.

% HNO_3 in mixture	Sp. gr.	% HNO_3 in mixture	Sp. gr.
0.00	1.8437	22.51	1.8215
0.57	1.8456	25.56	1.8112
1.05	1.8476	27.29	1.8053
4.67	1.8586	32.53	1.7863
7.17	1.8618	37.03	1.7700
7.37	1.8620	39.49	1.7601
7.75	1.8619	57.78	1.6879
9.10	1.8605	72.89	1.6227
11.33	1.8557	90.76	1.5408
12.71	1.8520	98.19	1.5080
16.52	1.8414	100.00	1.5009

(Marshall, J. Soc. Chem. Ind. 1902, **21**, 1508.)

Miscible with alcohol, with evolution of heat and formation of ethylsulphuric acid.
 $+\text{H}_2\text{O} = \text{H}_2\text{SO}_4$, also called tetrahydroxyl sulphuric acid. (Marignac, A. ch. (3) **39**, 184.)

Mpt. 8.35° . (Pickering.)

$+2\text{H}_2\text{O} = \text{H}_2\text{SO}_6$, also called perhydroxyl sulphuric acid.

Mpt. -38.9° . (Biron, J. Russ. Phys. Chem. Soc. 1899, **31**, 517.)

$+3\text{H}_2\text{O}$. (Pickering, Chem. Soc. 1890, **57**, 331.)

+4H₂O. Mpt. -75°. (Pickering, Chem. Soc. 1890, 57. 331.)

Sp. gr. and fr. pt. of hydrates of H₂SO₄.

Hydrate	H ₂ SO ₄	Sp. gr. of the liquid	Fr. pt.
H ₂ SO ₄ (pure)	100	1.842	+10.5
H ₂ SO ₄ +H ₂ O	84.48	1.777	+3.5
H ₂ SO ₄ +2H ₂ O	73.08	1.650	-70.0
H ₂ SO ₄ +4H ₂ O	57.65	1.476	-40.0
H ₂ SO ₄ +6H ₂ O	47.57	1.376	-50.0
H ₂ SO ₄ +8H ₂ O	40.50	1.311	-65.0
H ₂ SO ₄ +10H ₂ O	35.25	1.268	-88.0
H ₂ SO ₄ +11H ₂ O	33.11	1.249	-75.0
H ₂ SO ₄ +12H ₂ O	31.21	1.233	-55.0
H ₂ SO ₄ +13H ₂ O	29.52	1.219	-45.0
H ₂ SO ₄ +14H ₂ O	28.00	1.207	-40.0
H ₂ SO ₄ +15H ₂ O	26.63	1.196	-34.0
H ₂ SO ₄ +16H ₂ O	25.39	1.187	-25.6
H ₂ SO ₄ +18H ₂ O	23.22	1.170	-19.0
H ₂ SO ₄ +20H ₂ O	21.40	1.157	-17.0
H ₂ SO ₄ +25H ₂ O	17.88	1.129	-8.5
H ₂ SO ₄ +50H ₂ O	9.82	1.067	-3.5
H ₂ SO ₄ +75H ₂ O	6.77	1.045	0.0
H ₂ SO ₄ +100H ₂ O	5.16	1.032	+2.5
H ₂ SO ₄ +300H ₂ O	1.78	1.007	+4.5
H ₂ SO ₄ +1000H ₂ O	0.54	1.001	+0.5

(Pictet, C. R. 1894, 119. 645.)

Sulphuric acid, anhydrous, SO₃.

See Sulphur trioxide.

Disulphuric (Pyrosulphuric) acid, H₂S₂O₇.

Very deliquescent. Miscible with H₂O. Sol. in fuming H₂SO₄. Miscible in liquid SO₂. (Schultz-Sellack.)

H₂S₂O₇, 2H₂SO₄. Fumes on air. (Jacquelin, A. ch. (3) 30. 343.)

Tetrasulphuric acid, H₂S₄O₁₃.

Fumes on air. (Weber, Pogg. 159. 313.)

Sulphates.

Most sulphates are easily sol. in H₂O; but Ag₂SO₄, Hg₂SO₄, and CaSO₄ are only sl. sol., while BaSO₄, SrSO₄, and PbSO₄ are nearly insol. therein. All sulphates are sol. in conc. H₂SO₄. Basic sulphates are insol. in H₂O. Most sulphates are insol. in alcohol.

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 823.)

Aluminum sulphate, basic, 2Al₂O₃.SO₃+5H₂O.

Slowly sol. in 10 mols. HC₂H₃O₂. (Schlumberger, Bull. Soc. 1895, (3) 12. 41.)

+7H₂O. Easily sol. in 8 mols. dil. HCl+Aq. or in 10 mols. 10% acetic acid in 24 hours. (Schlumberger.)

+10H₂O. Insol. in H₂O; easily sol. in cold dil. mineral acids, and HC₂H₃O₂+Aq. (Crum, A. 89. 174.)

Min. *Felsöbanyite*.

+15H₂O. Min. *Paraluminite*.

8Al₂O₃.5SO₃+25H₂O. Insol. in H₂O in dil. acids. (Löwe, J. pr. 79. 428.)

5Al₂O₃.3SO₃+20H₂O. Easily sol. (Debray, Bull. Soc. (2) 7. 9.)

3Al₂O₃.2SO₃+9H₂O. Nearly in conc. H₂SO₄. (Bayer, Dingl. 363. 21.)

+20H₂O. Ppt.

4Al₂O₃.3SO₃+36H₂O. Insol. in dil. mineral acids, easily sol. in dil. HC₂H₃O₂+Aq. (Debray, Bull. Soc. (2) 7. 9.)

Al₂O₃.SO₃+6H₂O=(AlO)₂SO₄. Insol. in H₂O or HC₂H₃O₂+Aq. Sl. hot HCl, easily sol. in warm KC₂H₃O₂. (Böttger, A. 244. 225.)

+9H₂O. (Athanasesco, C. R. 101.)

Min. *Aluminite*.

[Al₂(OH)₃]₂SO₄+2H₂O.

Sol. in HCl+Aq. in the cold with. Very unstable. (Schlumberger, Bull. Soc. 1895, (3) 12. 60.)

3Al₂O₃.4SO₃+9H₂O. (Athanasesco 103. 271.)

+30H₂O. Sol. in 144 pts. cold, a pts. boiling H₂O. Easily sol. in H₂NO₃+Aq. (Rammelsberg, Pogg. 4.)

2Al₂O₃.3SO₃. Decomp. by H₂. 3Al₂O₃.SO₃ and Al₂(SO₄)₃. (Maus.)

Al₂O₃.2SO₃=Al₂O(SO₄)₂.

Min. *Alumaine*.

+H₂O. Sol. in small quantity of H₂O decomp. by a large quantity into (Al and Al₂(SO₄)₃). (Maus, Pogg. 11. 80.)

+12H₂O. Easily sol. in hot or cold. Sat. solution contains 45% salt at 15°. crystallises unchanged on evaporating. (guerite, C. R. 90. 354.)

Above basic compounds are miscible. (Pickering, C. N. 45. 121, 133, 146.)

Aluminum sulphate, Al₂(SO₄)₃.

100 pts. H₂O dissolve (a) pts. Al₂(SO₄)₃ and (b) pts. Al₂(SO₄)₃+18H₂O at:

	0°	10°	20°	30°	40°
a	31.3	33.5	36.15	40.36	45.73
b	86.85	95.8	107.35	127.6	167.6

	60°	70°	80°	90°	100°
a	59.09	66.23	73.14	80.83	98
b	262.6	348.2	467.3	678.8	1111

(Poggiale, A. ch. (3) 8. 467.)

See also +18H₂O.

Sp. gr. of Al₂(SO₄)₃+Aq.

Al ₂ (SO ₄) ₃	Sp. gr. at		
	15°	25°	35°
5	1.0569	1.0503	1.045
10	1.1071	1.1022	1.096
15	1.1574	1.1522	1.146
20	1.2074	1.2004	1.192
25	1.2572	1.2487	1.2407

(Reuss, B. 17. 2888.)

Sp. gr. of $\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$ at 15° containing:
 10 20 30% $\text{Al}_2(\text{SO}_4)_3 + 18\text{H}_2\text{O}$,
 0.635 1.1105 1.1710
 40 50% $\text{Al}_2(\text{SO}_4)_3 + 18\text{H}_2\text{O}$.
 1.2355 1.3050

Sp. gr. of sat. solution = 1.34.
 (Gerlach, Z. anal. 28. 493.)

Sp. gr. of $\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$ at 25° .

Strength of $\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$	Sp. gr.
1 normal	1.0550
$\frac{1}{2}$ "	1.0278
$\frac{1}{4}$ "	1.0138
$\frac{1}{8}$ "	1.0068

(Wagner, Z. phys. Ch. 1890, 5. 35.)

100 pts. of a mixture of 1 vol. $\text{H}_2\text{SO}_4 + 2$ vols. H_2O dissolve only 6.45 pts. $\text{Al}_2(\text{SO}_4)_3$. (Baud, C. R. 1903, 137. 494.)

$\text{Al}_2(\text{SO}_4)_3$ is completely pptd. from $\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$ by an excess of glacial $\text{HC}_2\text{H}_3\text{O}_2$. (Person, A. ch. (2) 63. 444.)

Solubility of $\text{Al}_2(\text{SO}_4)_3 + (\text{NH}_4)_2\text{Al}_2(\text{SO}_4)_4$.
 See under $(\text{NH}_4)_2\text{Al}_2(\text{SO}_4)_4$.

Solubility of $\text{Al}_2(\text{SO}_4)_3 + \text{K}_2\text{Al}_2(\text{SO}_4)_4$. See under $\text{K}_2\text{Al}_2(\text{SO}_4)_4$.

Solubility in $\text{Fe}_2(\text{SO}_4)_3 + \text{Aq}$ at 25° .

100 g. of the solution contain

g. $\text{Al}_2(\text{SO}_4)_3$	g. $\text{Fe}_2(\text{SO}_4)_3$
27.82	0
26.01	6.064
24.21	9.819
21.64	13.02
15.22	23.28
*10.70	31.91
10.23	31.90

*Solution sat. with respect to both salts.
 (Wirth and Bakke, Z. anorg. 1914, 87. 48.)
 See also under $\text{Fe}_2(\text{SO}_4)_3$.

Solubility of $\text{Al}_2(\text{SO}_4)_3 + \text{Li}_2\text{SO}_4$ at 30° .

Composition of				Solid phase
Solution		Residue		
% Li_2SO_4	% $\text{Al}_2(\text{SO}_4)_3$	% Li_2SO_4	% $\text{Al}_2(\text{SO}_4)_3$	
25.1	0	$\text{Li}_2\text{SO}_4, \text{H}_2\text{O}$
21.93	5.34	"
16.10	14.89	63.70	4.02	"
13.63	20.76	14.72	31.17	$\text{Li}_2\text{SO}_4, \text{H}_2\text{O} +$ $\text{Al}_2(\text{SO}_4)_3, 18\text{H}_2\text{O}$
13.24	21.71	61.24	7.22	$\text{Li}_2\text{SO}_4, 4\text{H}_2\text{O}$
11.73	22.08	6.92	33.54	$\text{Al}_2(\text{SO}_4)_3, 18\text{H}_2\text{O}$
6.75	24.34	3.77	37.06	"
3.44	26.12	"
0.00	28.0	"

(Schreinemakers and de Waal, Ch. Weekbl. 1906, 3. 539.)

100 g. of sat. solution of $\text{Al}_2(\text{SO}_4)_3$ in glycol contain 14.4 g. $\text{Al}_2(\text{SO}_4)_3$. (de Coninck, Bull. Ac. Roy. Belg. 1905. 359.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone. (Naumann, B. 1904, 37. 4328.)

+6 H_2O . Very slowly sol. in cold, completely sol. in hot H_2O .

+8 H_2O . (Margueritte-Delarcharbonny, C. R. 112. 229.)

+10 H_2O . Deliquescent. (v. Hauer, W. A. B. 13. 449.)

+16 H_2O . Sol. in conc. H_2SO_4 . (Baud, C. R. 1903, 137. 494.)

+17 H_2O . (Gawalowski, C. C. 1885. 721.)

+18 H_2O . Permanent. (Berzelius.)

100 g. of the aqueous solution contain 27.82 g. $\text{Al}_2(\text{SO}_4)_3$ at 25° . (Wirth, Z. anorg. 1913, 79. 361.)

Solubility of $\text{Al}_2(\text{SO}_4)_3 + 18\text{H}_2\text{O}$ in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

$\text{H}_2\text{SO}_4 + \text{Aq}$ % H_2SO_4	100 g. of the solution contain g. $\text{Al}_2(\text{SO}_4)_3$
0	27.82
5.23	29.21
9.90	26.21
18.70	20.44
25.50	15.40
40.70	5.07
52.25	1.216
63.70	1.243
73.64	2.915

(Wirth, Z. anorg. 1913, 79. 361.)

Hydrous salt is scarcely sol. in alcohol. (Berzelius.)

Min. *Alunogen*.

+27 H_2O . Efflorescent. (Margueritte-Delarcharbonny, C. R. 99. 800.)

Aluminum sulphate, acid, $\text{Al}_2\text{O}_3, 4\text{SO}_3 + 4\text{H}_2\text{O}$.

Extremely slowly sol. in cold, more rapidly in hot H_2O . (Baud, C. R. 1903, 137. 493.)

$\text{Al}_2\text{O}_3, 6\text{SO}_3 + 10\text{H}_2\text{O}$. Sol. in H_2O ; solution soon decomp. into $\text{Al}_2(\text{SO}_4)_3 + \text{H}_2\text{SO}_4$. (Silberberger, M. 1904, 25. 221.)

Aluminum ammonium sulphate (Ammonia alum), $(\text{NH}_4)_2\text{Al}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

100 pts. H_2O dissolve 2.9 pts. anhydrous salt at 0° ; 207.7 pts. anhydrous salt at 110.6° . (Mulder.)

100 pts. H_2O dissolve 8.74 pts. anhydrous salt at 17.5° . (Pohl, W. A. B. 6. 597.)

100 pts. H₂O at t° dissolve pts.
(NH₄)₂Al₂(SO₄)₄.

Solubility of NH₄ alum in presence of
(NH₄)₂SO₄ and Al₂(SO₄)₃.

t°	Pts. (NH ₄) ₂ Al ₂ (SO ₄) ₄	Pts. (NH ₄) ₂ Al ₂ (SO ₄) ₄ + 24H ₂ O
0	2.10	3.90
10	4.99	9.52
20	7.74	15.13
30	10.94	22.01
40	14.88	30.92
50	20.09	44.11
60	26.70	66.65
70	35.11	90.67
80	45.66	134.47
90	58.68	209.31
100	74.53	357.48

Mixture used	100 g. sat. solution contains	
	g. (NH ₄) ₂ SO ₄	g. Al ₂ (SO ₄) ₃
Sat. NH ₄ alum at 18.5°	1.42	3.69
20 cc. above sol- ution + 6 g. cryst. Al ₂ (SO ₄) ₃	0.45	16.09
20 cc. above sol- ution + 4g. (NH ₄) ₂ SO ₄	20.81	0.29

(Rüdorff, 1885, B. 18. 1160.)

(Poggiale, A. ch. (3) 8. 467.)

Insol. in alcohol. (Mulder.)
Solubility of Al(NH₄)(SO₄)₂ + 12H₂O in a
mixture of 93.3 g. H₂O and 23.33 g. glycerine
= 6.15 g. (Dunlop, Pharm. J. 1910, 88. 6.)
Solubility in 93.3 g. H₂O + 23.3 g. glycerine
+ 3.9 g. phenol = 5.59 g. Al(NH₄)(SO₄)₂
+ 12H₂O. (Dunlop.)
Min. *Tschermigite*.

Aluminum ammonium chromium sulphate,
Al₂(SO₄)₃, (NH₄)₂SO₄, Cr₂(SO₄)₃ +
48H₂O.

Sol. in H₂O; decomp. by boiling. (Vohl,
A. 94. 71.)

Aluminum caesium sulphate, Al₂Ca₂(SO₄)₄ +
24H₂O.

100 pts. H₂O at 17° dissolve 0.619 pt.
caesium alum. (Redtenbacher, J. pr. 94.
442.)

Solubility in 100 pts. H₂O at t° (calculated for
salt dried at 130°).

t°	Pts. alum	t°	Pts. alum	t°	Pts. alum
0	0.19	25	0.49	65	2.38
10	0.29	35	0.69	80	5.29
17	0.38	50	1.235

(Setterberg, A. 211. 104.)

Solubility in H₂O.

t°	Pts. anhydrous salt per litre	G. mole anhydrous salt per litre
25	4.7	0.013
30	5.89	0.0167
35	7.29	0.0207
40	9.00	0.0256

(Locke, Am. Ch. J. 1901, 28. 180.)

According to Locke (Am. Ch. J. 1901, 28.
174), Poggiale's tables for NH₄ and K alums
are evidently transposed, and the above data
are applied by Poggiale to K alum.

1 l. H₂O dissolves 91.9 g. anhydrous, or
191.9 g. hydrated salt, or 0.387 mols. an-
hydrous salt at 25°. (Locke, Am. Ch. J.
1901, 28. 175.)

Solubility in H₂O at t°.

t°	G. (NH ₄) ₂ Al ₂ (SO ₄) ₄ per 100 g. H ₂ O	G. (NH ₄) ₂ Al ₂ (SO ₄) ₄ + 24H ₂ O per 100 g. H ₂ O	G. mol. (NH ₄) ₂ Al ₂ (SO ₄) ₄ per 100 g. H ₂ O
0	2.10	3.90	0.0044
5	3.50	6.91	0.0074
10	4.99	9.52	0.0105
15	6.25	12.66	0.0132
20	7.74	15.13	0.0163
25	9.19	19.19	0.0194
30	10.94	22.01	0.0231
40	14.88	30.92	0.0314
50	20.10	44.10	0.0424
60	26.70	66.65	0.0569
95	109.7	00	0.2312

(Mulder, Poggiale, Locke; Marino, Gazz.
ch. it. 1905, 35. II, 351; Berkeley, Trans.
Roy. Soc. 1904, 203. A, 214, calc. by Seidell,
Solubilities.)

B.-pt. of sat. solution is 110.6°.
M.-pt. of (NH₄)₂Al₂(SO₄)₄ + 24H₂O = 92°.
(Tilden, Chem. Soc. 45. 409.); = 95°. (Locke,
L. c.)

Sp. gr. of aqueous solution at 15° contain
ing:

3% 6% 9%
1.0423 1.0141 1.0282 hydrous salt.

(Gerlach, Z. anal. 28. 495.)

solubility of $\text{Al}_2\text{Cr}_2(\text{SO}_4)_4$ in H_2O at t° .
 $\text{Al}_2\text{Cr}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$ in 100 g. solution.)

	% salt	t°	% salt
3	0.21	75	4.12
5	0.35	80	5.21
3	0.60	90	9.50
5	1.04	100.4	18.60
3	1.96		

Keley, Trans. Roy. Soc. 1904, 203. A, 214.)

Solubility in 100 g. H_2O at t° .

$\text{Al}_2(\text{SO}_4)_3$	t°	$\text{Al}_2(\text{SO}_4)_3$	t°	$\text{Al}_2(\text{SO}_4)_3$	t°	$\text{Al}_2(\text{SO}_4)_3$
0.19	26	0.50	52	1.45	78	5.15
0.20	27	0.51	53	1.51	79	5.40
0.21	28	0.52	54	1.58	80	5.78
0.22	29	0.55	55	1.65	81	6.05
0.23	30	0.57	56	1.71	82	6.4
0.24	31	0.59	57	1.77	83	6.7
0.25	32	0.60	58	1.86	84	7.0
0.26	33	0.62	59	1.92	85	7.4
0.27	34	0.65	60	2.06	86	7.7
0.28	35	0.69	61	2.14	87	8.0
0.29	36	0.72	62	2.25	88	8.3
0.30	37	0.75	63	2.37	89	8.6
0.31	38	0.77	64	2.50	90	8.8
0.32	39	0.80	65	2.65	91	9.0
0.34	40	0.85	66	2.78	92	9.2
0.35	41	0.87	67	2.96	93	9.5
0.36	42	0.91	68	3.13	94	9.9
0.38	43	0.96	69	3.34	95	10.1
0.39	44	1.01	70	3.50	96	10.4
0.40	45	1.06	71	3.67	97	10.8
0.41	46	1.10	72	3.85	98	11.1
0.42	47	1.17	73	4.07	99	11.5
0.43	48	1.21	74	4.30	100	12.0
0.45	50	1.27	75	4.50		
0.47	50	1.30	76	4.72		
0.49	51	1.39	77	4.95		

values from 0-7° obtained by interpolation
 of Setterberg's values for 0° (A. 1882, 211.)

From 80-100° they were calculated by
 interpolation.

Port and Huselton, J. Am. Chem. Soc. 1914,
 36, 2084.)

Alfolds in crystal H_2O at 106° (Tilden,
 Ann. Soc. 45. 409); at 120.5° (Erdmann);
 122° (Locke.)

Aluminum calcium sulphate, basic, Al_2O_3 ,
 6CaO , $3\text{SO}_3 + 32\text{H}_2\text{O}$.

Min. *Ettringite*. Mostly sol. in H_2O ; sol.
 in $\text{HCl} + \text{Aq}$.

Aluminum chromium sulphate, $\text{Al}_2\text{Cr}_2(\text{SO}_4)_4$.
 Insol. in H_2O .

$\text{Al}_2\text{Cr}_2(\text{SO}_4)_4 \cdot \text{H}_2\text{SO}_4$. Insol. in H_2O . (Étard
 C. R. 86. 1400.)

Aluminum chromium potassium sulphate,
 $\text{Al}_2(\text{SO}_4)_3$, $\text{Cr}_2(\text{SO}_4)_3$, $2\text{K}_2\text{SO}_4 + 48\text{H}_2\text{O}$.

Sol. in H_2O , but decomp. on boiling. (Vohl.)

Aluminum copper sulphate, $2\text{Al}_2\text{O}_3$, 9CuO ,
 $3\text{SO}_3 + 21\text{H}_2\text{O}$.

Min. *Cyanotrichite*. (Percy, Phil. Mag. (3)
 86. 103.)

Aluminum hydroxylamine sulphate,
 $\text{Al}_2(\text{SO}_4)_3$, $(\text{NH}_2\text{OH})_2\text{SO}_4 + 24\text{H}_2\text{O}$.

Sol. in H_2O . (Meyeringh, B. 10. 1946.)

Aluminum iron (ferrous) sulphate, $\text{Al}_2(\text{SO}_4)_3$,
 $\text{FeSO}_4 + 24\text{H}_2\text{O}$.

Sol. in H_2O . (Klauer, A. 14. 261.)

Min. *Halotrichite*.

$\text{Al}_2(\text{SO}_4)_3$, $2\text{FeSO}_4 + 27\text{H}_2\text{O}$. Sol. in H_2O .
 (Berthier.)

Al_2O_3 , 2SO_3 , 6FeSO_4 . Easily sol. in H_2O .
 (Phillips.)

$\text{Al}_2(\text{SO}_4)_3$, 2FeSO_4 , H_2SO_4 . Insol. in H_2O .
 (Étard, C. R. 87. 602.)

Aluminum iron (ferric) sulphate, $\text{Al}_2(\text{SO}_4)_3$,
 $\text{Fe}_2(\text{SO}_4)_3$.

Insol. in H_2O . (Étard, C. R. 86. 1399.)

$\text{Al}_2(\text{SO}_4)_3$, $\text{Fe}_2(\text{SO}_4)_3$, H_2SO_4 . As above
 (Étard.)

See $\text{Al}_2(\text{SO}_4)_3 + \text{Fe}_2(\text{SO}_4)_3$, under $\text{Al}_2(\text{SO}_4)_3$
 and $\text{Fe}_2(\text{SO}_4)_3$.

Aluminum ferrous potassium sulphate,
 $\text{Al}_2(\text{SO}_4)_3$, 12FeSO_4 , $2\text{K}_2\text{SO}_4 + 24\text{H}_2\text{O}$.

Permanent. Sl. sol. in H_2O . (Dufrenoy.)

Aluminum lead sulphate, $\text{Al}_2\text{Pb}_2(\text{SO}_4)_4 +$
 $20\text{H}_2\text{O}$.

Permanent; insol. in H_2O . (G. H. Bailey
 J. Chem. Soc. Ind. 6. 415.)

Aluminum lithium sulphate, $\text{Li}_2\text{Al}_2(\text{SO}_4)_4 +$
 $24\text{H}_2\text{O}$.

Sol. in 24 pts. cold, and 0.87 pt. hot H_2O .
 (Kralovansky, Schw. J. 54. 349.)

Does not exist. (Rammelsberg, J. B. 1847-
 48. 394; Arfvedson; Gmelin.)

Aluminum lithium potassium sulphate (?).

Sol. in H_2O , from which it crystallises on
 cooling. (Joss, J. pr. 1. 142.)

Aluminum magnesium sulphate, MgSO_4 ,
 $\text{Al}_2(\text{SO}_4)_3 + 22\text{H}_2\text{O}$.

Min. *Pickeringite*.

2MgSO_4 , $\text{Al}_2(\text{SO}_4)_3 + 22\text{H}_2\text{O}$. Min. *Pic-*
raluminite.

3MgSO_4 , $\text{Al}_2(\text{SO}_4)_3 + 36\text{H}_2\text{O}$. Very sol. in
 H_2O . (Klauer, A. 14. 284.)

Aluminum magnesium manganous sulphate,
 $\text{Al}_2(\text{SO}_4)_3, \text{MgSO}_4, \text{MnSO}_4 + 25\text{H}_2\text{O}$.
 As sol. in H_2O as K alum. (Kane.) Very
 sol. in H_2O . (Smith, Sill. Am. J. (2) 18. 379.)
 Min. *Bosjemanite*.

Aluminum manganous sulphate, $\text{Al}_2(\text{SO}_4)_3,$
 $\text{MnSO}_4 + 25\text{H}_2\text{O}$.
 Sol. in H_2O . (Berzelius.)
 $+24\text{H}_2\text{O}$. Min. *Apjohnite*.

Aluminum manganic sulphate, $2\text{Al}_2(\text{SO}_4)_3,$
 $\text{Mn}_2(\text{SO}_4)_3$.
 Insol. in H_2O . (Étard, C. R. 36. 1399.)

Aluminum nickel sulphate, $\text{Al}_2(\text{SO}_4)_3, 2\text{NiSO}_4,$
 H_2SO_4 .
 Insol. in H_2O , but gradually decomp.
 thereby. (Étard, C. R. 87. 602.)

Aluminum potassium sulphate, basic,
 $3(\text{Al}_2\text{O}_3, \text{SO}_3), \text{K}_2\text{SO}_4 + 6\text{H}_2\text{O} = \text{K}_2\text{SO}_4,$
 $3\text{Al}_2(\text{SO}_4)_3(\text{OH})_4$.
 Min. *Alunite*. Insol. in H_2O . Insol. in
 conc. $\text{HCl} + \text{Aq}$.

Sol. in boiling H_2SO_4 of 1.845 sp. gr., but
 more easily in a mixture of 12 g. H_2SO_4 and
 1.5 g. H_2O , and also in weaker acids, if heated
 to 210° . (Mitscherlich, J. pr. 81. 108.)
 $+9\text{H}_2\text{O}$. Min. *Löwigite*. Sl. sol. in boil-
 ing $\text{HCl} + \text{Aq}$. (Mitscherlich, J. pr. 83. 455.)
 Nearly insol. in HCl or conc. $\text{HNO}_3 + \text{Aq}$,
 but sol. in a mixture of 1 pt. H_2SO_4 and 1 pt.
 H_2O . (Debray, Bull. Soc. (2) 7. 9.)
 $\text{Al}_2\text{O}(\text{SO}_4)_3, \text{K}_2\text{SO}_4$. Sol. in H_2O , but de-
 comp. by heating.

With varying composition. Precipitates.
 Insol. in H_2O . Very sl. sol. in cold, gradually
 in hot acids. (Bley, J. pr. 39. 17.) Very
 difficultly sol. in warm conc. $\text{HCl} + \text{Aq}$, but
 easily sol. in $\text{KOH} + \text{Aq}$. (Naumann, B. 8.
 1630.)

Aluminum potassium sulphate (Potash alum),
 $\text{KAl}(\text{SO}_4)_2 + 12\text{H}_2\text{O}$ or $\text{K}_2\text{Al}_2(\text{SO}_4)_4 =$
 $\text{K}_2\text{SO}_4, \text{Al}_2(\text{SO}_4)_3 + 24\text{H}_2\text{O}$.

Sol. in H_2O with absorption of heat.
 When 100 pts. H_2O at 10.8° are mixed with
 14 pts. alum, the temp. is lowered 1.4° .
 (Rüdorff, B. 2. 68.)

Burnt alum is very slowly sol. in H_2O .
 100 pts. H_2O at t° dissolve P pts. $\text{K}_2\text{Al}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

t°	P	t°	P
12 5	7 6	50 0	46 7
21 25	10 4	62 5	230 0
25 0	22 0	75 0	920 0
37 5	41 1	87 5	1566 0

(Brand's, 1822.)
 Sol. in 18 pts. cold, and 1.6 pts. boiling H_2O (Four-
 croy); in 14.12 pts. cold, and 0.75 pt. boiling H_2O
 (Bergmann); in 15 pts. cold, and 0.75 pt. boiling H_2O
 (Dumas); in 11.7 pts. H_2O at 18.75° (Abl).
 100 pts. H_2O dissolve 14.79 pts. alum at 15.56° , and
 133.33 pts. at 100° . (Ure's Dict.)
 $\text{K}_2\text{Al}_2(\text{SO}_4)_4 + \text{Aq}$ sat. at 15° contains 10.939 pts. alum
 in every 100 pts. H_2O . (Michel and Krafft.)
 $\text{K}_2\text{Al}_2(\text{SO}_4)_4 + \text{Aq}$ sat. in cold contains 5.2% alum
 (Fourcroy). 6.7% (Boerhave).

100 pts. H_2O dissolve (a) pts. anhydrous
 alum, and (b) pts. crystallised at t° .

	0°	10°	20°	30°	40°	50°
a	2.62	4.50	6.57	9.05	12.35	15.9
b	5.22	9.16	13.66	19.29	27.3	36.5

	60°	70°	80°	90°	100°
a	21.1	26.95	35.2	50.3	70.83
b	51.3	71.97	103.1	187.8	421.9

(Poggiale, A. ch. (3) 8. 467.)

According to Locke (Am. Ch. J. 1901, 2.
 174), Poggiale's tables for NH_4 and K alum
 are evidently transposed, and the above
 data are applied by Poggiale to NH_4 alum.

100 pts. H_2O dissolve $\text{K}_2\text{Al}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$
 corresponding to pts. anhydrous
 $\text{K}_2\text{Al}_2(\text{SO}_4)_4$.

Temp.	Pts. $\text{K}_2\text{Al}_2(\text{SO}_4)_4$	Temp.	Pts. $\text{K}_2\text{Al}_2(\text{SO}_4)_4$
0	3.0	60	25
5	3.5	70	40
10	4.0	80	71
15	5.0	90	109
20	5.9	92.5	119 5
30	7.9	100	154
40	11.7	110	200
50	17.0	111.9	210.6

(Mulder, Scheik. Verhandel. 1864. 90.)
 100 pts. H_2O at 17° dissolve 13.5 pts.
 $\text{K}_2\text{Al}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$, or 7.36 pts. $\text{K}_2\text{Al}_2(\text{SO}_4)_4$.
 (Redtenbacher, J. pr. 94. 442.)

Forms supersaturated solutions very easily.
 Supersat. solutions are brought to crystalli-
 zation by addition of a crystal of alum or an
 isomorphous substance, as chrome or iron
 alum. Other substances as NaCl , etc. have
 no action. (Thomson, Chem. Soc. 36. 199.)
 1 l. H_2O dissolves 72.3 g. anhydrous, or
 138.4 g. hydrated salt, or 0.28 g. mol. of
 anhydrous salt at 25° . (Locke, Am. Ch.
 J. 1901, 26. 175.)

Solubility in H_2O at t° .
 (g. alum in 1000 g. H_2O .)

t°	g. alum	t°	g. alum
0	57.0	75	1280 9
5	76.3	76	1412 1
10	84.9	77	1517 9
15	103.6	78	1680 1
20	120.3	79	1775 2
25	131.3	80	1950 0
30	184.9	82	2273 5
36	204.3	84	2661 5
40	250.0	84.6	2816.0
45	290.2	85.1	3166.6
50	367.8	85.3	3337.2
55	457.7	85.6	3372 2
60	585.4	86	3997 8
65	708.4	87	4825.4
70	943.8	88	6639 6

(Marino, Gazz. ch. it. 1905, 36. (2) 351.)

Solubility in H ₂ O at t°.			
t°	g. K ₂ Al ₂ (SO ₄) ₄ per 100 g. H ₂ O	g. K ₂ Al ₂ (SO ₄) ₄ + 24H ₂ O per 100 g. H ₂ O	g. mol. K ₂ Al ₂ (SO ₄) ₄ per 100 g. H ₂ O
0	3.0	5.65	0.0058
5	3.5	6.62	0.0068
10	4.0	7.60	0.0077
15	5.0	9.59	0.0097
20	5.9	11.40	0.0114
25	7.23	14.14	0.0140
30	8.39	16.58	0.0162
40	11.70	23.83	0.0227
50	17.00	36.40	0.0329
60	24.75	57.35	0.0479
70	40.00	110.5	0.0774
80	71.0	321.3	0.01374
90	109.0	2,275.0	0.2110
92.5	119.0	∞	0.2318

(Mulder, Poggiale, Locke; Marino, Gazz. ch. it. 1905, **35**, (2) 351; and Berkeley, Proc. Roy. Soc. 1904, **203**, A, 214, calc. by Seidell, Solubilities, 1st Ed.)

M.-pt. of K₂Al₂(SO₄)₄ + 24H₂O = 84.5° (Tilden, Chem. Soc. **45**, 409.); = 92.5° (Erdmann); = 91° (Locke).

Sp. gr. of sat. K₂Al₂(SO₄)₄ + Aq at 8° = 1.045 (Anthon); at 15° = 1.0488 (Michel and Krafft); at 15° = 1.0456 (Stolba).

Sp. gr. of K₂Al₂(SO₄)₄ + Aq at 15° containing 5% K₂Al₂(SO₄)₄ = 1.0477. (Kohlrausch, W. Ann. **1879**, 1.)

Sp. gr. of K₂Al₂(SO₄)₄ + Aq at 15°. a = pts. K₂Al₂(SO₄)₄ + 24H₂O in 100 pts. solution; b = pts. K₂Al₂(SO₄)₄ in 100 pts. solution; c = pts. K₂Al₂(SO₄)₄ for 100 pts. H₂O.

a	b	c	Sp. gr.
4	2.1792	2.2277	1.0210
8	4.3584	4.5570	1.0420
12	6.5376	6.9950	1.0641
13	7.083	7.622	1.0690

(Gerlach, Z. anal. **27**, 280.)

Saturated solution boils at 111.9°, and contains 210.6 pts. K₂Al₂(SO₄)₄ + 24H₂O to 100 pts. H₂O. (Mulder.)

100 pts. H₂O contain 52 pts. K₂Al₂(SO₄)₄, and boils at 104.5°. (Griffiths.) Crust forms at 106.3°, when the solution contains 114.2 pts. K₂Al₂(SO₄)₄ to 100 pts. H₂O. (Gerlach, Z. anal. **26**, 426.)

B.-pt. of K₂Al₂(SO₄)₄ + Aq containing pts. K₂Al₂(SO₄)₄ to 100 pts. H₂O.

B.-pt.	Pts. K ₂ Al ₂ (SO ₄) ₄	B.-pt.	Pts. K ₂ Al ₂ (SO ₄) ₄
100.5°	17.0	104.0°	83.9
101.0	30.2	104.5	90.7
101.5	41.8	105.0	97.6
102.0	51.6	105.5	103.9
102.5	60.4	106.0	110.5
103.0	68.7	106.5	116.9
103.5	76.7	106.7	120.55

(Gerlach, Z. anal. **26**, 435.)

K₂Al₂(SO₄)₄ + Al₂(SO₄)₃.
K₂Al₂(SO₄)₄ is nearly insol. in sat. Al₂(SO₄)₃ + Aq. (Crum, A. **89**, 156.)

Solubility in Al₂(SO₄)₃ + Aq. Solid Phase = K alum + Al₂(SO₄)₃.

t°	g. Al ₂ (SO ₄) ₃ + 18H ₂ O in 1000 g. H ₂ O	g. K ₂ SO ₄ in 1000 g. H ₂ O
0	234.73	23.45
20	824.25	30.85
35	911.02	35.29
50	1,243.21	59.55
65	1,598.00	119.43
77	1,872.11	183.80

(Marino, Gazz. ch. it. 1905, **35**, (2) 351.)

Solubility is decreased by presence of Na alum. (Venable, C. N. **1879**, **40**, 198.)

Nearly completed pptd. from sat. aq. solution by addition of Fe or Cr alum. (v. Hauer, J. B. **1866**, 59.)

K₂Al₂(SO₄)₄ + MgSO₄.
K₂Al₂(SO₄)₄ + Aq sat. at 10°, and then sat. with MgSO₄ at 9°, contains for 100 pts. H₂O—

	At 10°		At 9°
Alum (anhydrous)	4.0	2.7	
MgSO ₄		31.2	31.1
		33.9	

(Mulder)

K₂Al₂(SO₄)₄ + K₂SO₄.
K₂Al₂(SO₄)₄ + Aq at 10°, and then sat. with K₂SO₄ at same temp., contains for 100 pts. H₂O—

	At 10°		At 9°
Alum (anhydrous)	4.0	0.86	
K ₂ SO ₄		9.16	9.7
		10.20	

(Mulder)

Solubility in $K_2SO_4 + Aq.$ Solid phase = $K \text{ alum} + K_2SO_4.$

t°	$Al_2(SO_4)_3 + 18H_2O$ in 1000 g. H_2O	g. K_2SO_4 in 1000 g. H_2O	t°	$Al_2(SO_4)_3 + 18H_2O$ in 1000 g. H_2O	g. K_2SO_4 in 1000 g. H_2O
0.	5.06	75.83	40	73.88	163.10
0.5	8.658	75.18	50	126.00	195.40
5.	16.07	85.78	60	249.70	238.80
10	18.52	96.50	70	529.01	323.74
15	20.56	109.30	80	1,044.04	517.27
30	39.60	147.80

(Marino, l. c.)

$K_2Al_2(SO_4)_4 + Na_2SO_4.$
 $K_2Al_2(SO_4)_4 + Aq$ sat. at 10°, and then sat. with Na_2SO_4 at 9°, contains for 100 pts. H_2O —

	At 10°		At 9°
Alum (anhydrous)	4.0	4.1	...
Na_2SO_4	...	8.8	8.4
		12.9	

(Mulder.)

Solubility of $K_2Al_2(SO_4)_4 + Th_2Al_2(SO_4)_4$ in H_2O at 25°.

G. $K_2Al_2(SO_4)_4$ per l.	G. $Th_2Al_2(SO_4)_4$ per l.	Solid phase Mol. % $K_2Al_2(SO_4)_4$	Sp. gr.
69.90	0.00	100	1.0591
74.56	0.48	99.61	1.0601
67.90	1.72	98.48	1.0598
65.30	4.52	95.45	1.0603
64.95	9.60	91.73	1.0605
53.23	18.44	82.54	1.0609
45.32	24.60	75.12	1.0609
38.02	32.48	65.73	1.0611
34.54	35.59	61.36	1.0611
28.35	42.99	51.93	1.0623
10.94	66.12	21.34	1.0654
0.00	75.46	0.00	1.0674

(Foch, Z. Kryst. Min. 1897, 28. 397.)

Insol. in alcohol of 0.905 sp. gr. or less. (Anthon, J. pr. 14. 125.)

Insol. in acetone. (Naumann, B. 1904, 37. 4328.)

Insol. in methyl acetate. (Naumann, B. 1900, 42. 3790.)

Solubility in H_2O is increased by glycerine. (Dunlop, Pharm. J, 1910, 31. 6.)

Min. *Kahmir*.
 $+8H_2O$. Stable in dry air. (Marino, l. c.)
 $+14H_2O$. Converted into ord. alum in air. (Marino.)

Aluminum rubidium sulphate, $Al + 24H_2O.$

100 pts. H_2O dissolve 2.27 p very sol. in hot H_2O . (Redts pr. 94. 442.)

Solubility in 100 pts. H_2O at t° for salt dried at 130°

t°	Pts. alum	t°	Pts. alum
0	0.71	25	1.85
10	1.09	35	2.67
17	1.42	50	4.98

(Setterberg, A. 211. 10)

Solubility in H_2O .

t°	Pts. per litre
25	18 1
30	21.9
35	26.6
40	32 2

(Locke, Am. Ch. J. 1901,

Melts in crystal H_2O at Chem. Soc. 45. 409); at 105 at 109° (Locke.)

Aluminum silver sulphate, $24H_2O.$

Decomp. by H_2O . (Churcote, C. N. 9. 155.)

Aluminum sodium sulphate, $A 24H_2O.$

Very sl. efflorescent.

Sol. in 2.14 pts. H_2O at 13°, or 100 45.7 pts. soda alum. Sol. in 1 pt. boiling Schw. J. 36. 183.)
100 pts. H_2O dissolve 110 pts. at 1 liquid of 1.295 sp. gr. (Ure.)

100 pts. H_2O dissolve 51 pts. 16°. (Augé, C. R. 110. 1139.)

100 pts. H_2O dissolve 110 p at 0°. (Tilden, Chem. Soc. 45.

100 g. H_2O dissolve at:
10° 15° 20° 25° 30°
36.7 38.7 40.9 43.145.8 g. at

(Smith, J. Am. Chem. Soc. 11 M-pt. of $Na_2Al_2(SO_4)_4 + 2$

(Tilden, Chem. Soc. 45. 409.); = Am. Ch. J. 1901, 26. 183.)

Insol. in absolute alcohol. Min. *Mendocite*.

allous sulphate, $\text{TiAl}(\text{SO}_4)_3$.

ols. of anhydrous salt are sol. 25°; or 1 l. H_2O dissolves 75 g. of anhydrous, or 117.8 g. of the hydrated (Locke, Am. Ch. J. 1901, 28.)

solubility in H_2O at t°.

G. $\text{Al}_2\text{Ti}_2(\text{SO}_4)_4$ in 100 g. H_2O	G. $\text{Al}_2\text{Ti}_2(\text{SO}_4)_4$ + $24\text{H}_2\text{O}$ in 100 g. H_2O
3.15	4.84
3.80	5.86
4.60	7.12
6.40	10.00
7.60	11.95
9.38	14.89
14.40	23.57
22.50	38.41
35.36	65.19

Solubilities, 1st Ed., p. 15.)

$\text{Ti}_2\text{SO}_4 + 96\text{H}_2\text{O}$. Sol. in H_2O .

zinc sulphate, $\text{Al}_2(\text{SO}_4)_3$, $\text{ZnSO}_4 + 6\text{H}_2\text{O}$. (Kane.)

Ammonium sulphate chromium chloride, $(\text{SO}_4)_2\text{CrCl}_2(\text{OH})_2 + 2\text{H}_2\text{O}$. (S. 1906, 89. 337.)

Ammonium sulphate sodium fluoride. Sol. in H_2O . (Weber, Dingl. 263.)

Ammonium sulphate, $(\text{NH}_4)_2\text{SO}_4$. Sol. in H_2O with absorption of heat. $(\text{NH}_4)_2\text{SO}_4$ mixed with 100 pts. of water the temperature from 13.2° to 3.4° (Rüdorff, B. 2. 68.)

Ammonium sulphate in H_2O at 19°. (Schiff, A 109. 326.)
 H_2O at 18.75°. (Abl.)
 H_2O at 15.6°, and in 1 pt. boiling H_2O .

At 62.5° dissolve 78 pts. $(\text{NH}_4)_2\text{SO}_4$.
At 15° dissolve 60.739 pts. $(\text{NH}_4)_2\text{SO}_4$. (S. 1906, 89. 337.)

pts. cold H_2O . (Vogel, N. Rep. 1.)
pts. cold H_2O at 10°. (Mulder, 1.)
pts. H_2O at 16–17°. (v. Hauer, 2. 221.)

100 pts. H_2O dissolve at:

0°	10°	20°	30°
71.00	73.65	76.30	78.95
40°	50°	60°	70°
81.60	84.25	86.90	89.55
80°	90°	100°	
92.20	94.85	97.50	

(Alluard, C. R. 59. 500.)

Solubility in 100 pts. H_2O at t°.

t°	Pts. $(\text{NH}_4)_2\text{SO}_4$	t°	Pts. $(\text{NH}_4)_2\text{SO}_4$	t°	Pts. $(\text{NH}_4)_2\text{SO}_4$
0	70.6	37	80.1	74	93.1
1	70.9	38	80.4	75	93.4
2	71.1	39	80.7	76	93.8
3	71.4	40	81.0	77	94.2
4	71.6	41	81.3	78	94.5
5	71.8	42	81.7	79	94.9
6	72.1	43	82.0	80	95.3
7	72.3	44	82.3	81	95.6
8	72.5	45	82.7	82	96.0
9	72.8	46	83.0	83	96.4
10	73.0	47	83.3	84	96.8
11	73.2	48	83.7	85	97.2
12	73.5	49	84.0	86	97.6
13	73.7	50	84.4	87	98.0
14	74.0	51	84.7	88	98.4
15	74.2	52	85.1	89	98.8
16	74.4	53	85.5	90	99.2
17	74.7	54	85.8	91	99.6
18	74.9	55	86.2	92	100.0
19	75.1	56	86.6	93	100.4
20	75.4	57	86.9	94	100.8
21	75.7	58	87.3	95	101.2
22	75.9	59	87.7	96	101.6
23	76.2	60	88.0	97	102.1
24	76.4	61	88.4	98	102.5
25	76.7	62	88.7	99	102.9
26	76.9	63	89.1	100	103.3
27	77.2	64	89.5	101	103.8
28	77.5	65	89.9	102	104.2
29	77.8	66	90.2	103	104.6
30	78.0	67	90.6	104	105.1
31	78.3	68	90.9	105	105.5
32	78.6	69	91.3	106	106.0
33	78.9	70	91.6	107	106.5
34	79.2	71	92.0	108	107.0
35	79.5	72	92.4	108.9	107.5
36	79.8	73	92.7

(Mulder, calculated from his own and other observations, Scheik. Verhandel. 1864. 60.)

100 g. $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ contain 41.4 g. $(\text{NH}_4)_2\text{SO}_4$ at 0°. (de Waal, Dissert. 1910); 44.27 g. at 30° (Schreinemakers, Z. phys. Ch. 71. 110); 47.81 g. at 70°. (de Waal.)
 $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ sat. at 15° has sp. gr. 1.248. (Michel and Kraft, A. ch. (3) 41. 471.)

Sp. gr. of (NH ₄) ₂ SO ₄ +Aq at 15°.					
% (NH ₄) ₂ SO ₄	Sp. gr.	% (NH ₄) ₂ SO ₄	Sp. gr.	% (NH ₄) ₂ SO ₄	Sp. gr.
1	1.0057	18	1.1035	35	1.2004
2	1.0115	19	1.1092	36	1.2060
3	1.0172	20	1.1149	37	1.2116
4	1.0230	21	1.1207	38	1.2172
5	1.0287	22	1.1265	39	1.2228
6	1.0345	23	1.1323	40	1.2284
7	1.0403	24	1.1381	41	1.2343
8	1.0460	25	1.1439	42	1.2402
9	1.0518	26	1.1496	43	1.2462
10	1.0575	27	1.1554	44	1.2522
11	1.0632	28	1.1612	45	1.2583
12	1.0690	29	1.1670	46	1.2644
13	1.0747	30	1.1724	47	1.2705
14	1.0805	31	1.1780	48	1.2766
15	1.0862	32	1.1836	49	1.2828
16	1.0920	33	1.1892	50	1.2890
17	1.0977	34	1.1948

(Schiff, calculated by Gerlach, Z. anal. 8. 280.)

Sp. gr. of (NH ₄) ₂ SO ₄ +Aq at 15°.					
% (NH ₄) ₂ SO ₄	Sp. gr.	% (NH ₄) ₂ SO ₄	Sp. gr.	% (NH ₄) ₂ SO ₄	Sp. gr.
5	1.0292	20	1.1160	31	1.1787
10	1.0581	30	1.1730		

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of (NH ₄) ₂ SO ₄ +Aq at 15°.					
% (NH ₄) ₂ SO ₄	Sp. gr.	% (NH ₄) ₂ SO ₄	Sp. gr.	% (NH ₄) ₂ SO ₄	Sp. gr.
3	1.0181	10	1.0600	30	1.1773
6	1.0359	20	1.1190	40	1.2352

(Gerlach, Z. anal. 28. 493.)

Sp. gr. of sat. solution = 1.248. (Gerlach.)

Sp. gr. of $(\text{NH}_4)_2\text{SO}_4 + \text{Aq.}$	
$\frac{1}{2}(\text{NH}_4)_2\text{SO}_4$ g. mol. in 1000 g. of solution	Sp. gr. 16°. 16'
0	1.000000
0.5514	1.000347
1.1251	1.000704
2.3114	1.001436
4.5840	1.002823
10.0893	1.006093
20.0138	1.012023
40.5236	1.024117
56.8536	1.033690

(Dijken, Z. phys. Ch. 1897, 24. 107.)

Sp. gr. of $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ at 20°.		
Normality of $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$	% $(\text{NH}_4)_2\text{SO}_4$	Sp. gr.
3.75	40.28	1.2289
2.964	32.99	1.1858
1.978	23.01	1.1319
0.876	10.88	1.0626
0.492	6.275	1.0352

(Forchheimer, Z. phys. Ch. 1900, 24. 22.)

Sp. gr. of (NH ₄) ₂ SO ₄ +Aq at 20°.			
Solution	Sp. gr.	weight of 10 ccm. of the solution	% (NH ₄) ₂ SO ₄
sat.	1.248	12.5062 g.	53.2
¾ "	1.196	11.9034 "	39.9
½ "	1.139	11.3377 "	26.6
¼ "	1.077	10.7232 "	13.3
⅛ "	1.039	...	6.65

(Wiener, Z. phys. Chem. 1911, 71. 120.)

B.-pt. of sat. solution: crust formed at 106.2°, solution containing 88.2 pts (NH₄)₂SO₄ to 100 pts. H₂O; highest temp. observed, 108.2°. (Gerlach, Z. anal. 28. 426.)

B.-pt. of (NH ₄) ₂ SO ₄ +Aq containing pts (NH ₄) ₂ SO ₄ to 100 pts. H ₂ O.			
B. pt.	Pts. (NH ₄) ₂ SO ₄	B. pt.	Pts. (NH ₄) ₂ SO ₄
100.5°	7.8	105.0°	71.8
101.0	15.4	105.5	78.7
101.5	22.8	106.0	85.5
102.0	30.1	106.5	92.3
102.5	37.2	107.0	99.1
103.0	44.2	107.5	106.9
103.5	51.1	108.0	112.6
104.0	58.0	108.2	115.3
104.5	64.9

(Gerlach, Z. anal. 28. 431.)

Sol. with decomp. in HCl+Aq.

Solubility in H ₂ SO ₄ +Aq at 25°.		
100 g. of the solution contain		Solid phase
Mol. H ₂ SO ₄	Mol. (NH ₄) ₂ SO ₄	
0.00	2.28	(NH ₄) ₂ SO ₄
0.24	3.25	
0.47	3.19	
0.97	3.15	
1.19	3.15	
1.43	3.22	(NH ₄) ₂ H(SO ₄) ₂
1.72	3.18	
2.20	3.02	
2.60	2.97	
2.71	3.00	
2.82	3.03	
2.96	3.10	
3.20	3.19	
3.32	3.25	
3.47	3.32	
3.54	3.20	(NH ₄)HSO ₄
3.76	2.84	
4.22	2.28	
5.09	1.44	

(D'Ans, Z. anorg. 1909, 65. 229.)

Composition of the solution			Solid phase
% wt. H ₂ SO ₄	% by wt. (NH ₄) ₂ SO ₄	% by wt. H ₂ O	
63	43.59	45.79	(NH ₄) ₂ SO ₄
70	43.25	46.05	"
18	44.10	42.72	(NH ₄) ₂ SO ₄ + 3(NH ₄) ₂ SO ₄ , H ₂ SO ₄
67	42.06	41.27	3(NH ₄) ₂ SO ₄ , H ₂ SO ₄
82	41.15	33.03	"
33	41.16	31.51	"
32	44.63	22.26	"
12	45.50	21.83	"
34	45.52	20.64	(NH ₄)HSO ₄
36	45.31	20.73	"
51	35.37	26.12	"
12	30.10	27.78	"
30	24.88	29.32	"
77	24.30	29.93	"
55	16.98	26.37	"
43	20.41	17.16	"
46	24.40	13.14	"
12	24.20	12.68	"
57	27.67	9.76	"
33	29.75	8.42	"
56	30.26	7.28	"
57	31.86	5.47	"
59	33.70	3.71	"
43	36.75	1.72	"
43	36.95	0.82	"

Van Dorp, Z. phys. Ch. 1910, 73. 285.)

Solubility of (NH ₄) ₂ SO ₄ in H ₂ SO ₄ +Aq at 25°.		
In 1000 g. of the solution		Solid phase
Mol. SO ₃	Mol. (NH ₄) ₂ SO ₄	
4.29	2.17	(NH ₄)HSO ₄
4.57	1.83	
4.85	1.60	
5.25	1.36	
5.66	1.22	
6.16	1.26	(NH ₄)HSO ₄
6.47	1.55	
6.51	1.95	
6.50	2.37	(NH ₄)HSO ₄ +(NH ₄)H ₂ (SO ₄) ₂
6.43	2.50	
7.28	1.64	(NH ₄)H ₂ (SO ₄) ₂
7.99	1.38	
(7.60)	(1.74)	(NH ₄)HS ₂ O ₇
(8.00)	(1.42)	
9.02	0.96	
9.21	0.832	(NH ₄)HS ₂ O ₇
9.60	0.977	
9.68	1.00	(NH ₄)HS ₂ O ₇ +?
9.67	1.26	
10.43	0.894	

(D'Ans, Z. anorg. 1913, 80. 241.)

Very easily sol., even in conc. NH₄OH + Aq. (Girard, Bull. Soc. (2) 43. 522.)

Solubility of (NH ₄) ₂ SO ₄ in NH ₄ OH + Aq at 25°.		
In 1000 g. of the solution		
Mol. (NH ₄) ₂ SO ₄	Mol. (NH ₃) ₂	Solid phase
3.28	0	
2.60	1.02	(NH ₄) ₂ SO ₄
2.13	1.95	
1.59	3.44	
1.16	5.35	
0.78	7.13	
0	9.47	(NH ₄) ₂ SO ₄ + NH ₄ Cl

(D'Ans and Schreiner, Z. anorg. 1910, 67. 438.)

100 pts. H₂O dissolve 46.5 pts. (NH₄)₂SO₄ and 26.8 pts. NH₄Cl at 21.5°.

Solubility of (NH ₄) ₂ SO ₄ in NH ₄ Cl + Aq at 30°.		
% NH ₄ Cl	% (NH ₄) ₂ SO ₄	Solid phase
0	44	
6.86	36.15	(NH ₄) ₂ SO ₄
14.62	28.6	"
17.60	25.69	(NH ₄) ₂ SO ₄ + NH ₄ Cl
17.93	25.81	"
19.07	23.22	NH ₄ Cl
19.97	21.3	"
22.3	16.33	"
24.06	12.72	"
29.5	0	"

(Schreinemakers, Arch. Néer. Sc. 1910, (2) 15. 92.)

See also under NH₄Cl.

Solubility of $(\text{NH}_4)_2\text{SO}_4 + \text{CuSO}_4$ in H_2O at 16° .

Solution	% CuSO_4	% $(\text{NH}_4)_2\text{SO}_4$
Both salts in excess	8.55	7.12
15 cc. sat. sol. + 3 g. $(\text{NH}_4)_2\text{SO}_4$	1.77	18.16
15 cc. sat. sol. + 3 g. $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$	15.85	5.65

(Rüdorff, B. 6. 482.)

Solubility of $(\text{NH}_4)_2\text{SO}_4 + \text{CuSO}_4$ in H_2O at 30° .

$\%(\text{NH}_4)_2\text{SO}_4$	$\% \text{CuSO}_4$	Solid phase
0	20.32	$\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$
2.45	20.19	"
5.79	20.53	$\text{CuSO}_4 \cdot 5\text{H}_2\text{O} + \text{CuSO}_4 \cdot (\text{NH}_4)_2\text{SO}_4 \cdot 6\text{H}_2\text{O}$
6.98	16.77	$\text{CuSO}_4 \cdot (\text{NH}_4)_2\text{SO}_4 \cdot 6\text{H}_2\text{O}$
8.19	13.65	"
9.33	11.03	"
17.53	4.05	"
29.27	1.57	"
38.32	0.77	"
43.29	0.49	$\text{CuSO}_4 \cdot (\text{NH}_4)_2\text{SO}_4 \cdot 6\text{H}_2\text{O} + (\text{NH}_4)_2\text{SO}_4$
44	0	$(\text{NH}_4)_2\text{SO}_4$

(Schreinemakers, Arch. Néer. Sc. 1910, 15.
12.)

See also under CuSO_4 .

$$(\text{NH}_4)_2\text{SO}_4 + \text{FeSO}_4.$$

Solubility of $(\text{NH}_4)_2\text{SO}_4 + \text{FeSO}_4$ in $\text{H}_2\text{O} + \text{Aq}$ at 30° .

Compositon of the solution		Solid phase
% by wt. FeSO ₄	% by wt. (NH ₄) ₂ SO ₄	
24.90	0	FeSO ₄ · 7H ₂ O
25.24	5.24	"
25.22	5.93	FeSO ₄ · 7H ₂ O + FeSO ₄ · (NH ₄) ₂ SO ₄ · 6H ₂ O
25.26	5.89	"
23.59	6.44	FeSO ₄ · (NH ₄) ₂ SO ₄ · 6H ₂ O
17.64	8.90	"
13.13	11.45	"
7.95	16.29	"
5.70	19.64	"
1.72	34.24	"
0.79	43.86	FeSO ₄ · (NH ₄) ₂ SO ₄ · 6H ₂ O + (NH ₄) ₂ SO ₄
0.79	43.90	"
0	44.27	(NH ₄) ₂ SO ₄

(Schreinemakers, Z. phys. Ch. 1910, **71**. 111.)

$$(\text{NH}_4)_2\text{SO}_4 + \text{Li}_2\text{SO}_4.$$

Solubility of $(\text{NH}_4)_2\text{SO}_4 + \text{Li}_2\text{SO}_4$

Temp. = 30°.

$(\text{NH}_4)_2\text{SO}_4$ %	Li_2SO_4 %	Solid phase
44.1	0	$(\text{NH}_4)_2\text{SO}_4$
40.8	3	"
39.5	6.6	$(\text{NH}_4)_2\text{SO}_4 + \text{NH}_4\text{LiSO}_4$
30	10	NH_4LiSO_4
21.6	15	"
15	20	"
12.5	21.9	$\text{NH}_4\text{LiSO}_4 + \text{Li}_2\text{SO}_4$
8.9	23	$\text{Li}_2\text{SO}_4, \text{H}_2\text{O}$
0	25.1	"

Temp. = 50°.

45.7	1	$(\text{NH}_4)_2\text{SO}_4$
43.05	5.86	$(\text{NH}_4)_2\text{SO}_4 + \text{NH}_4\text{I}$
19.65	16.35	NH_4LiSO_4
13.90	21.20	"
13.97	21.23	$\text{NH}_4\text{LiSO}_4 + \text{Li}_2\text{SO}_4$
11.45	21.75	$\text{Li}_2\text{SO}_4, \text{H}_2\text{O}$
9.63	22.79	"
8.58	23.09	"
7.56	23.86	"
0	24.3	"

(Schreinemakers and Cocheret, *Ch*
Weekbl. 1905, 2. 771.)

$$(\text{NH}_4)_2\text{SO}_4 + \text{MnSO}_4.$$

Solubility of $(\text{NH}_4)_2\text{SO}_4 + \text{MnSO}_4$ in 1
G. per 100 g. sat. solution.

Temp. = 30°.

MnSO_4	$(\text{NH}_4)_2\text{SO}_4$	Solid phase
39.3	0	$\text{MnSO}_4 \cdot 5\text{H}_2\text{O}$
38.49	3.64	$\text{MnSO}_4 \cdot 5\text{H}_2\text{O} + \text{Mn}$
		$(\text{NH}_4)_2\text{SO}_4 \cdot 6\text{H}_2\text{O}$
33.44	4.91	"
22.06	9.65	"
9.02	20.36	"
2.91	37.42	"
1.75	42.58	$\text{MnSO}_4 \cdot (\text{NH}_4)_2\text{SO}_4 \cdot$
		$+ (\text{NH}_4)_2\text{SO}_4$
1.77	43.24	$(\text{NH}_4)_2\text{SO}_4$
0	43.4	"

Temp. = 50°.

36.26	0	$\text{MnSO}_4, \text{H}_2\text{O}$
35.35	2.95	$\text{MnSO}_4, \text{H}_2\text{O} + 2\text{Mn}^{++}$
		$(\text{NH}_4)_2\text{SO}_4$
30.57	5.14	$2\text{MnSO}_4, (\text{NH}_4)_2$
16.86	17.62	"
6.92	35.98	"
6.29	39.71	"
5.70	43.24	$2\text{MnSO}_4, (\text{NH}_4)_2\text{SO}_4$
		$(\text{NH}_4)_2\text{SO}_4$
3.49	44.02	$(\text{NH}_4)_2\text{SO}_4$
0	45.7	"

(Schreinemakers, Chem. Weekbl. 19
131.)

(NH₄)₂SO₄+K₂SO₄.
100 pts. (NH₄)₂SO₄+K₂SO₄+Aq sat. at 16.17° contain 38.41 pts. of the two salts, of which 5.45 pts. are K₂SO₄, and 32.96 pts. (NH₄)₂SO₄. (v. Hauer, J. pr. 28. 137.)
100 pts. H₂O dissolve 50.6 pts. (NH₄)₂SO₄ and 7.2 pts. K₂SO₄ at 11°. (Mulder, J. B. 336. 67.)
(NH₄)₂SO₄ and K₂SO₄ replace each other in solution, so that by adding one of these salts to a seemingly saturated solution of the other, it is dissolved with pptn. of the other salt. (Rüdorff, B. 6. 485.)

Solubility of (NH₄)₂SO₄+K₂SO₄ at 19.1°.

Solution	%K ₂ SO ₄	%(NH ₄) ₂ SO ₄
Both salts in excess	39.3	37.97
1 cc. sat. sol.+4 g. K ₂ SO ₄	4.94	33.26
1 cc. sat. sol.+4 g. (NH ₄) ₂ SO ₄	2.05	40.80

(Rüdorff, B. 6. 482.)

Solubility of (NH₄)₂SO₄+K₂SO₄ in H₂O at 25°.

g. K ₂ SO ₄ per l.	g. (NH ₄) ₂ SO ₄ per l.	g. K ₂ SO ₄ per l.	g. (NH ₄) ₂ SO ₄ per l.
127.9	0.0	59.28	355.0
135.7	115.7	40.27	482.7
84.2	281.1	0.0	542.3

Results are also given for 14°, 15°, 16°, 30°, 38° and 47°.

(Fock, Z. Kryst. Min. 1897, 28. 365.)

(NH₄)₂SO₄+Th(SO₄)₂.

Solubility of (NH₄)₂SO₄+Th(SO₄)₂ at 16°. Pts. per 100 pts. H₂O.

(NH ₄) ₂ SO ₄	Th(SO ₄) ₂	Solid phase
2.13	3.361	Th(SO ₄) ₂ .9H ₂ O
4.80	5.269	"
10.02	8.947	"
16.56	13.330	Th(SO ₄) ₂ .9H ₂ O+Th(SO ₄) ₂ .(NH ₄) ₂ SO ₄ .4H ₂ O
28.00	10.359	Th(SO ₄) ₂ .(NH ₄) ₂ SO ₄ .4H ₂ O
45.20	9.821	" +Th(SO ₄) ₂ .2(NH ₄) ₂ SO ₄ .2H ₂ O
55.14	6.592	Th(SO ₄) ₂ .2(NH ₄) ₂ SO ₄ .2H ₂ O
69.05	5.750	"
82.88	4.583	Th(SO ₄) ₂ .3(NH ₄) ₂ SO ₄ .3H ₂ O
95.74	1.653	"

(Barre, A. ch. 1911, (8) 24. 239.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 826.)

Insol. in absolute alcohol. Sol. in 500 pts alcohol of 0.872 sp. gr., and in 62.5 pts. of 0.905 sp. gr. (Anthon, J. pr. 14. 125.)

Sol. in 217.4 pts. of 66.8% alcohol (sp. gr. = 0.88) at 24.3°. (Pohl, J. pr. 56. 219.)

Tolerably sol. in alcohol, the sp. gr. of which is greater than 0.860. Insol. in alcohol of sp. gr. less than 0.850.

Solubility in dil. alcohol.

When (NH₄)₂SO₄ is dissolved in dil. alcohol, two layers are formed, the compositions of which are as follows:

Sp. gr.	Lower layer 100 ccm. contain in g.		
	alcohol	water	salt
1.2240	...	71.43	74.16
1.1775	8.85	68.26	59.54
1.1661	10.62	67.70	56.56
1.1655	11.29	67.34	56.30
1.1735	11.42	66.54	59.20

Sp. gr.	Upper layer 100 ccm. containing.		
	alcohol	water	salt
0.9530	41.37	48.47	5.45
0.9512	44.20	45.95	4.97
0.9440	44.27	45.61	4.51
0.9098	52.64	36.78	1.56
0.8750	62.61	24.60	0.30
0.8549	67.04	18.36	0.09
0.8308	77.55	5.53	0.00

(Bodländer, Z. phys. Ch. 7. 3, 8.)

Solubility in ethyl alcohol+Aq.

Upper layer			
Temp.	g. H ₂ O	g. alcohol	g. salt
16.6°	52.80	40.21	6.99
33.0°	47.99	46.75	5.26
41.8°	47.34	47.67	4.99
55.7°	45.90	49.47	4.63

Lower layer			
16.6°	60.33	10.19	29.48
33.0°	61.02	9.80	29.18
41.8°	61.16	9.74	29.10
55.7°	61.59	9.46	28.95

(Traube, Z. phys. Ch. 1887, 1. 509.)

Solubility of $(\text{NH}_4)_2\text{SO}_4$ in alcohol at 30° .

Two liquid layers are formed between alcohol concentrations of 5.8 and 62%.

Composition of layers.

Upper layer			Lower layer		
% $(\text{NH}_4)_2\text{SO}_4$	% alcohol	% H_2O	% $(\text{NH}_4)_2\text{SO}_4$	% alcohol	% H_2O
2.2	56.6	41.2	37.1	5.8	57.1
2.6	54.5	42.9	35.7	6.3	58
3.4	52.3	44.3	33.8	7.4	58.8
13.2	31.8	55	21.7	18.4	59.9
17	25	—	17	25	58

At concentration of 62% alcohol, the liquid is homogeneous and contains 1.3% $(\text{NH}_4)_2\text{SO}_4$.

(Wibaut, Chem. Weekbl. 1909, 6. 401.)

Solubility of $(\text{NH}_4)_2\text{SO}_4$ in alcohol at 60° .

% $(\text{NH}_4)_2\text{SO}_4$	% alcohol	% H_2O
43.02	2.32	54.66
41.1	4.1	54.8
1.2	64.5	34.3
0.2	75.5	24.3

Between 4.1° and 64.5° , two liquid layers are formed.

Composition of layers.

Upper layer			Lower layer		
% $(\text{NH}_4)_2\text{SO}_4$	% alcohol	% H_2O	% $(\text{NH}_4)_2\text{SO}_4$	% alcohol	% H_2O
1.2	64.5	34.3	41.1	4.1	54.8
1.6	60	38.4	36.8	6	57.2
3.8	50	46.2	30.8	9	60.2
7.4	40	52.6	26.6	12	61.4
10	34.1	55.6	23.6	15	61.4

(Schreinemakers, Z. phys. Ch. 1907, 59. 641.)

Solubility in alcohol + Aq at 0°

% $(\text{NH}_4)_2\text{SO}_4$	% alcohol	% H_2O
41.4	0	58.6
30.0	9.41	60.59
0.11	73.03	26.83

Two layers are formed between alcohol concentrations of 9.41 and 73.03%.

(de Waal, Dissert. 1910.)

Solubility in propyl alcohol + Aq at 20°

% propyl alcohol	% $(\text{NH}_4)_2\text{SO}_4$
20	6.7
30	4.8
40	3.2
50	2.0
60	1.0
70	0.4

(Lanebarger, Am. Ch. J. 1892, 14. 38)

100 g. 95% formic acid dissolve 25.4 g. $(\text{NH}_4)_2\text{SO}_4$ at 16.5° . (Aschan, Ch. Ztg. 1912, 37. 1117.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Ammonium hydrogen sulphate, NH_4HSO_4 .

Sl. deliquescent. Sol. in 1 pt. cold H_2O . (Link.)

Very sl. sol. in alcohol. (Gerhardt, A. ch. (3) 20. 255.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

$(\text{NH}_4)_2\text{H}(\text{SO}_4)_3$. Not deliquescent. Sol. in H_2O . (Mitscherlich, Pogg. 39. 194.)

$(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{H}_2\text{SO}_4$. (D'Ans and Schreiner, Z. anorg. 1913, 80. 241.)

Ammonium pyrosulphate, $(\text{NH}_4)_2\text{S}_2\text{O}_7$.

Decomp. by H_2O . (Schulze.)

$\text{NH}_4\text{HS}_2\text{O}_7$. (D'Ans and Schreiner, Z. anorg. 1913, 80. 241.)

Ammonium octosulphate, $(\text{NH}_4)_8\text{S}_8\text{O}_{52}$.

Decomp. by H_2O . (Weber, B. 17. 497.)

Ammonium antimony sulphate,

$(\text{NH}_4)_2\text{SO}_4 \cdot \text{Sb}_2(\text{SO}_4)_3$.

Behaves toward H_2O and abs. alcohol as a mixture of the components. (Metz, Z. anorg. 1905, 48. 152.)

Decomp. very slowly by H_2O . (Germann, Arch. Pharm. 1898, 236. 479.)

Ammonium bismuth sulphate, $\text{NH}_4\text{Bi}(\text{SO}_4)_3 \cdot 4\text{H}_2\text{O}$.

Easily sol. in HCl , and HNO_3 + Aq, less sol. in conc. H_2SO_4 , and hot dil. acids. Slowly decomp. by cold $\text{HC}_2\text{H}_3\text{O}_2$, and dil. H_2SO_4 + Aq. (Lüddecke, A. 140. 277.)

Ammonium cadmium sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{CdSO}_4 \cdot 6\text{H}_2\text{O}$.

Can be recrystallized from a little H_2O . (v. Hauer.)

1 l. H_2O dissolves 723 g. anhydrous salt at 25° . (Locke, Am. Ch. J. 1902, 27. 439.)

SO_4 , $\text{CdSO}_4 + 10\text{H}_2\text{O}$. (André, C. 37.)

m calcium sulphate,

$(\text{NH}_4)_2\text{Ca}(\text{SO}_4)_2 + \text{H}_2\text{O}$.

o. by H_2O . (Fassbender, B. 11.)

$(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Rose, Pogg.

uble salt is stable between 0° and solutions containing an excess of CaSO_4 . It is not formed if the solution less than 35% $(\text{NH}_4)_2\text{SO}_4$. (Barre, 9, 143. 1605.)

$\text{Ca}_2(\text{NH}_4)_2$. Decomp. by H_2O . 1907, 40. 192.)

uble salt is formed in the presence ss of CaSO_4 and at temp. above 80° . R. 1909, 143. 1605.)

$(\text{NH}_4)_2(\text{SO}_4)_2 + \text{H}_2\text{O}$. Decomp. by H_2O . 1907, 40. 192.)

m calcium cupric sulphate,

$(\text{NH}_4)_2(\text{SO}_4)_4 + 2\text{H}_2\text{O}$.

able. (D'Ans, B. 1908, 41. 1778.)

m calcium potassium sulphate,

$\text{CaK}(\text{SO}_4)_2 + \text{H}_2\text{O}$.

o. by cold H_2O . (Fassbender, B.

m cerous sulphate, $(\text{NH}_4)_2\text{Ce}_2(\text{SO}_4)_4$.

l. in cold than in hot H_2O . (Czud-

H_2O dissolve at:

35° 35.1° 45.2°

128 5.184 4.993 g. anhydrous salt,

55.3° 55.2°

2.240 2.187 g. anhydrous salt,

85.2°

2 1.184 g. anhydrous salt.

Wolff, Z. anorg. 1905, 45. 102.)

SO_4 , $\text{Ce}_2(\text{SO}_4)_3$. (Barre, C. R. 873.)

m ceric sulphate, $3(\text{NH}_4)_2\text{SO}_4$.

efflorescent. Easily sol. in H_2O . eff, A. 168. 50.)

SO_4 , $2\text{Ce}(\text{SO}_4)_2 + 3\text{H}_2\text{O}$. Sl. sol. (Mendelejeff.)

m chromous sulphate, $\text{NH}_4\text{Cr}(\text{SO}_4)_2$.

mol. anhydrous salt is sol. in 1 l. 5° . (Locke, Am. Ch. J. 1901, 26.

100 ccm. of sat. aqueous solution 5 g. of the salt at 20° . Insol. in (Laurent, C. R. 1911, 131. 112.)

Ammonium chromic sulphate, $(\text{NH}_4)_2\text{SO}_4$, $\text{Cr}_2(\text{SO}_4)_3$.

Not attacked by boiling H_2O or conc. $\text{HCl} + \text{Aq}$. Very slowly attacked by boiling $\text{KOH} + \text{Aq}$ (sp. gr. = 1.3). Insol. in $\text{CrCl}_2 + \text{Aq}$ or $\text{SnCl}_2 + \text{Aq}$. (Klobb, Bull. Soc. (3) 9. 664.)

$+5\text{H}_2\text{O}$. Is ammonium chromosulphate, which see.

$+24\text{H}_2\text{O}$. **Chrome Alum.**

Violet modification. Efflorescent. Sol. in cold H_2O , but solution is decomp. on heating with formation of green modification. The dil. solution of green modification is gradually converted into violet modification by standing. Alcohol ppts. it from aqueous solution. (Schrötter, Pogg. 53. 526.)

100 cc. H_2O dissolve 10.78 g. anhydrous, or 21.21 g. hydrated salt at 25° . Melts in crystal H_2O at 45° . (Locke, Am. Ch. J. 1901. 26. 174.)

Solubility in H_2O .

Saturation is very slowly reached owing to transition between violet and green modifications. If time of saturation is taken at $2\frac{1}{2}$ hours, 100 g. of the solution contain at:

0°	30°	40°
3.77	10.6	15.5 g. $(\text{NH}_4)_2\text{Cr}_2(\text{SO}_4)_4$.

This is assumed to be the solubility of the violet modification.

In 300 hours, 15.96 g. salt are dissolved at 30° , and 24.64 g. in 250 hours at 40° . (Koppel, B. 1906, 39. 3741.)

Calc. from electrical conductivity measurements, a solution containing 3.8 g. of the sulphate in 100 g. contains 48% of the green compound at 40° and 61% at 55° . With solutions of 6-7 times the above concentration equilibrium is reached at 40° with 30-40% green alum. (Koppel.)

Sp. gr. of aqueous solution of violet modification at 15° , containing:

4	8	12%
1.020	1.0405	1.0610

Sat. solution at 15° has sp. gr. = 1.070. (Gerlach.)

Green modification. Sol. in H_2O and alcohol. When in aqueous solution, it gradually changes to violet modification.

Sp. gr. of aqueous solution of green modification at 15° , containing:

10	20	30%
1.044	1.091	1.142

40	50	60%
1.197	1.255	1.317

70	80	90%
1.384	1.456	1.532

(Gerlach, Z. anal. 28. 498.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

$3(\text{NH}_4)_2\text{SO}_4$, $\text{Cr}_2(\text{SO}_4)_3$. Only sl. attacked by boiling H_2O . Not attacked by boiling conc. $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Klobb, Bull. Soc. (3) 9. 663.)

Ammonium cobaltous sulphate,
 $(\text{NH}_4)_2\text{Co}(\text{SO}_4)_2 + 6\text{H}_2\text{O}$.
100 pts. H_2O dissolve at:
0° 10° 18° 23° 35°
8.9 11.6 15.2 17.1 19.6 pts. anhydrous salt,

40° 45° 50° 60° 75°
22.3 25 28.7 34.5 43.3 pts. anhydrous salt.
(Tobler, A. 95. 193.)

100 pts. saturated solution contain at:
20° 40° 60° 80°
14.9 20.8 25.6 33 pts. anhydrous salt.
(v. Hauer, J. pr. 74. 433.)

1 l. H_2O dissolves 147.2 g. anhydrous salt
at 25°. Tobler's results are inaccurate.
(Locke, Am. Ch. J. 1902, 27. 459.)
Pptd. from aqueous solution by alcohol.

Ammonium cobaltic sulphate,
 $(\text{NH}_4)_2\text{Co}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.
Sol. in H_2O with decomp. (Marshall,
Chem. Soc. 59. 760.)

Ammonium cobaltous cupric sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CoSO}_4, \text{CuSO}_4 + 12\text{H}_2\text{O}$.
Quite easily sol. in hot H_2O , but on long
boiling a basic salt is pptd. (Vohl, A. 94. 58.)

Ammonium cobaltous ferrous sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CoSO}_4, \text{FeSO}_4 + 12\text{H}_2\text{O}$.
Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium cobaltous magnesium sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CoSO}_4, \text{MgSO}_4 + 12\text{H}_2\text{O}$.
Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium cobaltous manganous sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CoSO}_4, \text{MnSO}_4 + 12\text{H}_2\text{O}$.
Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium cobaltous nickel sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CoSO}_4, \text{NiSO}_4 + 12\text{H}_2\text{O}$.
Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium cobaltous zinc sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CoSO}_4, \text{ZnSO}_4 + 12\text{H}_2\text{O}$.
Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium cupric sulphate, $(\text{NH}_4)_2\text{SCu}_2$,
 $\text{CuSO}_4 + 6\text{H}_2\text{O}$.
Efflorescent in warm air.
Sol. in 1.5 pts. boiling H_2O , and separates almost
wholly on cooling. (Vogel, J. pr. 2. 194
Sol. in 1.55 pts. H_2O at 18.75°. (Abl.)

100 pts. H_2O at 19° dissolve 26.6 pts., and
sat. solution has sp. gr. = 1.1337. (Schiff, A.
109. 426.
100 g. sat. solution at 30° contain 30.36 g.
anhydrous salt. (Schreinemakers, Arch. Néer.
Sci. 1910, (2) 15. 92.)

Solubility of $(\text{NH}_4)_2\text{Cu}(\text{SO}_4)_2, 6\text{H}_2\text{O} +$
 $(\text{NH}_4)_2\text{Ni}(\text{SO}_4)_2, 6\text{H}_2\text{O}$ in H_2O at 13-14°.
Mols. per 100 mols. H_2O .

Cu salt	Ni salt	% Cu salt in solid phase
0	0.521	0
0.1476	0.295	10.29
0.2664	0.2089	30.59
0.4165	0.1449	52.23
0.4785	0.1202	78.80
1.0350	0	100

(Fock, Z. Kryst. Min. 1897, 28. 365.)

Solubility of $(\text{NH}_4)_2\text{Cu}(\text{SO}_4)_2, 6\text{H}_2\text{O} +$
 $\text{K}_2\text{Cu}(\text{SO}_4)_2, 6\text{H}_2\text{O}$ in H_2O at 13-14°.
Mols. per 100 mols. H_2O .

K salt	NH_4 salt	% K salt in solid phase
0	1.035	0
0.897	0.8618	5.06
0.2269	0.6490	16.78
0.2570	0.5887	30.40
0.2946	0.5096	36.63
0.3339	0.3319	50.15
0.4560	0.1961	69.93
0.4374	0	100.

(Fock.).

Solubility of $(\text{NH}_4)_2\text{Cu}(\text{SO}_4)_2, 6\text{H}_2\text{O} +$
 $(\text{NH}_4)_2\text{Zn}(\text{SO}_4)_2, 6\text{H}_2\text{O}$ in H_2O at 13-14°.
Mols. per 100 mols. H_2O .

Cu salt	Zn salt	% Cu salt in solid phase
0.0422	0.8069	2.39
0.0666	0.5638	4.52
0.1218	0.5115	9.03
0.2130	0.4924	14.67
0.3216	0.4022	22.62
1.035	0	100.

(Fock.)

$(\text{NH}_4)_2\text{SO}_4, 2\text{CuSO}_4$. Very sol. in H_2O
(Klobb, C. R. 115. 230.)

Ammonium cupric ferrous sulphate.
Sol. in H_2O without decomposition. (Vohl
A. 94. 61.)

Ammonium cupric magnesium sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{CuSO}_4, \text{MgSO}_4 + 12\text{H}_2\text{O}$.
Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium cupric magnesium potassium sul-
phate, $(\text{NH}_4)_2\text{SO}_4, \text{CuSO}_4, \text{MgSO}_4, \text{K}_2\text{SO}_4 +$
 $12\text{H}_2\text{O}$.
Sol. in H_2O . (Schiff.)
 $2(\text{NH}_4)_2\text{SO}_4, \text{CuSO}_4, 2\text{MgSO}_4, \text{K}_2\text{SO}_4 +$
 $18\text{H}_2\text{O}$. Sol. in H_2O . (Schiff.)

cupric manganous sulphate,
 $\text{MnSO}_4 \cdot \text{CuSO}_4 \cdot 12\text{H}_2\text{O}$.
 O. (Vohl, A. 94. 57.)

cupric nickel sulphate,
 $\text{NiSO}_4 \cdot \text{CuSO}_4 \cdot 12\text{H}_2\text{O}$.
 O. (Vohl.)

cupric potassium sulphate,
 $\text{K}_2\text{SO}_4 \cdot \text{CuSO}_4 \cdot 6\text{H}_2\text{O}$.
 O. (Schiff.)

cupric zinc sulphate,
 $\text{ZnSO}_4 \cdot \text{CuSO}_4 \cdot 12\text{H}_2\text{O}$.
 O. (Vohl.)

cupric sulphate ammonia,
 $\text{SO}_4 \cdot \text{CuO} \cdot 2\text{NH}_3$.
 5 pts. cold H_2O , but decomp. on
 air or dilution Insol. in alcohol.

didymium sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{Dy}_2\text{O}_3 \cdot 8\text{H}_2\text{O}$.
 8 pts. H_2O , and less easily in
 Aq. (Marignac.)
 ly sol. in H_2O (Cleve, Bull. Soc.)

erbium sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{Er}_2\text{O}_3 \cdot 8\text{H}_2\text{O}$.
 O. (Cleve.)

gallium sulphate,
 $\text{Ga}_2(\text{SO}_4)_3 \cdot 24\text{H}_2\text{O}$.
 d water and dilute alcohol. Conc.
 uds up on boiling, but clears on
 il. solution separates out a basic
 hot or cold H_2O . (Boisbaudran.)

glucinum sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{Ge}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$.
 O. (Atterberg.)

indium sulphate,
 $\text{In}_2(\text{SO}_4)_3 \cdot 24\text{H}_2\text{O}$.
 H_2O dissolve 200 pts. salt at 16° ,
 at 30° .
 alcohol.
 crystal H_2O at 36° . (Rössler, J.)
 (Rössler.)

iridium sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{Ir}_2\text{O}_3 \cdot 24\text{H}_2\text{O}$.
 l. in H_2O . (Marino, Z. anorg.
 1.)

iron (ferrous) sulphate,
 $\text{Fe}(\text{SO}_4)_2 \cdot 6\text{H}_2\text{O}$.
 sol. in H_2O than $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$.
 r. 2. 192.)

100 pts. H_2O dissolve at:
 0° 12° 20° 30° 36°
 12.2 17.5 21.6 28.1 31.8 pts. anhydrous salt,
 45° 55° 60° 65° 75°
 36.2 40.3 44.6 49.8 56.7 pts. anhydrous salt.
 (Tobler, A. 95. 193.)

100 pts. H_2O at 16.5° dissolve 35.9 pts.
 hydrous salt.
 1 l. H_2O dissolves 351 pts. or 1.044 g. mol.
 anhydrous salt at 25° . (Locke, Am. Ch. J.
 1902, 27. 459.)

Sol. in H_2O without decomp. Aq. solution
 at 30° contains 13.13% FeSO_4 and 11.45%
 $(\text{NH}_4)_2\text{SO}_4$. (Schreinemakers, C. C. 1910, I.
 801.)

Sp. gr. of $(\text{NH}_4)_2\text{FeSO}_4 \cdot \text{Aq}$ at 19° .
 $\% = \%(\text{NH}_4)_2\text{FeSO}_4 \cdot 6\text{H}_2\text{O}$.

%	Sp. gr	%	Sp. gr	%	Sp. gr
1	1.006	11	1.066	21	1.130
2	1.013	12	1.073	22	1.136
3	1.018	13	1.080	23	1.143
4	1.024	14	1.085	24	1.150
5	1.030	15	1.092	25	1.156
6	1.036	16	1.097	26	1.164
7	1.042	17	1.104	27	1.171
8	1.047	18	1.110	28	1.179
9	1.054	19	1.116	29	1.185
10	1.060	20	1.124	30	1.193

(Schiff calculated by Gerlach, Z. anal. 8.
 280.)

Insol. in acetone.

Ammonium ferric sulphate, basic.

Extremely difficultly sol. in $\text{HCl} + \text{Aq}$. Not
 decomp. by $\text{KOH} + \text{Aq}$. (Berzelius.)
 $5(\text{NH}_4)_2\text{O} \cdot 3\text{Fe}_2\text{O}_3 \cdot 12\text{SO}_3 + 18\text{H}_2\text{O}$ or
 $2(\text{NH}_4)_2\text{O} \cdot \text{Fe}_2\text{O}_3 \cdot 4\text{SO}_3 + 4\text{H}_2\text{O}$. Sol. in 2.4
 pts. cold H_2O . (Maus, Pogg. 11. 79.)

Ammonium iron (ferric) sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{Fe}_2(\text{SO}_4)_3$.

Attacked slowly by cold H_2O . (Lachaud
 and Lepierre.)

Nearly insol. in H_2O . (Weinland, Z. anorg.
 1913, 84. 363.)

+ $24\text{H}_2\text{O}$. Iron alum. Sol. in 3 pts. H_2O
 at 15° . (Forchhammer, Ann. Phil. 5. 406.)

100 cc. H_2O dissolve 44.15 g. anhydrous, or
 124.40 g. hydrated salt at 25° , or 1.659 g.
 mols. anhydrous salt are sol. in 1 l. H_2O at
 25° . (Locke, Am. Ch. J. 1901, 28. 174.)

Sp. gr. of aqueous solution at 15° contain-
 ing:

5 10 15% $(\text{NH}_4)_2\text{Fe}_2(\text{SO}_4)_6 + 24\text{H}_2\text{O}$,
 1.023 1.047 1.071

20 25 30% $(\text{NH}_4)_2\text{Fe}_2(\text{SO}_4)_6 + 24\text{H}_2\text{O}$,
 1.096 1.122 1.148

35 40% $(\text{NH}_4)_2\text{Fe}_2(\text{SO}_4)_6 + 24\text{H}_2\text{O}$.
 1.175 1.203

40% solution is sat. at 15°. (Gerlach, Z. anal. 28. 496.)

Melts in crystal H_2O at 40°. (Locke.)
 $3(NH_4)_2SO_4, Fe_2(SO_4)_3$. Insol. in cold H_2O . (Lachaud and Lepierre.)

Ammonium iron (ferroferric) sulphate,
 $4(NH_4)_2SO_4, FeSO_4, Fe_2(SO_4)_3 + 3H_2O$.

Sl. sol. in cold H_2O ; decomp. into basic salt by hot H_2O ; insol. in alcohol. (Lachaud and Lepierre, C. R. 114. 916.)

Ammonium ferrous magnesium sulphate,
 $4(NH_4)_2SO_4, 3FeSO_4, MgSO_4 + 24H_2O$.

Sol. in H_2O . (Schiff, A. 107. 64.)
 $2(NH_4)_2SO_4, FeSO_4, MgSO_4 + 12H_2O$. Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium ferrous manganous sulphate,
 $2(NH_4)_2SO_4, FeSO_4, MnSO_4 + 12H_2O$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium ferrous nickel sulphate,
 $2(NH_4)_2SO_4, FeSO_4, NiSO_4 + 12H_2O$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium ferrous zinc sulphate,
 $2(NH_4)_2SO_4, FeSO_4, ZnSO_4 + 12H_2O$.

Sol. in H_2O . (Bette, A. 14. 286.)

Ammonium lanthanum sulphate, $(NH_4)_2SO_4, La_2(SO_4)_3 + 8H_2O$.

Sl. sol. in H_2O . (Marignac.)
 Quite sol. in H_2O . (Cleve.)
 $+ 2H_2O$. (Barre, C. R. 1910, 151. 872.)
 $5(NH_4)_2SO_4, 2La_2(SO_4)_3$. Sl. sol. in $(NH_4)_2SO_4 + Aq$ of concentrations above 60%. (Barre, A. ch. 1911, (8) 24. 246.)
 $5(NH_4)_2SO_4, La_2(SO_4)_3$. (Barre.)

Ammonium lead sulphate, $(NH_4)_2SO_4, PbSO_4$.
 Decomp. by H_2O into its constituents. (Wöhler and Litton, A. 43. 126.)

Decomp. by H_2O . Only stable in contact with solutions containing:

13.86 pts. $(NH_4)_2SO_4$ per 100 pts. H_2O at 20°.

19.25 pts. $(NH_4)_2SO_4$ per 100 pts. H_2O at 50°.

24.31 pts. $(NH_4)_2SO_4$ per 100 pts. H_2O at 75°.

29.42 pts. $(NH_4)_2SO_4$ per 100 pts. H_2O at 100°.

(Barre, C. R. 1909, 149. 294.)

Ammonium lithium sulphate, NH_4LiSO_4 .

Solubility in H_2O = 35.25% at -10°, and 36.18% at 70°. (Schreinemakers, C. C. 1906, I. 217.)

This is the only double salt which $(NH_4)_2SO_4$ forms with Li_2SO_4 below 100°. (Spielrein, C. R. 1913, 157. 48.)

Ammonium magnesium sulphate,
 $(NH_4)_2Mg(SO_4)_2 + 6H_2O$.

100 pts. H_2O dissolve 15.9 pts. anhydrous double salt at 13°. (Mulder.)

100 pts. H_2O dissolve at:

0°	10°	15°	20°	30°
9.0	14.2	15.7	17.9	19.1

pts. anhydrous s

45°	50°	55°	60°	75°
25.6	30.0	31.9	36.1	45.3

pts. anhydrous s
 (Tobler, A. 96. 193.)

More sol. in H_2O than $(NH_4)_2SO_4$ or $MgSO_4$ (Graham.)

1 l. H_2O dissolves 199 pts. anhydrous : at 25°. Tobler's results are inaccurate (Locke, Am. Ch. J. 1902, 27. 459.)

100 g. H_2O dissolve at:

34°	41° F.
18.22	20.72 g. $(NH_4)_2SO_4, MgSO_4, 6H_2O$

50°	59° F.
22.48	24.08 g. $(NH_4)_2SO_4, MgSO_4, 6H_2O$

60°	70° F.
24.81	28.26 g. $(NH_4)_2SO_4, MgSO_4, 6H_2O$

81° F.

33.33 g. $(NH_4)_2SO_4, MgSO_4, 6H_2O$.

(Lothian, Pharm. J. 1910, (4) 30. 546.)

Lothian's results for solubility in H_2O probably incorrect because of inaccuracy experimental method. (Seidell, Pharm. 1911, (4) 33. 846.)

Solubility of $(NH_4)_2Mg(SO_4)_2$ in H_2O at t

t°	g. anhydrous salt per 100 g.	
	solution	H_2O
0	10.58	11.83
10	12.75	14.61
20	15.23	17.96
25	16.45	19.69
30	17.84	21.71
40	20.51	25.86
50	23.18	30.17
60	26.02	35.17
80	32.58	46.32
100	39.66	65.72

(Porlezza, Att. Acc. Linc. 1914, (5) 21. 509.)

Min. *Cerbolite*.

Ammonium magnesium nickel sulphate,
 $2(NH_4)_2SO_4, MgSO_4, NiSO_4 + 12H_2O$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium magnesium potassium zinc sulphate, $2(NH_4)_2SO_4, 3MgSO_4, 3K_2SO_4, 2ZnSO_4 + 30H_2O$.

Sol. in H_2O . (Schiff, A. 107. 64.)

$(NH_4)_2SO_4, 2MgSO_4, 2K_2SO_4, ZnSO_4, 18H_2O$. Sol. in H_2O . (Schiff.)

$(NH_4)_2SO_4, MgSO_4, K_2SO_4, ZnSO_4 + 12H_2O$. Sol. in H_2O . (Schiff.)

Ammonium magnesium zinc sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{MgSO}_4, \text{ZnSO}_4 + 12\text{H}_2\text{O}$.
 Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium manganous sulphate, $(\text{NH}_4)_2\text{SO}_4, \text{MnSO}_4 + 6\text{H}_2\text{O}$.

Deliquescent. Easily sol. in H_2O . (Jahn.)
 1 l. H_2O dissolves 372 g. anhydrous salt at 25° . (Locke, Am. Ch. J. 1902, 27. 459.)
 $(\text{NH}_4)_2\text{SO}_4, 2\text{MnSO}_4$. Readily decomp. by H_2O . (Lepierre, C. R. 1895, 120. 924.)

Ammonium manganic sulphate, $(\text{NH}_4)_2\text{SO}_4, \text{Mn}_2(\text{SO}_4)_3$.

Decomp. by H_2O . Insol. in ether, C_6H_6 , and conc. H_2SO_4 . Sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Lepierre, Bull. Soc. 1895, (3) 13. 596.)
 $+24\text{H}_2\text{O}$. Decomp. by H_2O . (Mitscherlich.)

Ammonium manganous nickel sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{MnSO}_4, \text{NiSO}_4 + 12\text{H}_2\text{O}$.
 Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium manganous zinc sulphate,
 $2(\text{NH}_4)_2\text{SO}_4, \text{MnSO}_4, \text{ZnSO}_4 + 12\text{H}_2\text{O}$.
 Sol. in H_2O . (Vohl.)

Ammonium mercuric sulphate, $(\text{NH}_4)_2\text{SO}_4, 3\text{HgSO}_4 + 2\text{H}_2\text{O}$.

(Hirzel, J. B. 1850. 333.)
 $(\text{NH}_4)_2\text{SO}_4, \text{HgSO}_4$. Difficultly sol. in H_2O . Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$.

Ammonium mercurous sulphate ammonia,
 $3\text{Hg}_2\text{O}, 2(\text{NH}_4)\text{HgSO}_4, 2\text{NH}_3$.

Insol. in hot or cold, dil. or conc. H_2SO_4 and HNO_3 . Sol. in HCl . (Tarugi, Gazz. ch. it. 1903, 33. (1) 131.)

Ammonium nickel sulphate, $(\text{NH}_4)_2\text{SO}_4, \text{NiSO}_4 + 6\text{H}_2\text{O}$.

Sol. in 4 pts. cold H_2O . (Link, 1796.)

100 pts. H_2O dissolve at:

8.5°	10°	16°	20°	30°
1.8	3.2	5.8	5.9	8.3

pts. anhydrous salt,

40°	50°	59°	68°	85°
11.5	14.4	16.7	18.8	28.6

pts. anhydrous salt.
 (Tobler, A. 95. 193.)

100 pts. sat. solution contain at 20° , 9.4; at 40° , 13.2; at 60° , 18.6; at 80° , 23.1 pts. anhydrous salt. (v. Hauer, J. pr. 74. 433.)

1 l. H_2O dissolves 75.7 g. anhydrous salt at 25° . (Locke, Am. Ch. J. 1902, 27. 459.)

Nearly insol. in a weak acid solution of $(\text{NH}_4)_2\text{SO}_4$. (Thompson, C. C. 1863. 957.)

Ammonium nickel zinc sulphate, $2(\text{NH}_4)_2\text{SO}_4, \text{NiSO}_4, \text{ZnSO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Ammonium nickel sulphate ammonia,
 $(\text{NH}_4)_2\text{SO}_4, \text{NiSO}_4, 6\text{NH}_3 + 3\text{H}_2\text{O}$.
 (André, C. R. 106. 936.)

Ammonium platonic sulphate, $2(\text{NH}_4)_2\text{SO}_4, \text{Pt}_2(\text{SO}_4)_3 + 25\text{H}_2\text{O}$.

Sol. in H_2O . (Prost, Bull. Soc. (2) 46. 156.)

Ammonium potassium sulphate, $(\text{NH}_4)_2\text{SO}_4, \text{K}_2\text{SO}_4 + 4\text{H}_2\text{O}$.

Soluble in H_2O . 100 pts. H_2O at 16° dissolve 13.68 pts. salt. (Thomson, 1831.)
 Min. *Taylorite*.

Ammonium praseodymium sulphate,
 $(\text{NH}_4)_2\text{SO}_4, \text{Pr}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$.

Sl. sol. in H_2O . (von Scheele, Z. anorg. 1898, 18. 359.)

Ammonium rhodium sulphate,
 $(\text{NH}_4)_2\text{SO}_4, \text{Rh}_2(\text{SO}_4)_3 + 24\text{H}_2\text{O}$.

Very sol. H_2O ; melts in crystal H_2O at $102-103^\circ$. (Piccini, Z. anorg. 1901, 27. 67.)

Ammonium samarium sulphate, $(\text{NH}_4)_2\text{SO}_4, \text{Sm}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Cleve, Bull. Soc. (2) 43. 166.)

Ammonium scandium sulphate, $(\text{NH}_4)_2\text{SO}_4, \text{Sc}_2(\text{SO}_4)_3$.

Sol. in H_2O . (Cleve.)

Sol. in H_2O and in dil. $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (R. J. Meyer, Z. anorg. 1914, 86. 279.)

Ammonium sodium sulphate, $\text{NH}_4\text{NaSO}_4 + 2\text{H}_2\text{O}$.

100 pts. H_2O dissolve 46.6 pts. of cryst. salt at 15° , and the solution has a sp. gr. of 1.1749.

Sp. gr. of aqueous solution containing:

31.8	24.44	15.9%	$\text{NH}_4\text{NaSO}_4 + 2\text{H}_2\text{O}$
1.1749	1.1380	1.0849	
12.72	6.36	%	$\text{NH}_4\text{NaSO}_4 + 2\text{H}_2\text{O}$
1.0679	1.0337		

(Schiff, A. 114. 68.)

Ammonium strontium sulphate.

Insol. in excess of $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. (Rose, Pogg. 110. 296.)

$(\text{NH}_4)_2\text{SO}_4, \text{SrSO}_4$. This double salt is only stable in contact with nearly sat. solutions of $(\text{NH}_4)_2\text{SO}_4$. (Barre, C. R. 1909, 149. 293.)

Ammonium tellurium sulphate,
 $(\text{NH}_4)\text{HSO}_4, 2\text{TeO}_2, \text{SO}_3 + 2\text{H}_2\text{O}$.

As K salt. (Metzner, A. ch. 1898, (7) 15. 203.)

Ammonium thallic sulphate, $\text{NH}_4\text{Tl}(\text{SO}_4)_2$.

(Marshall, C. C. 1902, II. 1089.)

$+4\text{H}_2\text{O}$. Decomp. by H_2O . Easily sol. in dil. acids. (Fortini, Gazz. ch. it. 1905, 35. (2) 450.)

Ammonium thorium sulphate, $2(\text{NH}_4)_2\text{SO}_4 \cdot \text{Th}(\text{SO}_4)_2$.

Easily sol. in H_2O and sat. $(\text{NH}_4)_2\text{SO}_4 + \text{Aq.}$ (Cleve.)

+ $2\text{H}_2\text{O}$. (Barre.)

$(\text{NH}_4)_2\text{SO}_4 \cdot \text{Th}(\text{SO}_4)_2 + 4\text{H}_2\text{O}$. (Barre, A. ch. 1911, (8) 24. 240.)

$3(\text{NH}_4)_2\text{SO}_4 \cdot \text{Th}(\text{SO}_4)_2 + 3\text{H}_2\text{O}$. (B.)

Ammonium titanium sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{TiO} \cdot \text{SO}_4 + \text{H}_2\text{O}$.

Very sol. in H_2O with decomp.

Insol. in conc. H_2SO_4 . (Rosenheim, Z. anorg. 1901, 26. 252.)

$(\text{NH}_4)_2\text{O} \cdot 2\text{TiO}_2 \cdot 2\text{SO}_3 + 3\text{H}_2\text{O}$. Slowly decomp. by H_2O . (Blondel, Bull. Soc. 1899, (3) 21. 262.)

Ammonium titanium sesquisulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot 3\text{Ti}_2(\text{SO}_4)_3 + 18\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in HCl .

Insol. in H_2SO_4 . Decomp. by boiling with conc. H_2SO_4 . (Stähler, B. 1905, 38. 2623.)

Ammonium uranous sulphate, $2(\text{NH}_4)_2\text{SO}_4 \cdot \text{U}(\text{SO}_4)_2$.

Easily sol. in H_2O . (Rammelsberg.)

Sol. in H_2O but solution rapidly decomp. (Kohlschütter, B. 1901, 34. 3630.)

Ammonium uranyl sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot (\text{UO}_2)_2\text{SO}_4 + 2\text{H}_2\text{O}$.

Quite difficultly sol. in H_2O . (Arfvedson.)

Ammonium vanadous sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{VSO}_4 + 6\text{H}_2\text{O}$.

Decomp. in the air.

Sol. in H_2O . (Piccini, Z. anorg. 1899, 19. 205.)

Less sol. in H_2O than $\text{VSO}_4 + 7\text{H}_2\text{O}$. (Piccini and Marino, Z. anorg. 1902, 32. 60.)

Ammonium vanadic sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{V}_2(\text{SO}_4)_3 + 12\text{H}_2\text{O}$.

Insol. in H_2O .

Insol. in H_2SO_4 . Decomp. by boiling with conc. H_2SO_4 .

Sol. in HCl . (Stähler, B. 1905, 38. 3980.)

+ $24\text{H}_2\text{O}$. Very sol. in H_2O ; decomp. in the air. (Piccini, Z. anorg. 1896, 11. 108.)

100 pts. H_2O dissolve 39.76 pts. salt at 10° .

Sp. gr. of sat. solution at $4^\circ/20^\circ = 1.687$. (Piccini, Z. anorg. 1897, 13. 446.)

1 l. H_2O dissolves 31.69 g. anhydrous or 78.51 g. hydrated salt at 25° , or 1.210 g. mols. anhydrous salt are sol. in 1 l. H_2O at 25° .

Mpt. of crystals = 45° . (Locke, Am. Ch. J. 1901, 26. 175.)

Ammonium vanadyl sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{VOSO}_4 + 3\frac{1}{2}\text{H}_2\text{O}$.

Easily sol. in H_2O and in a mixture of alcohol and conc. H_2SO_4 , but cannot be recryst. therefrom. (Koppel and Behrendt, Z. anorg. 1903, 35. 176.)

$(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{VOSO}_4 + \text{H}_2\text{O}$. Deliquescent. Slowly but abundantly sol. in H_2O , but cannot be recryst. from it except by addition of H_2SO_4 . (Koppel and Behrendt, Z. anorg. 1903, 35. 172.)

Ammonium yttrium sulphate, $2(\text{NH}_4)_2\text{SO}_4 \cdot \text{Y}_2(\text{SO}_4)_3 + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve.)

Ammonium zinc sulphate, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{ZnSO}_4 + 6\text{H}_2\text{O}$.

100 pts. H_2O dissolve pts. $(\text{NH}_4)_2\text{SO}_4 \cdot \text{ZnSO}_4$ at:

0°	10°	13°	15°	20°
7.3	8.8	10.0	12.5	12.6 pts. salt,
30°	45°	60°	75°	85°
16.5	21.7	29.7	37.8	46.2 pts. salt.

(Tobler, A. 95. 193.)

1 l. H_2O dissolves 140.8 g. anhydrous salt at 25° . (Locke, Am. Ch. J. 1902, 27. 459.)

+ $7\text{H}_2\text{O}$. (André, C. R. 104. 987.)

Ammonium zirconium sulphate.

Sol. in cold or hot H_2O or in acids. (Berzelius.)

Ammonium sulphate antimony fluoride, $(\text{NH}_4)_2\text{SO}_4 \cdot 2\text{SbF}_3$.

Very sol. in H_2O . (Mayer, B. 1894, 27. R. 922.)

Ammonium sulphate chromic chloride, $2(\text{NH}_4)_2\text{SO}_4 \cdot \text{CrCl}_3 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Weinland, B. 1907, 40. 3765.)

Ammonium sulphate hydrogen peroxide, $(\text{NH}_4)_2\text{SO}_4 \cdot \text{H}_2\text{O}_2$.

Efflorescent in air.

Easily sol. in H_2O . (Willstätter, B. 1903, 36. 1829.)

Antimony sulphate basic, $7\text{Sb}_2\text{O}_3 \cdot 2\text{SO}_3 + 3\text{H}_2\text{O}$.

Insol. in, and not decomp. by hot or cold H_2O . (Adie, Chem. Soc. 57. 540.)

$5\text{Sb}_2\text{O}_3 \cdot 2\text{SO}_3 + 7\text{H}_2\text{O}$. Insol. in H_2O . (Hensgen, R. t. o. 4. 401.)

$2\text{Sb}_2\text{O}_3 \cdot \text{SO}_3 + x\text{H}_2\text{O}$. Not decomp. by cold H_2O . (Adie.)

$\text{Sb}_2\text{O}_3 \cdot \text{SO}_3 = (\text{SbO})_2\text{SO}_4$. Decomp. by hot H_2O . (Peligot, J. B. 1847. 426.)

+ H_2O . As above. (Adie.)

$\text{Sb}_2\text{O}_3 \cdot 2\text{SO}_3$, and + H_2O , and + $2\text{H}_2\text{O}$. Scarcely decomp. by cold, slowly by boiling H_2O . Slowly sol. in dil. $\text{HCl} + \text{Aq.}$ (Adie.)

Antimony sulphate, $\text{Sb}_2(\text{SO}_4)_3$.

Very deliquescent. Combines with H_2O to a hard mass with evolution of heat; with more H_2O it becomes liquid, and by repeated treat-

with much boiling H_2O it is wholly p. into H_2SO_4 and Sb_2O_3 . (Hensgen, . 4. 401.)

any sulphate, acid, Sb_2O_3 , $4SO_3$, comp. by H_2O . (Adie.)
 $Sb_2O_3 + 8$, or $9SO_3$. Decomp. by H_2O .

any barium sulphate, $Ba_2(SO_4)_3$, $BaSO_4 + 6H_2O$.
 Ca comp. (Kühl, Z. anorg. 1907, 54.)

any caesium sulphate, $SbCs(SO_4)_3$.
 ly decomp. by cold H_2O . (Gutman, Pharm. 1908, 246. 188.)

any calcium sulphate, $Ba_2(SO_4)_3$, $CaSO_4 + 6H_2O$.
 comp. by H_2O . (Kühl, Z. anorg. 1907, .)

any lithium sulphate, $SbLi(SO_4)_3$.
 comp. by H_2O . (Gutman, Arch. Pharm. 246. 187.)

any potassium sulphate, $KSb(SO_4)_3$.
 comp. by H_2O . (Gutmann, Arch. . 1898, 236. 478.)

any rubidium sulphate, $SbRb(SO_4)_3$.
 ly decomp. by cold H_2O . (Gutman.)

any silver sulphate, $SbAg(SO_4)_3$.
 ly decomp. by H_2O . (Kühl, Z. anorg. 54. 258.)
 ly decomp. by cold H_2O . (Gutman Pharm. 1908, 246. 189.)

any sodium sulphate, $NaSb(SO_4)_3$.
 ly decomp. by H_2O . (Gutmann, Pharm. 1898, 236. 478.)

any strontium sulphate, $Sb_2(SO_4)_3$, $SO_4 + 6H_2O$.
 Ca comp. (Kühl, Z. anorg. 1907, 54.)

any thallium sulphate, $SbTl(SO_4)_3$.
 ly decomp. by cold H_2O . Decomp. ic. H_2SO_4 . (Gutman, Arch. Pharm. 246. 189.)

c sulphate.
 Arsenic sulphur trioxide.

i sulphate, $BaSO_4$.

43,000 pts. H_2O (Kirwan); in 200,000 pts. H_2O ritte, C. R. 33. 308).
 ts. H_2O dissolve 0.002 pt. $BaSO_4$. (Ure's

$BaCl_2 + Aq$ containing 1 pt. BaO to 71,000 pts. H_2O , when treated with H_2SO_4 , becomes turbid in $\frac{1}{2}$ hour. (Harting, J. pr. 22. 52.)

$Ba(NO_3)_2 + Aq$ containing 1 pt. BaO to 25,000 pts. H_2O gives a distinct cloud with H_2SO_4 or $Na_2SO_4 + Aq$; with 50,000–100,000 pts. H_2O a slight turbidity is produced; with 200,000–400,000 pts. H_2O the mixture becomes turbid in a few minutes; while with 800,000 pts. H_2O no action is visible. (Lassaigne, J Chim. Méd. 8. 526.)

Sol. in 800,000 pts. H_2O (Calvert); in 400,000 pts. cold or hot H_2O (Fresenius).

Calculated from the electrical conductivity of the solution, $BaSO_4$ is sol. in 429,700 pts. H_2O at 18.4° , and 320,000 pts. at 37.7° . (Holleman, Z. phys. Ch. 12. 131.)

1 l. H_2O dissolves 1.72 mg. at 2° ; 1.97 mg. at 10° ; 2.29 mg. at 19.0° ; 2.60 mg. at 26° ; 2.91 mg. at 34° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

Calculated from the electrical conductivity of the solution, $BaSO_4$ is sol. in 425,000 pts. H_2O at 18.3° . Results of Fresenius and Hintz (Z. anal. 1896, 35. 170) are incorrect. (Küster, Z. anorg. 1896, 12. 267.)

Sat. aq. solution contains 2.29 mg. $BaSO_4$ per liter at 25° when particles of salt are not less than 1.8μ in diameter.

Sat. aq. solution contains 4.15 mg. $BaSO_4$ per liter at 25° when particles of salt are 0.1μ in diameter. ($\mu = 0.0001$ cm.) (Hulett, Z. phys. Ch. 1901, 37. 398–9.)

In general the influence of the size of the grain on the solubility of the substance is negligible when the solubility exceeds 2%. The increase of normal solubility by using finely divided solids, amounts to 80% in the case of $BaSO_4$. (Hulett, Z. phys. Ch. 1904, 47. 366.)

1 l. H_2O dissolves 2.3 mg. $BaSO_4$ at 18° . (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)

Calculated from electrical conductivity of $BaSO_4 + Aq$.

0.0190 milli-equivalents are sol. in 1 liter H_2O at 18° ; 0.0212 at 25° ; 0.0288 at 50° ; 0.0334 at 100° .

(Melcher, J. Am. Chem. Soc. 1910, 32. 55.)

Not attacked by cold HCl or $HNO_3 + Aq$ after several hours, and only in traces after several days. On boiling, traces of $BaSO_4$ dissolve, and the liquid after cooling can be precipitated by $BaCl_2$ or $H_2SO_4 + Aq$, but not by H_2O alone. (Rose, Pogg. 95. 108.)

By washing $BaSO_4$ long enough with H_2O containing HCl or HNO_3 [$HC_2H_3O_2$ (Siegle)], the filtrate can be precipitated by H_2SO_4 or $BaCl_2$. (Piria, J. B. 1856. 334.)

1000 pts. 3% $HCl + Aq$ dissolve 0.06 pt. $BaSO_4$ in the cold, and still more on boiling.

230 ccm. $HCl + Aq$ of 1.02 sp. gr. dissolve 0.048 g. $BaSO_4$ from 0.679 g. $BaSO_4$ when boiled $\frac{1}{4}$ hour.

168 ccm. $HCl + Aq$ of 1.03 sp. gr. dissolve 0.0075 g. $BaSO_4$ from 0.577 g. $BaSO_4$ when boiled 5 minutes. (Siegle, J. pr. 69. 142.)

Solubility of BaSO₄ in HCl + Aq.

No. cc. HCl + Aq containing 1 mg. equiv. HCl	Mg. BaSO ₄ per 1 mg. equiv. of HCl	g. per 100 cc. solution	
		HCl	BaSO ₄
2.0	0.133	1.82	0.0067
1.0	0.089	3.65	0.0089
0.5	0.056	7.29	0.0101
0.2	0.017	18.23	0.0086

(Banthisch, J. pr. 1884, 29. 54.)

100,000 pts. H₂O dissolve 0.124 pt. BaSO₄; 1000 pts. HNO₃ + Aq of 1.167 sp. gr. dissolve 2 pts. BaSO₄; 1000 pts. HNO₃ + Aq of 1.032 sp. gr. dissolve 0.062 pt. BaSO₄. (Calvert, Chem. Gaz. 1856. 55.)

When 0.4 g. BaSO₄ is heated ¼ hour with 150 ccm. HNO₃ + Aq of 1.02 sp. gr., 0.165 g. is dissolved. (Siegle, J. pr. 69. 142.)

Solubility of BaSO₄ in HNO₃ + Aq.

No. cc. HNO ₃ + Aq containing 1 mg. equiv. HNO ₃	Mg. BaSO ₄ per 1 mg. equiv. of BaSO ₄	G. per 100 cc. solution	
		HNO ₃	BaSO ₄
2.0	0.140	3.15	0.0070
1.0	0.107	6.31	0.0107
0.5	0.085	12.61	0.0170
0.2	0.048	31.52	0.0241

(Banthisch, J. pr. 1884, 29. 54.)

Acetic acid has less solvent power than other acids. 80 ccm. HC₂H₃O₂ + Aq of 1.02 sp. gr. boiled with 0.4 g. BaSO₄ ¼ hour dissolve 0.002 g. (Siegle, J. pr. 69. 142.)

Sol. in boiling conc. H₂SO₄. (See BaH₂(SO₄)₂.)

Sol. in fuming H₂SO₄. (See BaS₂O₇.)

Sol. in 2500 pts. boiling 40% HBr + Aq; in 6000 pts. boiling 40% HI + Aq. (Haslam, C. N. 53. 87.)

Sol. in considerable amount in metaphosphoric acid + Aq. (Scheerer and Drechsel, J. pr. (2) 7. 68.)

Not attacked by boiling conc. KOH + Aq if CO₂ is not present. (Rose, Pogg. 95. 104.)

Very sl. decomp. by standing a long time with cold conc. alkali carbonates + Aq.

Decomp. by boiling Na₂CO₃ or K₂CO₃ + Aq, not by (NH₄)₂CO₃ + Aq. (See Storer's Dict. for analytical data.)

Very sl. sol. in NH₄Cl + Aq, 1 pt. dissolving in 230,000 pts. sat. NH₄Cl + Aq.

500 ccm. sat. NH₄NO₃ + Aq with 50 ccm. sat. NH₄Cl + Aq dissolve 2 g. BaSO₄. 100 ccm. sat. NH₄NO₃ + Aq with 100 ccm. sat. NH₄Cl + Aq dissolve only 0.08 g. BaSO₄, therefore above solubility is due to free chlorine. (Mittentzwey, J. pr. 75. 214.)

BaSO₄ cannot be precipitated from solutions containing free Cl₂. (Erdmann, J. pr. 75. 215.)

Pptn. is retarded sl. by tartaric and racemic acids. (Spiller.)

Na metaphosphate prevents pptn. of BaSO₄, but not ortho- or pyrophosphate. (Scheerer, J. pr. 75. 114.)

Not precipitated in presence of alkali citrates. (Spiller.)

Much less sol. in NH₄Cl + Aq than in NH₄NO₃ + Aq. Insol. in warm conc. Na₂S₂O₃ + Aq. (Diehl, J. pr. 79. 431.)

Not appreciably sol. in H₂O containing ammonium or sodium chloride. (Brett, Wittstein, Wackenroder.)

Not appreciably sol. in H₂O at 250°, or in H₂O containing Na₂S. (Senarmont.)

Solubility is increased by alkali nitrates, but not appreciably by NaCl, KClO₃, or Ba(NO₃)₂. (Fresenius, Z. anal. 9. 52.) Scarcely sol. in boiling conc. (NH₄)₂SO₄ + Aq. (Fresenius.)

Solubility in H₂O increased by presence of MgCl₂. (Fresenius); cerium salts (Marignac).

Sol. in Fe₂Cl₆ + Aq. (Lunge, Z. anal. 19. 141.)

Solubility in various salts + Aq at 20-25°.

g. salt per l.	Mg. BaSO ₄ dissolved per l. is		
	FeCl ₃	AlCl ₃	MgCl ₂
1	58	33	30
2.5	72	43	30
5	115	60	33
10	123	94	34
25	150	116	50
50	160	170	50
100	170	175	50

(Fraps, Am. Ch. J. 1902, 27. 290.)

Solubility in sat. solution of various salts + Aq.

Salt	G. BaSO ₄ sol. in 1 l. of the solvent
NaNO ₃	0.2940
NaCl	0.00783
NH ₄ Cl	0.00627

(Ehlert, Z. Elektrochem. 1912, 18. 728)

Conc. CrCl₃ + Aq dissolves 40-120 times as much BaSO₄ as H₂O, when boiled therewith for 5 days; conc. CrCl₃ + Aq acidified with HCl, 450 times as much in 10 days. (Kuster, Z. anorg. 1905, 43. 348.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 827.)

Sol. in H₂O₂ + Aq. (Gawalowski, C. C. 1906, II. 7.)

100 cc. 95% formic acid dissolve 0.01 g. BaSO₄ at 18.5°. (Aschan, Ch. Ztg. 1913, 37. 1117.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); in ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Min. Barite.

Barium hydrogen sulphate, $\text{BaH}_2(\text{SO}_4)_2$.

100 pts. H_2SO_4 dissolve 2.22 pts. BaSO_4 . Lies-Bodart and Jacquemin, C. R. 46. 1206; .69 pts. BaSO_4 (Struve, Z. anal. 9. 34).

Boiling H_2SO_4 dissolves 10-12% freshly precipitated BaSO_4 without separating crystals on cooling. H_2SO_4 at 100° dissolves more than boiling H_2SO_4 , and becomes cloudy when heated to boiling. (Schultz, Pogg. 133. 46.)

1 g. BaSO_4 pptd. from BaCl_2 is sol. in 3153 . 91% H_2SO_4 ; from $\text{Ba}(\text{NO}_3)_2$ is sol. in 1519 . 91% H_2SO_4 . (Varenne and Pauleau, C. R. 9. 1016.)

100 pts. hot conc. H_2SO_4 dissolve approx. pts. BaSO_4 . (Rohland, Z. anorg. 1910, 66. 06.)

10 cc. of sat. BaSO_4 + absolute H_2SO_4 contain approx. 2.851 g. BaSO_4 . (Bergius, J. phys. Ch. 1910, 72. 355.)

Equilibrium in the system $\text{BaSO}_4 + \text{H}_2\text{SO}_4 + \text{H}_2\text{O}$ at 25° .

Composition of the solution

% H_2SO_4	g BaSO_4 per l.	Solid phase
73.83	0.030	BaSO_4
78.04	0.135	"
80.54	0.285	"
83.10	0.800	"
85.78	3.215	$\text{BaSO}_4, 2\text{H}_2\text{SO}_4, \text{H}_2\text{O}$
88.08	12.200	"
93.17	49.665	$\text{BaSO}_4, \text{H}_2\text{SO}_4$

Volkhonski, C. C. 1910, I. 1954; C. A. 1911. 617.)

Decomp. by H_2O , alcohol, or ether.

+ $2\text{H}_2\text{O}$. (Schultz.)

$\text{BaSO}_4, 2\text{H}_2\text{SO}_4 + \text{H}_2\text{O}$. (Volkhonski.)

Barium pyrosulphate, BaS_2O_7 .

100 pts. fuming H_2SO_4 dissolve 15.89 pts. BaSO_4 . (Struve, Z. anal. 9. 34.)

Very deliquescent.

Decomp. with H_2O with hissing. (Schultze-ellack, B. 4. 111.)

Barium calcium sulphate, $3\text{BaSO}_4, \text{CaSO}_4$.

Min. *Dreclite*.

Barium platonic sulphate (?)

Insol. in H_2O or boiling HCl or HNO_3 + Aq. Sol. in hot conc. H_2SO_4 or aqua regia. (E. Davy.)

Barium tin (stannic) sulphate, $\text{BaSO}_4, \text{Sn}(\text{SO}_4)_2 + 3\text{H}_2\text{O}$.

Decomp. by H_2O . Insol. in HCl . (Weinland and Kühl, Z. anorg. 1907, 54. 249.)

Barium titanium sulphate, $2\text{BaSO}_4, 3\text{Ti}(\text{SO}_4)_3$.

Ppt. Decomp. by H_2O , giving titanio acid. Weinland, Z. anorg. 1907, 54. 255.)

Barium sulphate potassium chloride, $3\text{BaSO}_4, \text{KCl}$.

Ppt. (Silberberger, M. 1904, 25. 233.)

Bismuth sulphate, basic, $(\text{BiO})_2\text{SO}_4$.

Insol. in H_2O . Sol. in HNO_3 or H_2SO_4 + Aq.

+ $2\text{H}_2\text{O}$. (Heintz, Pogg. 63. 55.)

$4\text{Bi}_2\text{O}_3, 3\text{SO}_3 + 15\text{H}_2\text{O}$. Insol. in H_2O . (Leist.)

$(\text{BiO})\text{HSO}_4 + \text{H}_2\text{O}$. Insol. in H_2O . Sol. in dil. H_2SO_4 + Aq.

+ $2\text{H}_2\text{O}$. Decomp. by H_2O with separation of $(\text{BiO})_2\text{SO}_4 + 2\text{H}_2\text{O}$. (Heintz.)

$3\text{Bi}_2\text{O}_3, 2\text{SO}_3 + 2\text{H}_2\text{O}$. Insol. in H_2O . (Athanasesco, C. R. 103. 271.)

$5\text{Bi}_2\text{O}_3, 11\text{SO}_3 + 17\text{H}_2\text{O}$. This sulphate crystallizes out from sulphuric acid of any strength between $\text{H}_2\text{SO}_4, 6\text{H}_2\text{O}$ and $\text{H}_2\text{SO}_4, 12\text{H}_2\text{O}$. (Adie, Proc. Chem. Soc. 1899, 15. 226.)

$\text{Bi}_2\text{O}_3, 2\text{SO}_3, 2\frac{1}{2}\text{H}_2\text{O}$ is in equilibrium at 50° with 5.4-51.4% H_2SO_4 + Aq.

$\text{Bi}_2\text{O}_3, \text{SO}_3$ is in equilibrium at 50° with acid solutions weaker than 5.4% H_2SO_4 .

(Allan, Am. Ch. J. 1902, 27. 287.)

Bismuth sulphate, $\text{Bi}_2(\text{SO}_4)_3$.

Very hygroscopic. Takes up H_2O with strong evolution of heat to form $2\text{Bi}_2(\text{SO}_4)_3 + 7\text{H}_2\text{O}$, which becomes $\text{Bi}_2(\text{SO}_4)_3 + 3\text{H}_2\text{O}$ at 100° . Decomp. by boiling H_2O into $\text{Bi}_2\text{O}_3, \text{SO}_3 + \text{H}_2\text{O}$. (Hensgen, J. B. 1886. 552.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Bismuth sulphate, acid, $\text{Bi}_2\text{O}_3, 4\text{SO}_3$.

$\text{Bi}_2\text{O}_3, 4\text{SO}_3$ is in equilibrium at 50° with 51.4-90% H_2SO_4 + Aq. (Allan, Am. Ch. J. 1902, 27. 287.)

+ H_2O . Crystallizes out from sulphuric acid at temp. above 170° . (Adie, Proc. Chem. Soc. 1899, 15. 226.)

+ $3\text{H}_2\text{O}$. Crystallizes from sulphuric acid of any strength between $\text{H}_2\text{SO}_4, \text{H}_2\text{O}$ and $\text{H}_2\text{SO}_4, 2\text{H}_2\text{O}$. (Adie.)

+ 7, or $9\text{H}_2\text{O} = \text{BiH}(\text{SO}_4)_2 + 3\text{H}_2\text{O}$. Insol. in H_2O . Easily sol. in acids, especially HCl , and HNO_3 + Aq. (Leist, A. 160. 29.)

+ $7\text{H}_2\text{O}$. Crystallizes out from sulphuric acid of any strength between $\text{H}_2\text{SO}_4, 3\text{H}_2\text{O}$ and $\text{H}_2\text{SO}_4, 5\text{H}_2\text{O}$.

+ $10\text{H}_2\text{O}$. Crystallizes out from sulphuric acid at temp. below 170° . (Adie.)

Bismuth potassium sulphate, $\text{Bi}_2(\text{SO}_4)_3, 3\text{K}_2\text{SO}_4$ (?)

Decomp. by H_2O ; insol. in sat. K_2SO_4 + Aq. (Heintz.)

$\text{Bi}_2(\text{SO}_4)_3, 2\text{K}_2\text{SO}_4$.

$\text{BiK}(\text{SO}_4)_2 = \text{Bi}_2(\text{SO}_4)_3, \text{K}_2\text{SO}_4$. Insol. in cold H_2O ; decomp. by boiling. (Brigham, Am. Ch. J. 14. 170.)

Bismuth sodium sulphate, $\text{Bi}_4\text{Na}_6(\text{SO}_4)_6$.
(Lüdecke, A. 140. 277.)

Boron sulphate.
See Borosulphuric acid.

Bromomolybdenum sulphate.
See under Bromomolybdenum compounds.

Cadmium sulphate, basic, 2CdO , SO_3 , and $+\text{H}_2\text{O}$.
Difficultly sol. in H_2O . (Stromeyer.) Sl. sol. in hot H_2O . (Habermann, M. 5. 432.)
 4CdO , SO_3 . (Pickering, Chem. Soc. 1907, 91. 1987.)

Cadmium sulphate, CdSO_4 .
Sat. $\text{CdSO}_4 + \text{Aq}$ contains at:
0° 10° 24° 30° 65°
35.9 37.5 41.5 42.0 49.7% CdSO_4 ,
86° 94° 130° 165° 188° 200°
43.5 91.6 27.7 14.7 7.1 2.3% CdSO_4 .
 CdSO_4 easily forms supersat. solutions. (Étard, A. ch. 1894, (7) 2. 552.)

Solubility in H_2O .

t°	Per cent CdSO_4 in sat. solution	Solid phase
-18	43.35	Ice
-10	43.27	"
0	43.01	$\text{CdSO}_4 \cdot \frac{1}{2}\text{H}_2\text{O}$
+10	43.18	"
15	43.20	"
20	43.37	"
30	43.75	"
40	43.99	"
60	44.99	"
62	45.06	"
72	46.2	"
73.5	46.6	"
74.5	46.7	"
75	46.5	$\text{CdSO}_4 \cdot \text{H}_2\text{O}$
77	42.2	"
78.5	41.5	"
85	39.6	"
90	38.7	"
95	38.1	"
100	37.8	"

(Mylius and Funk, B. 1897, 30. 825.)
See also under $\text{CdSO}_4 + 2\frac{1}{2}\text{H}_2\text{O}$, and $7\text{H}_2\text{O}$.

Sp. gr. at 0°/4° of $\text{CdSO}_4 + \text{Aq}$ containing 14.0 g. CdSO_4 in 1000 g. $\text{H}_2\text{O} = 1.0122$.
Sp. gr. at 12°/4° of $\text{CdSO}_4 + \text{Aq}$ containing 14.0 g. CdSO_4 in 100 g. $\text{H}_2\text{O} = 1.0121$.
Sp. gr. at 12°/4° of $\text{CdSO}_4 + \text{Aq}$ containing 57.2 g. CdSO_4 in 1000 g. $\text{H}_2\text{O} = 1.0514$.
Sp. gr. at 0°/4° of $\text{CdSO}_4 + \text{Aq}$ containing 183.1 g. CdSO_4 in 1000 g. $\text{H}_2\text{O} = 1.1552$.
Sp. gr. at 13°/4° of $\text{CdSO}_4 + \text{Aq}$ containing 183.1 g. CdSO_4 in 1000 g. $\text{H}_2\text{O} = 1.1529$.
(Fouqué, Ann. Observ. 1868, 9. 172.)

Sp. gr. of $\text{CdSO}_4 + \text{Aq}$ at 18°.

% CdSO_4	1	5	10	15
Sp. gr.	1.0084	1.0486	1.1026	1.1607

% CdSO_4 20 25 30 35
Sp. gr. 1.2245 1.295 1.3725 1.4575

% CdSO_4 36
Sp. gr. 1.4743
(Grotrian, W. Ann. 1883, 18. 193.)

Sp. gr. of $\text{CdSO}_4 + \text{Aq}$ at room temp. containing:

7.14	14.66	22.011% CdSO_4
1.0681	1.1591	1.2681

(Wagner, W. Ann. 1883, 18. 268.)

Sp. gr. of $\text{CdSO}_4 + \text{Aq}$ at 25°.

Concentration of $\text{CdSO}_4 + \text{Aq}$	Sp. gr.
1-normal	1.0973
$\frac{1}{5}$ "	1.0487
$\frac{1}{5}$ "	1.0244
$\frac{1}{5}$ "	1.0120

(Wagner, Z. phys. Ch. 1890, 5. 36.)

Sp. gr. of $\text{CdSO}_4 + \text{Aq}$.

% CdSO_4	t°	Sp. gr. at t°	Sp. gr. at 15°
0.0289	17.29	0.99908	0.99893
...	23.65	0.99776	...
0.0498	18.00	...	0.99915
0.0999	18.00	...	0.99961
0.495	18.00	...	1.0034
0.981	18.00	...	1.0084

(Wershofen, Z. phys. Ch. 1890, 5. 494.)

Sp. gr. at 16°/4° of $\text{CdSO}_4 + \text{Aq}$ containing 29.4654% $\text{CdSO}_4 = 1.36289$; containing 21.3671% $\text{CdSO}_4 = 1.24211$. (Schönrock, Z. phys. Ch. 1893, 11. 781.)

Sp. gr. of $\text{CdSO}_4 + \text{Aq}$ at 18°/4°.

% CdSO_4	25.121	18.172
Sp. gr.	1.297	1.200

% CdSO_4 9.952 5.639
Sp. gr. 1.101 1.055
(de Muynck, W. Ann. 1894, 53. 561.)

$\text{CdSO}_4 + \text{Aq}$ containing 13.40% CdSO_4 has sp. gr. 20°/20° = 1.1429.
 $\text{CdSO}_4 + \text{Aq}$ containing 16.79% CdSO_4 has sp. gr. 20°/20° = 1.1847.
(Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 282.)

gr. of CdSO₄ at 18.2°, when p = per cent strength of solution; d = observed density; w = volume conc. in grams per cc. ($\frac{pd}{100} = w$.)

p	d	w
39.86	1.5639	0.6231
31.53	1.4080	0.4439
26.85	1.3310	0.3574
24.17	1.2901	0.3118
18.35	1.2084	0.2217
13.27	1.1437	0.1518
9.97	1.1045	0.1102
7.46	1.0764	0.0803
6.12	0.0619	0.0650
2.52	0.0242	0.0259
1.45	0.0132	0.0147
0.464	0.0033	0.0046

(Barnes, J. phys. Ch. 1898, 2. 543.)

Sp. gr. of CdSO₄ + Aq sat. at 25° and 1 atm. 1.617. (Sinnige, Z. phys. Ch. 1909, 67. 8.)
See also under CdSO₄ + 2²/₃H₂O, and 4H₂O.
Insol. in liquid NH₃. (Franklin, Am. Ch. 1898, 20. 827.)
Insol. in acetone. (Naumann, B. 1904, 37. 29); methyl acetate. (Naumann, B. 1909, 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)
+H₂O. See Mylius and Funk, under 1SO₄.
+1¹/₂ H₂O. (Worobieff, Bull. Soc. 1896, 16. 1754.)
+2²/₃H₂O. 1 pt. H₂O dissolves 0.59 pt. anhydrous salt at 23°, and not much more on heating. Sat. solution boils at 102°. Precipitated by alcohol. (v. Hauer.)

100 g. H₂O dissolve g. CdSO₄ at t°.

t°	G. CdSO ₄
0	75.52
5	75.65
7	65.73
9	75.85
11.5	75.94
13	76.04
15	76.11
16	76.16
17	76.13
18	76.14
19	76.18
25	76.79

Johnstamm and Cohen, W. Ann. 1898, 65. 352.)

100 pts. H₂O dissolves at:
13.7° 14.98° 15.0° 16.0°
76.06 76.09 76.14 76.18 pts. anhydrous salt,
16.96° 18.0° 19.0° 25.0°
76.26 76.32 76.39 76.81 pts. anhydrous salt.
(Steinwehr, W. Ann. 1902, (4) 9. 1050.)

100 g. H₂O dissolve 76.02 g. CdSO₄ at 25°. (Stortenbecker, Z. phys. Ch. 1900, 34. 109.)
Solubility of CdSO₄ + 2²/₃H₂O at 25° and varying pressures.

Pressure in atmospheres	G. CdSO ₄ in 100 g. H ₂ O
1	76.80
500	77.85
500	78.08
1000	78.77
1000	78.68

Det. by another method

Pressure in atmospheres	G. CdSO ₄ in 100 g. H ₂ O
250	77.53
500	78.02
750	78.60
1000	78.96

(Cohen and Sinnige, Trans. Farad. Soc. 1910, 5. 269.)

Sp. gr. at 21.6°/0° of CdSO₄ + Aq containing 11.47% CdSO₄ + ²/₃H₂O = 1.0944. (Kannonnikoff, J. pr. 1885, (2) 31. 346.)
100 g. H₂O dissolve 57.61 g. CdSO₄ + 10.63 g. FeSO₄ at 25°. (Stortenbecker, Z. phys. Ch. 1900, 34. 109.)
+4H₂O. (Lescœur, A. ch. 1895, (7) 4. 222.)
Sp. gr. at 15° of CdSO₄ + Aq containing 10 g. CdSO₄ + 4H₂O in 100 c.c of solution = 1.0790; containing 20 g. CdSO₄ + 4H₂O in 100 cc. of solution = 1.1522. (Traube, J. pr. 1885, (2) 31. 207.)
Could not be obtained. (Mylius and Funk.)
+7H₂O.

Solubility in H₂O.

t°	% CdSO ₄
-17	44.45
-16	44.5
-12	45.3
-10	46.1
-7	47.5
-5	48.5
-4.5	48.7

(Mylius and Funk, B. 1897, 30. 828.)

Cadmium caesium sulphate, CdSO₄, Cs₂SO₄ + 6H₂O.

Sol. in H₂O. (Tutton, Chem. Soc. 63. 337.)
1 l. H₂O dissolves 1399 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Cadmium calcium potassium sulphate,
 $\text{Ca}_2\text{CdK}_2(\text{SO}_4)_4 + 2\text{H}_2\text{O}$.
 (D'Ans, B. 1908, 41. 1778.)

Cadmium cerium sulphate, $\text{CdSO}_4 \cdot \text{Ce}_2(\text{SO}_4)_3 + 6\text{H}_2\text{O}$.
 Sol. in H_2O . (Wyruboff.)

Cadmium hydrazine sulphate,
 $\text{CdH}_2(\text{SO}_4)_2 \cdot 2\text{N}_2\text{H}_4$.
 1 pt. is sol. in 202.5 pts. H_2O at 12° .
 Not attacked by dil. acids. Easily sol. in
 $\text{NH}_4\text{OH} + \text{Aq}$. (Curtius, J. pr. 1894, (2) 50.
 331.)

Cadmium magnesium sulphate, $\text{CdSO}_4 \cdot \text{MgSO}_4 + 14\text{H}_2\text{O}$.
 Very efflorescent. Sol. in H_2O . (Schiff, A.
 104. 325.)

Cadmium potassium sulphate, $\text{K}_2\text{SO}_4 \cdot \text{CdSO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$.
 Sol. in H_2O . (v. Hauer, Pogg. 133. 176.)
 100 pts. H_2O dissolve 42.50 pts. anhydrous
 salt at 26° .
 100 pts. H_2O dissolves 42.80 pts. anhydrous
 salt at 31° .
 100 pts. H_2O dissolve 43.45 pts. anhydrous
 salt at 40° .
 100 pts. H_2O dissolve 44.90 pts. anhydrous
 salt at 64° . (Wyruboff, Bull. Soc. Min.
 1901, 24. 68.)
 $+2\text{H}_2\text{O}$. 100 pts. H_2O dissolve 42.89 pts.
 anhydrous salt at 16° ; 46.82 pts. at 31° ;
 47.40 pts. at 40° . (Wyruboff.)
 $+4\text{H}_2\text{O}$. Efflorescent. (Wyruboff, Bull.
 Soc. Min. 1891, 14. 235.)
 $+6\text{H}_2\text{O}$. Very efflorescent, and easily
 decomp. (Schiff.)
 Does not exist. (Wyruboff.)

Cadmium rubidium sulphate, $\text{CdSO}_4 \cdot \text{Rb}_2\text{SO}_4 + 6\text{H}_2\text{O}$.
 Efflorescent. Sol. in H_2O . (Tutton.)
 1 l H_2O dissolves 767 g. anhydrous salt
 at 25° . (Locke, Am. Ch. J. 1902, 27. 459.)

Cadmium sodium sulphate, $\text{CdSO}_4 \cdot \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$.
 Sol. in H_2O . (v. Hauer.)

Solubility of $\text{CdNa}_2(\text{SO}_4)_2 + 2\text{H}_2\text{O}$ in 100 g
 H_2O at t°

t°	Grams CdSO_4	Grams Na_2SO_4
24	35.49	24.04
25	35.88	24.46
30	36.28	24.605
35	36.69	24.99
40	37.24	25.455

(Koppel, Z. phys. Ch. 1905, 52. 413.)
 Decomp. by H_2O below 20.5° .

Solubility of $\text{CdNa}_2(\text{SO}_4)_2 \cdot 2\text{H}_2\text{O} + \text{CdSO}_4$
 $\frac{1}{2}\text{H}_2\text{O}$ in 100 g. H_2O at t° .

t°	Grams CdSO_4	Grams Na_2SO_4	t°	Grams CdSO_4	Grams Na_2SO_4
0	73.51	4.85	15	73.76	9.45
5	73.38	8.67	20	73.81	9.45
10	72.76	3.55	25	73.71	10.48
			30	73.01	11.21
			40	73.28	13.13

(Koppel)

Solubility of $\text{CdNa}_2(\text{SO}_4)_2 \cdot 2\text{H}_2\text{O} + \text{Na}_2\text{SO}_4$
 $10\text{H}_2\text{O}$ in 100 g. H_2O at t°

t°	Grams CdSO_4	Grams Na_2SO_4
-14	8	72.68
0		66.325
5		61.78
10		55.34
12		51.615
15		46.60
19	8	36.13
20		36.25
24		27.82
25		25.59
30		14.62
		8.32
		11.625
		12.97
		14.785
		15.95
		17.99
		22.16
		23.52
		29.17
		31.06
		44.145

(Koppel.)

Solubility of $\text{CdNa}_2(\text{SO}_4)_2 \cdot 2\text{H}_2\text{O} + \text{Na}_2\text{SO}_4$
 (anhydrous) in 100 g. H_2O at t°

t°	Grams CdSO_4	Grams Na_2SO_4
35	13.26	47.06
40	16.25	46.27

(Koppel.)

Cadmium sulphate ammonia, $\text{CdSO}_4 \cdot 6\text{NH}_3$.
 Sol. in H_2O with separation of CdO . (Rosa,
 Pogg. 20. 152.)
 $\text{CdSO}_4 \cdot 4\text{NH}_3 + 4\text{H}_2\text{O}$. Decomp. by H_2O .
 (Malaguti and Sarzeau, A. ch. (3) 9. 431.)
 $+2\text{H}_2\text{O}$. Ppt. (André, C. R. 104. 987.)
 $+2\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O with separation of
 basic sulphate. (Müller, A. 149. 70.)
 $\text{CdSO}_4 \cdot 3\text{NH}_3$. (Isambert, C. R. 1870, 70.
 457.)

Cadmium sulphate cupric oxide, $\text{CdSO}_4 \cdot 3\text{CuO} + x\text{H}_2\text{O}$.
 (Recoura, C. R. 1901, 132. 1415.)
 $2\text{CdSO}_4 \cdot 3\text{CuO} + 8\text{H}_2\text{O}$. (Mailhe, A. ch.
 1902, (7) 27. 383.)
 $+12\text{H}_2\text{O}$. (Mailhe.)
 $6\text{CdSO}_4 \cdot 20\text{CuO} + x\text{H}_2\text{O}$. (Recoura, C. R.
 1901, 132. 1415.)

Cadmium sulphate hydrazine, $\text{CdSO}_4 \cdot 2\text{N}_2\text{H}_4$.

Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$ but cannot be
 cryst. therefrom. (Fransen, Z. anorg. 1909,
 60. 282.)

1 sulphate hydrogen chloride,
 $\text{SO}_4, 4\text{HCl}+4\text{H}_2\text{O}.$
deliquescent. (Baskerville and Harris,
hem. Soc. 1901, **23**. 896.)
 $4, 8\text{HCl}.$ Very deliquescent. (Bas-
nd Harris.)

sulphate, $\text{Cs}_2\text{SO}_4.$
deliquescent.
3. H_2O dissolve 158.7 pts. Cs_2SO_4 at
. H_2O at 17–18° dissolve 163.5 g.
(Tutton, Chem. Soc. 1894, **65**.)

Solubility in $\text{H}_2\text{O}.$

Cs_2SO_4 100 g.	G. Cs_2SO_4 per l.	t°	G. Cs_2SO_4 per 100 g.		G. Cs_2SO_4 per l.
			Solu- tion	H_2O	
H_2O	G. Cs_2SO_4 per l.	t°	Solu- tion	H_2O	G. Cs_2SO_4 per l.
167.1	3.42	60	66.7	199.9	3.78
173.1	3.49	70	67.2	205.0	3.83
178.7	3.56	80	67.8	210.3	3.88
184.1	3.62	90	68.3	214.9	3.92
189.9	3.68	100	68.8	220.3	3.97
194.9	3.73	108.6	69.2	224.5	4.00

, Trans. Roy. Soc. 1904, **203**. A.
210.)

ity in $\text{Na}_2\text{SO}_4+\text{Aq}.$
olution contains 54.7% Cs_2SO_4+
 Na_2SO_4 at 25°. (Foote, J. Am.
oc. 1911, **33**. 467.)
in alcohol. (Bunsen.)
in methyl acetate. (Naumann, B.
3790.)
in acetone. (Naumann, B. 1904, **37**.
idmann, C. C. 1899, II. 1014.)
ity in glycol at ord. temp.=3.0–
(de Coninck, Belg. Acad. Bull.
l.)

hydrogen sulphate, $\text{CsHSO}_4.$
 $\text{H}_2\text{O}.$

pyrosulphate, $\text{Cs}_2\text{S}_2\text{O}_7.$
p. by $\text{H}_2\text{O}.$

octosulphate, $\text{Cs}_2\text{S}_8\text{O}_{22}.$
p. by $\text{H}_2\text{O}.$ (Weber, B. 17. 2497.)

calcium sulphate, $\text{Ca}_2\text{Cs}_2(\text{SO}_4)_3.$
table, (D'Ans, B. 1908, **41**. 1776.)

chromium sulphate, $\text{Cs}_2\text{Cr}_2(\text{SO}_4)_4.$
 $\text{H}_2\text{O}.$
in crystal H_2O at 116°. (Locke.)

Solubility in $\text{H}_2\text{O}.$

Temp.	G. per litre	G. mols. anhydrous salt per l.
25°	5.7	0.015
30°	9.6	0.025
35°	12.06	0.032
40°	15.3	0.0405

(Locke, Am. Ch. J. 1901, **26**. 180.)

**Cæsium cobaltous sulphate, $\text{Cs}_2\text{SO}_4, \text{CoSO}_4+$
 $6\text{H}_2\text{O}.$**

Sol. in $\text{H}_2\text{O}.$ (Tutton, Chem. Soc. **63**.
337.)

1 l. H_2O dissolves 418.8 g. anhydrous
salt at 25°. (Locke, Am. Ch. J. 1902, **27**.
459.)

**Cæsium cobaltic sulphate, $\text{Cs}_2\text{Co}_2(\text{SO}_4)_4+$
 $24\text{H}_2\text{O}.$**

Melts in crystal H_2O at 116°. (Locke,
Am. Ch. J. 1901, **26**. 183.)

**Cæsium copper sulphate, $\text{Cs}_2\text{SO}_4, \text{CuSO}_4+$
 $6\text{H}_2\text{O}.$**

Sol. in $\text{H}_2\text{O}.$ (Tutton.)
1 l. H_2O dissolves 460 g. anhydrous salt
at 25°. (Locke, Am. Ch. J. 1902, **27**. 459.)

**Cæsium gallium sulphate, $\text{Cs}_2\text{Ga}_2(\text{SO}_4)_4+$
 $24\text{H}_2\text{O}.$**

(Soret, Arch. sc. phys. nat. 1888, (3) **20**.
531.)

**Cæsium indium sulphate, $\text{Cs}_2\text{In}_2(\text{SO}_4)_4+$
 $24\text{H}_2\text{O}.$**

75.7 g. anhydrous (117.39 hydrated) salt
or 0.172 g. mols of anhydrous salt are sol.
in 1 l. H_2O at 25°. (Locke, Am. Ch. J. 1901,
26. 175.)

100 pts. H_2O dissolve 3.04 pts. at 16.5°.
(Chabriè and Rengade, C. R. 1900, **131**. 1301.)

**Cæsium iridium sulphate, $\text{Cs}_2\text{SO}_4,$
 $\text{Ir}_2(\text{SO}_4)_3+24\text{H}_2\text{O}.$**

Mpt. 109–110°
Very sl. sol. in cold. More easily sol. in
hot $\text{H}_2\text{O}.$ (Marino, Z. anorg. 1904, **42**. 218.)

**Cæsium iron (ferrous) sulphate, $\text{Cs}_2\text{SO}_4,$
 $\text{FeSO}_4+6\text{H}_2\text{O}.$**

Sol. in $\text{H}_2\text{O}.$ (Tutton.)
1 l. H_2O dissolves 1011 g. anhydrous salt
at 25°. (Locke, Am. Ch. J. 1902, **27**. 459.)

**Cæsium iron (ferric) sulphate, $\text{Cs}_2\text{Fe}_2(\text{SO}_4)_4$
 $+24\text{H}_2\text{O}.$**

Melts in crystal H_2O at 71°. (Locke.)

Burnt gypsum easily forms supersat. solutions containing nearly 1% CaSO_4 . It forms supersat. solutions more readily at 0° , and that tendency decreases with increase of temp., hence figures in (d) which contained burnt gypsum. (Goldammer, C. C. 1888. 708.)

Calculated from electrical conductivity of $\text{CaSO}_4 + \text{Aq}$, 1 l. H_2O dissolves 2.07 g. CaSO_4 at 18° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

The anhydrous salt varies in solubility. Solubility depends (1) upon temp. and time of drying, (2) upon the relative amount of salt, (3) upon time of shaking. Possibly α and β modifications. (Pottlitz, C. C. 1894, II. 515.)

2.04 gr. are dissolved in 1 liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

At 15° and after 5 minutes shaking, the highest degree of supersaturation which can be obtained with pure sol. calcium sulphate = 9.47 g. of the anhydrous salt or 11.976 g. $\text{CaSO}_4 + 2\text{H}_2\text{O}$ in 1 l. of H_2O . (Cavazzi, C. C. 1905, I. 1694.)

Solubility of CaSO_4 in 100 pts. H_2O at high temp.

t°	Pts. CaSO_4	t°	Pts. CaSO_4	t°	Pts. CaSO_4
140	0.078	175-185	0.027	250	0.016
165	0.056	240	0.018

(Tilden and Shenstone, Phil. Trans. 1884. 31.)

Pptn. of CaSO_4 which has been started by heating solution to $140-150^\circ$ continues even after solution has cooled. (Storer.)

CaSO_4 is completely insol. in sea water or pure H_2O at temperatures between 140° and 150° . (Cousté.)

Solubility of CaSO_4 in sea water at temperatures over 100° . t° = temp.; P = pressure in atmospheres; % = per cent CaSO_4 in sat. solution.

t°	P	%	t°	P	%
103	1	0.500	118.5	1.50	0.226
103.8	1	0.477	121.2	1.5	0.183
105.15	1	0.432	124	2	0.140
108.6	1.25	0.395	127.9	2	0.097
111	1.25	0.355	130°	2.5	0.060
113.2	1.25	0.310	133.3	2.5	0.023
115.8	1.50	0.267

(Cousté, Ann. Min. (5) 5. 80.)

Solubility of CaSO_4 in H_2O at various pressures.

100 g. sat. $\text{CaSO}_4 + \text{Aq}$ at 1 atmos. pressure and 15° contain 0.206 g. CaSO_4 ; at 20 atmos. pressure and 15° contain 0.227 g. CaSO_4 ; at 1 atmos. pressure and 16.2° contain 0.213 g. CaSO_4 . (Möller, Pogg. 117. 386.)

Soluble anhydrite:

1 l. H_2O dissolves 22.8 milliequivalents at 100° .

1 l. H_2O dissolves 6.4 milliequivalents at 156° .

Anhydrite:

1 l. H_2O dissolves 9.2 milliequivalents at 100° .

1 l. H_2O dissolves 2.7 milliequivalents at 156° .

1 l. H_2O dissolves 0.7 milliequivalents at 218° .

(Melcher, J. Am. Chem. Soc. 1910, 32. 61.)

See also under gypsum, p. 953.

Maximum solubility is at 37.5° . (Cameron, J. phys. Chem. 1901, 5. 572.)

Sp. gr. of sat. $\text{CaSO}_4 + \text{Aq}$ at $15^\circ = 1.0022$ (Stolba, J. pr. 97. 503.)

Sp. gr. of sat. $\text{CaSO}_4 + \text{Aq}$ at $31^\circ = 1.0081$

1 pt. CaSO_4 is sol. in 218 pts. H_2O containing CO_2 . (Beyer, Arch. Pharm. (2) 180. 193.)

Sl. sol. in cold $\text{HCl} + \text{Aq}$; completely sol. in boiling dil. HCl or $\text{HNO}_3 + \text{Aq}$. (Rose, Pogg. 95. 108.)

Solubility of CaSO_4 in $\text{HCl} + \text{Aq}$.

t°	% HCl	100 cc. dissolve g. of CaSO_4	t°	% HCl	100 cc. dissolve g. of CaSO_4
25	0.77	0.6405	25	6.12	1.6539
25	1.56	0.8821	101	0.77	1.1309
25	3.06	1.2639	102	3.06	3.1730
25	4.70	1.5342	103	6.12	4.6902

(Lunge, J. Soc. Chem. Ind. 4. 31.)

Solubility in $\text{HNO}_3 + \text{Aq}$ at 25° .

g. HNO_3 per 100 cc.	g. CaSO_4 per 100 cc. solution	g. HNO_3 per 100 cc.	g. CaSO_4 per 100 cc. solution
0	0.208	6	1.48
1	0.56	8	1.70
2	0.82	10	1.84
3	1.02	12	1.96
4	1.20		

(Banthisch, J. pr. 1884, 29. 52.)

For solubility in H_2SO_4 see $\text{CaH}_2(\text{SO}_4)_2$.

Solubility in $\text{H}_3\text{PO}_4 + \text{Aq}$ at 25° .

G. P_2O_5 per l.	G. CaSO_4 per l.	Sp. gr. $25^\circ 25'$
0.0	2.126	
5.0	3.138	1.002
10.5	3.734	1.007
21.4	4.456	1.016
46.3	5.760	1.035
105.3	7.318	1.075
145.1	7.920	1.106
204.9	8.383	1.145
312.0	7.965	1.221
395.7	6.848	1.230
494.6	5.573	1.344

(Taber, J. phys. Chem. 1906, 10. 625.)

Solubility in formic acid at 25° .

100 cc. of solution of acid containing 4

issolve 0.24 g. CaSO_4 . (Banthisch, J. pr. 884, 29. 52.)

Solubility of CaSO_4 in chloracetic acid t 25°. 100 cc. of solution of acid containing % dissolve 0.22 g. CaSO_4 ; 10%, 0.25 g. Banthisch, J. pr. 1884, 29. 52.)

Solubility in H_2O is increased by presence f NH_4Cl (Vogel, J. pr. 1. 196), ammonium acetate (Wittstein, Repert. 57. 18), $\text{NH}_4)_2\text{SO}_4$, $(\text{NH}_4)_2\text{B}_4\text{O}_7$ (Popp, A. Suppl. . 11); also KNO_3 (Vogel, Jun.), Na_2SO_4 Henry, J. Pharm. 12. 31), NaCl (Tromms- orf, N. J. Pharm. 18, 1. 234.)

Decomp. by alkali carbonates + Aq. (See torer's Dict.)

1 g. CaSO_4 is sol. in 162 ccm. sat. $\text{KCl}+\text{Aq}$ t 8°; in 147 ccm. sat. $\text{NaCl}+\text{Aq}$ at 8.5°; in 3 ccm. sat. $\text{NH}_4\text{Cl}+\text{Aq}$ at 12.5°; in 94 ccm. st. KNO_3+Aq ; in 92 ccm. sat. NaNO_3+Aq ; i 320 ccm. sat. $\text{NH}_4\text{NO}_3+\text{Aq}$; in 54 ccm. , sat. $\text{NH}_4\text{NO}_3+\text{Aq}$; in about 2000 ccm. st. $\text{K}_2\text{SO}_4+\text{Aq}$. (Droese.)

More sol. in Fe_2Cl_6 , Cr_2Cl_6 , CuCl_2 , ZnCl_2 + q than in H_2O , but not more sol. in CaCl_2 + q. (Gladstone.)

$\text{NH}_4\text{Cl}+\text{Aq}$.

1 g. CaSO_4 is sol. in 92 ccm. sat. $\text{NH}_4\text{Cl}+\text{q}$ at 13.5°; in 94 ccm. $\frac{1}{2}$ sat. $\text{NH}_4\text{Cl}+\text{Aq}$ at 3.5–15.5°; in 200 ccm. $\frac{1}{3}$ sat. $\text{NH}_4\text{Cl}+\text{Aq}$ at 3.5°; in 183 ccm. $\frac{1}{4}$ sat. $\text{NH}_4\text{Cl}+\text{Aq}$ at 100°. Fassbender, B. 9. 1360.)

olubility of CaSO_4 in 25% $\text{NH}_4\text{Cl}+\text{Aq}$.

t°	% CaSO_4	t°	% CaSO_4
8	1.030	60	1.333
9	1.023	80	1.026
25	1.096	120	1.000
39	1.126

Tilden and Shenstone, Roy. Soc. Proc. 38. 335.)

Solubility in $\text{NH}_4\text{Cl}+\text{Aq}$ increases with per- ntage of NH_4Cl , but if solution contains ore than 60 g. NH_4Cl per l. more CaO dis- lves than SO_3 . With 333 g. NH_4Cl per the solution contains 4.9 g. SO_3 and 4.4 g. CaO , while the SO_3 content requires only 4 g. CaO . (Ditte, C. R. 1898, 126. 694.)

solubility of CaSO_4 in $\text{NH}_4\text{Cl}+\text{Aq}$ at 25°.

Grams NH_4Cl per liter	Grams CaSO_4 per liter
10.8	3.90
24.4	5.38
46.7	7.07
94.5	8.80
149.7	10.30
198.6	10.85
210.0	10.88
275.0	10.60
325.0	9.40
375.3 (saturated)	7.38

Cameron and Brown, J. phys. Chem. 1905, 9. 211.)

CaCl_2+Aq .

Solubility of CaSO_4 in CaCl_2+Aq at t°.

t°	% CaCl_2	100 ccm. dissolve g. of CaSO_4	t°	% CaCl_2	100 ccm. dissolve g. of CaSO_4
23	3.54	0.1225	25	16.91	0.0702
24	6.94	0.0963	101.0	3.54	0.1370
25	10.36	0.0886	102.5	10.36	0.1426
25	15.90	0.0734	103.5	16.91	0.1301

(Lunge, l. c.)

Solubility of CaSO_4 in H_2O containing various amts. of CaCl_2 at 20°. 100 pts. H_2O con- taining pts. CaCl_2 dissolve pts. CaSO_4 .

Pts. CaCl_2	Pts. CaSO_4	Pts. CaCl_2	Pts. CaSO_4
0.00	0.225	19.80	0.041
11.50	0.078	51.00	0.000
14.39	0.063	67.05	0.000

(Tilden and Shenstone.)

Solubility of CaSO_4 in CaCl_2+Aq at t°.

t°	% CaCl_2	% CaSO_4	t°	% CaCl_2	% CaSO_4
15	15.00	0.063	94	15.16	0.110
21	14.70	0.068	138	14.70	0.071
39	15.00	0.091	170	14.82	0.031
72	14.90	0.100	195	14.70	0.022

(Tilden and Shenstone, l. c.)

Solubility in CaCl_2+Aq at 25°.

g. per l. of solution		g. per l. of solution	
CaCl_2	CaSO_4	CaCl_2	CaSO_4
0.00	2.06	51.53	1.02
7.49	1.24	97.02	0.84
11.96	1.18	192.71	0.47
25.77	1.10	280.30	0.20
32.05	1.08	367.85	0.03

(Cameron and Seidell, J. phys. Ch. 1901, 5. 643.)

1000 pts. of 1% CaCl_2+Aq . dissolve 1.1414 pts. CaSO_4 ; 40% CaCl_2 , 0.2130 pts. CaSO_4 . (Orloff, Chem. Soc. 1903, 24, (2) 211.)

Solubility in $\text{CaO}_2\text{H}_2 + \text{Aq}$ at 25° .

G. CaSO_4 per l.	G. CaO per l.	Solid phase
0.0	1.166	$\text{Ca}(\text{OH})_2$
0.391	1.141	"
0.666	1.150	"
0.955	1.215	"
1.214	1.242	"
1.588	1.222	$\text{Ca}(\text{OH})_2$ and $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$
1.634	0.939	$\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$
1.722	0.611	"
1.853	0.349	"
1.918	0.176	"
2.030	0.062	"
2.126	0.0	"

(Cameron and Bell, J. Am. Chem. Soc. 1906, 28. 1221.)

$\text{MgCl}_2 + \text{Aq.}$

Sol. in 324 pts. $\text{MgCl}_2 + \text{Aq}$ (34.1% MgCl_2) at 19° . (Karsten.)
1 g. CaSO_4 is sol. in 146 ccm. $\frac{1}{2}$ sat. $\text{MgCl}_2 + \text{Aq}$ at 13.5° . (Fassbender.)
1 l. $\frac{1}{2}$ sat. $\text{MgCl}_2 + \text{Aq}$ dissolves 6.83 g. $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ at 13.5° . (Droeze.)

Solubility of CaSO_4 in $\text{MgCl}_2 + \text{Aq.}$

t°	% MgCl_2	% CaSO_4
9	19.7	0.765
39	11.1	2.744
80	9.99	1.038

(Tilden and Shenstone, *l. c.*)

Solubility in $\text{MgCl}_2 + \text{Aq}$ at 26° .

g. per l. of solution			g. per l. of solution		
MgCl_2	CaSO_4	H_2O	MgCl_2	CaSO_4	H_2O
0.0	2.08	997.9	121.38	8.62	972.2
8.50	4.26	996.5	206.98	6.57	949.9
19.18	5.69	994.5	337.0	2.77	908.7
46.64	7.59	989.1	441.0	1.39	878.6

(Cameron and Seidell, J. phys. Ch. 1901, 5. 645.)

1 l. sat. $\text{MgCl}_2 + \text{Aq}$ at 25° containing 476.5 g. MgCl_2 dissolves 1.09 g. CaSO_4 . (Cameron and Brown, J. phys. Ch. 1905, 9. 214.)

$\text{NH}_4\text{NO}_3 + \text{Aq.}$

1 g. CaSO_4 is sol. in 320 ccm. sat. $\text{NH}_4\text{NO}_3 + \text{Aq}$ at $8-9^\circ$; in 54 ccm. $\frac{2}{3}$ sat. $\text{NH}_4\text{NO}_3 + \text{Aq}$ at 13.5° ; in 103 ccm. $\frac{2}{3}$ sat. $\text{NH}_4\text{NO}_3 + \text{Aq}$ at 13.5° . (Fassbender.)

Solubility of CaSO_4 in $\text{NH}_4\text{NO}_3 + \text{Aq}$ at 25° .

G. NH_4NO_3 per l.	G. CaSO_4 per l.
10	3.18
25	3.93
55	5.80
100	7.65
150	8.88
200	9.65
300	10.80
400	11.40
550	12.02
750	12.20
1000	11.81
1200	11.10
1400	10.02
saturated	7.55

(Cameron and Brown, J. phys. Chem. 1906, 9. 213.)

$\text{Ca}(\text{NO}_3)_2 + \text{Aq.}$

Solubility of CaSO_4 in $\text{Ca}(\text{NO}_3)_2 + \text{Aq}$ at 25° .

Weight of 1000 cc. of solution	G. $\text{Ca}(\text{NO}_3)_2$ per l.	G. CaSO_4 per l.
998.1	0	2.064
1013.8	25	1.238
1031.7	50	1.196
1067.3	100	1.134
1136.9	200	0.929
1203.5	300	0.759
1265.6	400	0.569
1328.1	500	0.403
1352.0	544	0.346

(Seidell and Smith, J. phys. Chem. 1904, 8. 498.)

$\text{Mg}(\text{NO}_3)_2 + \text{Aq.}$

Solubility of CaSO_4 in $\text{Mg}(\text{NO}_3)_2 + \text{Aq}$ at 25° .

Weight of 1000 cc. of solution grams	G. $\text{Mg}(\text{NO}_3)_2$ per l.	G. CaSO_4 per l.
998.1	0	2.064
1020.5	25	5.772
1039.8	50	7.884
1078.6	100	9.920
1149.8	200	13.340
1219.0	300	14.000
1282.1	400	14.683
1355.3	514	15.040

(Seidell and Smith, J. phys. Chem. 1904, 8. 497.)

1 l. sat. $\text{Mg}(\text{NO}_3)_2 + \text{Aq}$ at 25° containing 615.1 g. $\text{Mg}(\text{NO}_3)_2$ dissolves 15.26 g. CaSO_4 . (Cameron and Brown, J. phys. Ch. 1906, 9. 214.)

KNO₃+Aq.

1 g. CaSO₄ is sol. in 94 ccm. sat. KNO₃+Aq at 13.5°; in 82 ccm. sat. KNO₃+Aq at 15.5°; in 68 ccm. nearly sat. KNO₃+Aq at 20°. (Fassbender.)

Solubility in KNO₃+Aq at 25°.

Wt. of 1000 ccm. of solution grams	G. KNO ₃ per l.	G. CaSO ₄ per l.
998.1	0.0	2.084
1008.1	12.5	3.284
1015.4	25.0	4.080
1032.1	50.0	5.255
1062.5	100.0	6.855
1092.4	150.0	7.907
1122.4	200.0	8.688
1153.9	260.0	a { 6.278 12.112

a Probably due to formation of double salt of calcium and potassium sulphates, CaK₂(SO₄)₂+H₂O.

(Seidell and Smith, J. phys. Chem. 1908, 8. 496.)

NaNO₃+Aq.

1 g. CaSO₄ is sol. in 92 ccm. sat. NaNO₃+Aq at 8.5°; in 318 ccm. 1/2 sat. NaNO₃+Aq at 13.5°. (Fassbender.)

100 ccm. sat. NaNO₃+Aq dissolve 1.086 g. CaSO₄+2H₂O; 100 ccm. 1/2 sat. NaNO₃+Aq dissolve 0.314 g. CaSO₄+2H₂O. (Droeze, B. 10. 338.)

Solubility in NaNO₃+Aq at 25°.

Wt. of 1000 ccm. of solution grams	G. NaNO ₃ per l.	G. CaSO ₄ per l.
998.1	0	2.084
1016.3	25	4.252
1034.0	50	5.500
1058.4	100	7.100
1133.6	200	8.790
1191.6	300	9.282
1363.9	600	7.886
1390.4	655	7.238

(Seidell and Smith, J. phys. Chem. 1904, 8. 495.)

1 l. sat. NaNO₃+Aq at 25°, containing 668.4 g. NaNO₃, dissolves 5.52 g. CaSO₄. (Cameron and Brown, J. phys. Ch. 1905, 9. 214.)

Solubility in KBr+Aq at 21°.

G. KBr per l.	G. CaSO ₄ per l.	G. KBr per l.	G. CaSO ₄ per l.
0	2.05	100	6.3
10	3.1	125	6.7
20	3.6	150	7.0
40	4.5	200	7.3
60	5.2	250	Double salt.
80	5.9		

(Ditte, A. ch. 1898, (7) 14. 294.)

KCl+Aq.

1 g. CaSO₄ is sol. in 162 ccm. sat. KCl+Aq at 8°; in 295 ccm. 1/2 sat. KCl+Aq at 9°.

Solubility in KCl+Aq at 21°.

g. per l.		g. per l.	
KCl	CaSO ₄	KCl	CaSO ₄
0	2.05	60	6.6
10	3.6	80	7.2
20	4.5	100	7.5
40	5.8	125	Double Salt

(Ditte, A. ch. 1898, (7) 14. 294.)

Solubility in KI+Aq at 21°.

G. KI per l.	G. CaSO ₄ per l.	G. KI per l.	G. CaSO ₄ per l.
0	2.05	100	5.1
10	2.8	125	5.45
20	3.2	150	5.8
40	3.9	200	5.95
60	4.5	250	6.00
80	4.85	300	Double salt.

(Ditte, l. c.)

NaCl+Aq.

Sol. in 122 pts. sat. NaCl+Aq. (Anthon.)

Insol. in sat. NaCl+Aq, but more sol. in dil. NaCl+Aq than in H₂O. Maximum solubility in NaCl+Aq is when the sp. gr. is 1.033.

1 g. CaSO₄ is sol. in 147 ccm. of sat. NaCl+Aq at 8.5°; in 150 ccm. of sat. NaCl+Aq at 13.5°; in 149 ccm. of 1/2 sat. NaCl+Aq at 13.5°; in 244 ccm. of 1/2 sat. NaCl+Aq at 13.5°. (Fassbender.)

100 ccm. sat. NaCl+Aq dissolve 0.6785 g. CaSO₄+2H₂O at 8.5°; 0.6665 g. CaSO₄+2H₂O at 13.5°. 100 ccm. 1/2 sat. NaCl+Aq dissolve 0.671 g. CaSO₄+2H₂O at 13.5°; 1/2 sat. NaCl+Aq dissolve 0.4085 g. CaSO₄+2H₂O at 13.5°. (Droeze.)

Solubility of CaSO₄ in NaCl+Aq at t°.

t°	% NaCl	% CaSO ₄	t°	% NaCl	% CaSO ₄
20	19.90	0.823	130	19.92	0.392
44	19.93	0.830	165	20.04	0.250
67	19.95	0.832	169	20.05	0.244
85	19.90	0.823	179	20.10	0.229
101	20.08	0.682	225	21.00	0.178

(Tilden and Shenstone, Roy. Soc. Proc. 38. 331.)

Solubility of CaSO_4 in $\text{NaCl} + \text{Aq}$ at t° .

t°	% NaCl	100 ccm. dissolve g. of CaSO_4	t°	% NaCl	100 ccm. dissolve g. of CaSO_4
21.5	3.53	0.5115	17.5	17.46	0.7369
19.5	7.35	0.6429	101.0	3.53	0.4891
21	11.12	0.7215	102.5	14.18	0.6248
18	14.18	0.7340		17.46	0.6299

(Lunge, J. Soc. Chem. Ind. 4. 31.)

100 pts. H_2O containing pts. NaCl dissolve
pts. CaSO_4 at 20° .

Pts. NaCl	Pts. CaSO_4	Pts. NaCl	Pts. CaSO_4	Pts. NaCl	Pts. CaSO_4
0.00	0.225	5.05	6.34	24.40	0.820
0.52	0.301	10.00	7.38	35.10	0.734
2.03	0.441	20.00	0.823	35.86	0.709
5.02	6.15

(Tilden and Shenstone.)

Solubility in $\text{NaCl} + \text{Aq}$ at 28° .

g per l.		wt of 1 cc solution	g per l.		wt of 1 cc. solution
NaCl	CaSO_4		NaCl	CaSO_4	
0.00	2.12	0.9998	176.50	7.12	1.1196
9.11	6.66	1.0644	228.76	6.79	1.1488
143.99	7.18	1.0981	264.17	6.50	1.1707
148.34	7.16	1.012	320.49	5.72	1.2034

(Cameron, J. phys. Ch. 1901, 5. 556.)

Solubility in $\text{NaCl} + \text{Aq}$ at 15° .

G. CaSO_4 per l.	G. NaCl per l.
2.3	0.6
2.5	1.1
3.1	5.1
3.7	10.6
4.8	31.1
5.6	51.4
7.4	139.9

(Cameron, J. phys. Ch. 1901, 5. 559.)

Solubility in $\text{NaCl} + \text{Aq}$ at 26° .

NaCl in 100 g. H_2O	CaSO_4 in 100 g. H_2O
0.0000	0.2126
9.4307	0.6846
15.2056	0.7581
15.6859	0.7575
18.8570	0.7605
25.0478	0.7439
29.3509	0.7219
36.5343	0.6515

(Cameron, J. phys. Ch. 1901, 5. 564.)

Solubility in $\text{NaCl} + \text{Aq}$.

30°		52°		70°		92°	
G. NaCl per l.	G. CaSO_4 per l.	G. NaCl per l.	G. CaSO_4 per l.	G. NaCl per l.	G. CaSO_4 per l.	G. NaCl per l.	G. CaSO_4 per l.
0.5	2.5	0.5	2.3	0.5	2.2	0.02	0.02
10.3	3.6	1.1	2.4	10.0	3.4	1.02	1.02
30.3	5.0	5.0	2.9	29.6	4.9	5.62	5.62
47.3	6.1	10.1	3.5	48.8	5.8	10.13	10.13
73.4	6.9	29.6	5.0	132.7	7.4	29.54	29.54
126.9	7.3	48.3	5.8	195.0	7.6	48.85	48.85
192.4	7.7	75.7	6.6			74.96	74.96
		131.6	7.1			128.77	128.77
		195.9	7.4			195.17	195.17

(Cameron, J. phys. Ch. 1901, 5. 562.)

1 l. sat. $\text{NaCl} + \text{Aq}$ at 25° containing 315
g. NaCl dissolves 5.52 g. CaSO_4 . (Cameron
and Brown, J. phys. Ch. 1905, 9. 214.)Solubility in $\text{NaCl} + \text{Aq}$.

G. NaCl per l. of $\text{NaCl} + \text{Aq}$	G. anhydrous CaSO_4 dissolved per litre	
	at 14°	at 30°
0.0	1.70	2.10
2.925	2.32	2.70
5.850	2.79	3.15
11.70	3.41	3.75
14.62	3.68	4.00
29.25	4.40	4.70
58.50	5.72	6.00
87.75	6.58	6.85
102.3	6.90	7.15
117.0	7.10	7.30
131.6	7.20	7.30
146.2	7.10	7.13
160.8	7.00	7.05
175.6	6.80	6.80
204.7	6.30	6.30
234.0	5.90	5.90
263.2	5.50	5.52
292.6	5.30	5.30

(d'Anselme, Bull. Soc. 1903, (3) 20. 373)

Solubility in $\text{NaCl} + \text{Aq}$.

G. NaCl in 100 cc. solution	G. $\text{CaSO}_4 + 2\text{H}_2\text{O}$
0.00	0.200 g
2.44 g.	0.635 g
4.77 g.	0.826 g
9.50 g.	1.056 g
14.22 g.	1.193 g
23.15 g.	1.275 g
31.30 g.	1.583 g

(Cloer, Bull. Soc. 1903, (3) 20. 167.)

Solubility in NaCl + Aq at t° .

When a sat. solution of NaCl is shaken with a mixture of solid NaCl and CaSO_4 , the calcium sulphate dissolved, calculated from the amount of CaO in solution, is greater than that calculated from the free acid in solution. Similar results are obtained when solid calcium sulphate alone is shaken with a sat. solution of NaCl.

In 100 g. of the solution			
Cl	CaSO_4 calc. from CaO	CaSO_4 calc. from SO_4	
15.253	0.4464	0.4334	
15.920	0.4477	0.4426	
15.987	0.4609	0.4542	
16.123	0.4938	0.4730	
16.270	0.5093	0.4832	
16.324	0.5305	0.5047	
16.381		0.5091	
16.459	0.5435	0.5749	
16.486	0.5578	0.5631	
16.524	0.5603	0.5587	
16.670	0.5399	0.5519	
17.128	0.4066	0.3414	

Smith, Bull. Soc. 1906, (3) 35. 780.)

In a temp. range from 25° – 80° CaSO_4 forms no double salt in solutions of NaCl. At any concentration with respect to NaCl, the maximum solubility occurs with 155 g. NaCl per l. and amounts to 7.3 g. CaSO_4 at 25° (Cameron, J. phys. Chem. 1907, 11.

See under Gypsum, p. 653.

Solubility of CaSO_4 in NaCl + Aq in contact with solid $\text{Ca}(\text{HCO}_3)_2$.

CaSO_4 l.	G. $\text{Ca}(\text{HCO}_3)_2$ per l.	G. NaCl per l.
298	0.0603	0.000
200	0.0724	3.628
160	0.0885	11.490
160	0.1006	39.620
140	0.0603	79.520
120	0.0563	121.900
160	0.0482	193.800
120	0.0402	267.600

Smith and Seidell, J. phys. Chem. 1901, 5. 653.)

$(\text{NH}_4)_2\text{SO}_4$ + Aq

in 287 pts. $(\text{NH}_4)_2\text{SO}_4$ + Aq (1.4, us, Z. anal. 30. 593.)

CaSO_4 is sol. in 327 ccm. $(\text{NH}_4)_2\text{SO}_4$ at 9° ; in 369 ccm. $1/7$ sat. $(\text{NH}_4)_2\text{SO}_4$ + 3.5° . (Fassbender)

Solubility in sat. $(\text{NH}_4)_2\text{SO}_4$, or Na_2SO_4 is less than in H_2O . (Droeze, B. 10. 330.)

Solubility in $(\text{NH}_4)_2\text{SO}_4$ + Aq at 25° .

g. per l. solution			g. per l.		
$(\text{NH}_4)_2\text{SO}_4$	CaSO_4	wt. of 100 cc. solution	$(\text{NH}_4)_2\text{SO}_4$	CaSO_4	wt. of 100 cc. solution
0.00	0.208	99.91	6.575	0.144	100.36
0.129	0.204	99.91	13.15	0.146	100.82
0.258	0.199	99.92	26.30	0.162	101.76
0.821	0.181	99.95	84.9	0.233	105.34
1.643	0.166	99.99	169.8	0.333	110.32
3.287	0.154	100.10	339.6	0.450	119.15

(Sullivan, J. Am. Chem. Soc. 1905, 27. 529.)

Solubility in $(\text{NH}_4)_2\text{SO}_4$ + Aq at 50° .

Sp. gr.	$(\text{NH}_4)_2\text{SO}_4$ per l.	CaSO_4 per l.	Solid phase
1.0026	0	2.168	
1.0113	15.65	1.609	
1.0440	30.67	1.750	
1.0819	91.6	2.542	
1.1108	160.4	3.402	
1.1385	221.6	4.068	$\text{CaSO}_4 + 2\text{H}_2\text{O}$
1.1653	280.6	4.690	
1.1972	340.6	5.084	
1.1972	415.6	5.336	
1.1964	416.5	5.354	
1.2043	428.4	4.632	
1.2187	479.4	3.524	$\text{CaSO}_4, (\text{NH}_4)_2\text{SO}_4 + 2\text{H}_2\text{O}$
1.2437	530.8	2.152	
1.2480	558.0	1.986	
1.2502	564.7	1.98	
1.2508	566.0	1.08	$(\text{NH}_4)_2\text{SO}_4$
1.2510	566.7	0	

(Bell and Taber, J. phys. Chem. 1906, 10. 120.)

Solubility of CaSO_4 in $(\text{NH}_4)_2\text{SO}_4$ + Aq at t° .

Excess of $(\text{NH}_4)_2\text{SO}_4$			Excess of CaSO_4		
t°	% CaSO_4	% $(\text{NH}_4)_2\text{SO}_4$	t°	% CaSO_4	% $(\text{NH}_4)_2\text{SO}_4$
6	0.1529	41.82	3	0.3782	36.62
40.5	0.1569	44.55	31	0.4070	35.50
58	0.1662	46.07	60	0.5083	34.97
78	0.1968	47.51	75	0.5898	34.86
100	0.2546	49.45	80	0.6108	34.88
			84	0.5725	32.40
			100	0.4895	25.97

(Barre, C. R. 1909, 148. 1605.)

The solubility of CaSO_4 in H_2O is considerably increased by the presence of $(\text{NH}_4)_2\text{SO}_4$.

but decreased by the presence of K_2SO_4 .
(Barre, C. R. 1909, 148. 1606.)

CuSO₄+Aq.
Solubility in CuSO₄+Aq at 25°.

Sp. gr. of the solution 25°/25°	g. CuSO ₄ per l.	g. CaSO ₄ per l.
1.002	1.144	2.068
1.005	3.564	1.986
1.007	6.048	1.944
1.009	7.279	1.858
1.016	14.814	1.760
1.021	19.729	1.736
1.030	29.543	1.688
1.041	39.407	1.718
1.051	49.382	1.744
1.061	58.880	1.782
1.098	97.950	1.931
1.146	146.725	2.048
1.192	196.021	2.076
1.218	224.916	2.088

(Bell and Taber, J. phys. Ch. 1907, 11. 637.)

MgSO₄+Aq.
Insol. in sat. MgSO₄+Aq.
1 g. CaSO₄ is sol. in 1162 ccm. $\frac{1}{10}$ sat. MgSO₄+Aq at 13.5°. (Fassbender, B. 9. 1360.)
Sol. in 635 pts. sat. MgSO₄+Aq at 19°. (Karsten.)
Absolutely insol. in sat. MgSO₄+Aq, and pptd. from aqueous solution by the addition of MgSO₄. (Droeze, B. 10. 340.)
1 l. $\frac{1}{10}$ sat. MgSO₄+Aq dissolves 0.86 g. CaSO₄+2H₂O. (Droeze.)

Solubility in MgSO₄+Aq at 25°.

g. per l.		Sp. gr. at 25°/25°	g. per l.		Sp. gr. at 25°/25°
MgSO ₄	CaSO ₄		MgSO ₄	CaSO ₄	
0.0	2.046	1.0032	149.67	1.597	1.1377
3.20	1.620	1.0055	165.7	1.549	1.1479
6.39	1.507	1.0090	171.2	1.474	1.1537
10.64	1.471	1.0118	198.8	1.422	1.1813
21.36	1.478	1.0226	232.1	1.254	1.2095
42.68	1.558	1.0419	265.6	1.070	1.2382
64.14	1.608	1.0626	298.0	1.860	1.2624
85.67	1.617	1.0833	330.6	0.647	1.2877
128.28	1.627	1.1190	355.0	0.501	1.3023

(Cameron and Bell, J. phys. Ch. 1906, 10. 210.)

K₂SO₄+Aq.
1 g. CaSO₄ is sol. in 2325 ccm. sat. K₂SO₄+Aq. at 13.5°; in 664 ccm. $\frac{1}{10}$ sat. K₂SO₄+Aq at 13.5°.

Solubility in K₂SO₄+Aq at 25°.

g. per l.		wt. of 1 cc. of solution
K ₂ SO ₄	CaSO ₄	
0.0	2.08	0.9961
4.88	1.60	1.0036
5.09	1.56	1.0038
9.85	1.45	1.0075
19.57	1.49	0.151
28.35	1.55	1.0229
30.66	1.57	1.0236
32.47*	1.58	

*Solid phase syngenite.
(Cameron and Breazeale, J. phys. Ch. 1904, 8. 335.)

Solubility in K₂SO₄+Aq. at 25°.
In 1000 g. of the solution
mole K₂SO₄ 3.223 mole CaSO₄ 0.223
(D'Ans, Z. anorg. 1909, 62. 151.)

Solubility of CaSO₄ in K₂SO₄+Aq at t°.

Excess of K ₂ SO ₄			Excess of CaSO ₄	
t°	% $\frac{CaSO_4}{Ca}$	%K ₂ SO ₄	% $\frac{CaSO_4}{Ca}$	%K ₂ SO ₄
0	0.1296	2.00	0.0229	6.99
18	0.1531	2.79	0.0271	9.81
51	0.1754	4.21	0.0300	14.18
80	0.1922	5.00	0.0349	17.55
99	0.1980	5.39	0.0371	19.70

(Barre, C. R. 1909, 148. 1606.)
Ag₂SO₄+Aq.
1 l. of the solution contains 2.31 g. CaSO₄+7.23 g. Ag₂SO₄=9.54 g. mixed salts at 17°. Sp. gr.=1.0083.
1 l. of the solution contains 2.61 g. CaSO₄+8.11 g. Ag₂SO₄=10.72 g. mixed salts at 25°. Sp. gr.=1.010. (Euler, Z. phys. Ch. 1904, 49. 313.)

Na₂SO₄+Aq.
1 g. CaSO₄ is sol. in 398 ccm. sat. Na₂SO₄+Aq at 10.5°.

Solubility of CaSO₄ in Na₂SO₄+Aq at 22°.

G. CaSO ₄ per l.	G. Na ₂ SO ₄ per l.
2.084	0.000
1.583	2.771
1.433	13.820
1.408	16.360
1.569	39.310
1.841	77.320
2.185	133.00
2.414	193.800
*2.578	*222.580

*Both CaSO₄ and Na₂SO₄ as solid phases in contact with the solution.
(Cameron and Seidell, J. phys. Chem. 1901, 5. 650.)

Solubility in Na ₂ SO ₄ +Aq at 25°.		
wt. of 1000 ccm. of solution grams	g. Na ₂ SO ₄ per l.	g. CaSO ₄ per l.
1001.26	2.390	1.650
1007.59	9.535	1.457
1011.45	14.132	1.388
1020.46	24.369	1.471
1031.48	36.979	1.563
1039.12	46.150	1.650
1079.47	94.220	1.980
1096.47	115.084	2.096
1142.66	146.612	2.234
1176.47	205.105	2.503
1212.00	257.100	2.650

Cameron and Breazeale, J. phys. Chem. 1904, 8. 340.)

1 l. sat. Na₂SO₄+Aq at 25° containing 254.09, Na₂SO₄ dissolves 2.58 g. CaSO₄. Cameron and Brown, J. phys. Ch. 1905, 9 214.)

Hydration is retarded by dil. solutions and accelerated by conc. solution of sodium, potassium, ammonium and magnesium sulphates. (Rohland, Z. Elektrochem. 1908, 14. 422.)

More than 10 times as much CaSO₄ dissolves in sat. Na₂S₂O₃+Aq as in H₂O. (Diehl.

Insol. in alcohol. of 0.905 sp. gr. or less. (Anthon, J. pr. 14. 125.)

Solubility in 10% alcohol=0.0970 g. CaSO₄ per 100 g. solution. (Magnanini, Gazz. Ch. it. 1901, 31. (2) 544.)

Sol. in dil. alcoholic solutions of NH₄NO₃, KNO₃, NaNO₃, NH₄Cl, KCl, and NaCl. (Margueritte, C. R. 38. 308.)

Sol. to considerable extent in NH₄C₂H₃O₂+Aq, especially if freshly pptd. More sol. in NH₄C₂H₃O₂+Aq than in NH₄Cl+Aq. (Weppen, J. pr. 11. 182.)

More sol. in NH₄C₂H₃O₂+Aq than in other NH₄ salts. (Cohn, J. pr. (2) 35. 43.)

More sol. in NaC₂H₃O₂+Aq or KCl+Aq than in H₂O. (Mulder.)

Solubility in N/200 potassium hydrogen tartrate+Aq=0.2323 g. CaSO₄ per 100 g. solution. (Magnanini, Gazz. ch. it. 1901, 31. (2) 544.)

72.61 millimols. per l. of CaSO₄+2H₂O are sol. at 25° in ammonium citrate+Aq (concentration=0.5 millimols. per l.)

36.39 millimols per l. of CaSO₄+2H₂O are sol. at 25° in sodium citrate+Aq. (Concentration=0.25 millimols per l. (Rindell, Z. phys. Ch. 1910, 70. 452.)

100 pts. glycerine dissolve 0.957 pt. CaSO₄+2H₂O, and solubility increases with the temp. (Asselin, C. R. 76. 884.)

100 g. glycerine (sp. gr. 1.256) dissolve 5.17 g. CuSO₄ at 15-16°. (Ossendowski, Pharm. J. 1907, 79. 575.)

Solubility in 10% alcoholic N/200 KHC₂H₃O₆+Aq=0.0866 g. CaSO₄ per 100 g. solution.

Solubility in N/200 KHC₂H₃O₆+Aq+5% tartaric acid=0.2556 g. CaSO₄ per 100 g. solution.

Solubility in 10% alcoholic N/400 KHC₂H₃O₆+5% tartaric acid=0.1086 g. CaSO₄ in 100 g. solution. (Magnanini.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethylacetate. (Naumann, B. 1910, 43. 314.)

Solubility in sugar+Aq at t°.

% sugar	G. CaSO ₄ dissolved in 1 l. sugar solutions					
	30°	40°	50°	60°	70°	80°
0	...	2.157	1.730	1.730	1.652	1.710
10	2.041	1.730	1.730	1.574	1.574	1.613
20	1.808	1.652	1.419	1.380	1.419	1.263
27	1.550	1.438	1.361	1.283	1.283	0.972
35	1.263	1.050	1.088	1.108	0.914	...
42	1.030	...	0.777	0.816	0.855	0.729
49	...	0.564	0.739	0.564	0.603	0.486
55	...	0.486	0.505	0.486	0.369	0.330

(Stolle, Z. Ver. Zuckerind, 1900, 50. 331).

Min. Anhydrite.
+2H₂O. Min. Gypsum.

Gypsum. A sat. aq. solution of gypsum of particles not less than 2μ contains 2.085 g. CaSO₄ per litre at 25°.

A sat. aq. solution of gypsum of particles not smaller than 0.3μ contains 2.476 g. CaSO₄ per liter at 25°. (μ=0.0001 cm.) (Hulett and Allen, Z. phys. Ch. 1901, 37. 391 and 393.)

Solubility in H₂O at t°.

t°	g. CaSO ₄ in 100 ccm. of the solution	Density of the solution at t°
0	0.17590	1.001970
10	0.19285	1.001727
18	0.20160	1.000590
25	0.20805	0.999109
30	0.20905	0.997891
35	0.20960	0.996122
40	0.20970	0.994390
45	0.20835	0.992370
55	0.20095	0.987960
65.3	0.19320	0.982560
75	0.18475	0.977724
100	0.16195	...

(Hulett and Allen, J. Am. Chem. Soc. 1902, 24. 674.)

1 l. H₂O dissolves 2.13 g. CaSO₄+2H₂O at 25°. (Euler, Z. phys. Ch. 1904, 49. 314.)

2023 mg. are dissolved in 1 l. of sat. solution at 18°. (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

1 l. H₂O dissolves 2.267 g. CaSO₄+2H₂O at 0°; 2.684 g. at 35°; 2.662 g. at 50°; and 2.155 g. at 100°. (Cavazzi, C. C. 1905, 1. 1693.)

Calcium sodium sulphate, $\text{CaNa}_2(\text{SO}_4)_2$.

Min. *Glauberite*. Gradually sol. in H_2O , but crystals of $\text{CaSO}_4 + 2\text{H}_2\text{O}$ soon separate out. (Fritzsche.)

Insol. in alcohol, and conc. $\text{NaC}_2\text{H}_3\text{O}_2 + \text{aq}$; decomp. by H_2O . (Folkhard, C. N. 43.)

$\text{CaNa}_2(\text{SO}_4)_2 + 2\text{H}_2\text{O}$. Decomp. by H_2O . (Fritzsche.)

Calcium titanium sulphate, $\text{CaSO}_4 \cdot \text{Ti}(\text{SO}_4)_2$.

Ppt.; decomp. by H_2O giving titanous acid. (Weinland, Z. anorg. 1907, 54. 254.)

Calcium uranium sulphate.

Min. *Uranochalcite*.

Min. *Medjidite*. Easily sol. in dil. $\text{HCl} + \text{aq}$.

Cerous sulphate, $\text{Ce}_2(\text{SO}_4)_3$.

Anhydrous cerous sulphate is much more sol. in H_2O than the hydrated salt.

Easily sol. in cold H_2O if added thereto in small amounts. If large amount of $\text{Ce}_2(\text{SO}_4)_3$ is treated with a little H_2O it hardens with evolution of heat, and becomes very difficultly soluble. 100 pts. H_2O dissolve 161 pts. $\text{Ce}_2(\text{SO}_4)_3$ at 0° and 17.86 pts. at 19° .

$\text{Ce}_2(\text{SO}_4)_3 + \text{Aq}$ sat. in cold deposits $\text{Ce}_2(\text{SO}_4)_3$ at 75° , and only 2.25 pts. remain in solution at 100° . (Jolin, Bull. Soc. (2) 21. 536.)

100 pts. H_2O dissolve 8.31 pts. $\text{Ce}_2(\text{SO}_4)_3$ at 20° ; 8.08 pts. at 45° ; 4.95 pts. at 60° ; 0.504 pt. at 100° . (Bühlig, J. pr. (2) 12. 240.)

60 pts. anhydrous salt dissolve quickly at $1-3^\circ$ in 100 pts. H_2O .

At 15° the solution solidifies, and the mother liquor contains only 27.88% $\text{Ce}_2(\text{SO}_4)_3$. At 15° the maximum attainable strength is 31.62% $\text{Ce}_2(\text{SO}_4)_3$. (Brauner, Chem. Soc. 53. 357.)

100 pts. H_2O dissolve 10.747 pts. $\text{Ce}_2(\text{SO}_4)_3$ at 16° ; 9.648 pts. at 19° ; 6.949 pts. at 33° .

The solubility of $\text{Ce}_2(\text{SO}_4)_3$ in H_2O is diminished by the addition of $(\text{NH}_4)_2\text{SO}_4$, K_2SO_4 , or Na_2SO_4 . (Barre, C. R. 1910, 151. 872.)

Sp. gr. of $\text{Ce}_2(\text{SO}_4)_3 + \text{Aq}$ was found to be constant whether $\text{Ce}_2(\text{SO}_4)_3$ or $\text{Ce}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ was used. The following results were obtained at 15° .

Pts. $\text{Ce}_2(\text{SO}_4)_3$ to 100 pts. H_2O	Sp. gr.	Pts. $\text{Ce}_2(\text{SO}_4)_3$ to 100 pts. H_2O	Sp. gr.
3.17	1.03005	12.66	1.11917
6.11	1.05812	14.56	1.13665
8.35	1.07910	15.64	1.14623
9.61	1.09085	21.19	1.19640
10.55	1.09939	31.62	1.28778
11.66	1.10987

(Brauner, Chem. Soc. 53. 357.)

4.5 pts. $\text{Ce}_2(\text{SO}_4)_3$ dissolve in 100 pts. H_2SO_4 . (Wyruboff, Bull. Soc. (3) 2. 745.)

Solubility in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ at 16° .

Per 100 pts. H_2O	
$(\text{NH}_4)_2\text{SO}_4$	$\text{Ce}_2(\text{SO}_4)_3$
0.00	10.747
3.464	1.026
9.323	0.782
19.240	0.748
29.552	0.701
45.616	0.497
55.083	0.194
63.920	0.090
72.838	0.035

(Barre, A. ch. 1911, (8) 24. 252.)

Solubility in $\text{Na}_2\text{SO}_4 + \text{Aq}$ at $19^\circ-20^\circ$.

Per 100 pts. H_2O	
Na_2SO_4	$\text{Ce}_2(\text{SO}_4)_3$
0.00	9.64
0.328	0.637
0.684	0.259
1.091	0.0937
1.392	0.057
1.699	0.0303
2.640	0.012
3.589	0.0065
5.660	0.0046
7.710	0.0037

(Barre, A. ch. 1911, (8) 24. 251.)

Solubility in $\text{K}_2\text{SO}_4 + \text{Aq}$ at 16° .

Per 100 pts. H_2O	
K_2SO_4	$\text{Ce}_2(\text{SO}_4)_3$
0.00	10.747
0.178	0.956
0.510	0.432
0.726	0.250
1.290	0.0419

(Barre, A. ch. 1911, (8) 24. 248.)

$+4\text{H}_2\text{O}$. 100 g. H_2O dissolve at:

35° 40° 50° 57°
8.5 6.04 3.43 2.34 g. $\text{Ce}_2(\text{SO}_4)_3$.

65° 70° 82° 100.5° bpt. of sat. solution.
1.883 1.38 1.01 0.43 g. $\text{Ce}_2(\text{SO}_4)_3$.

(Koppel, Z. anorg. 1904, 41. 399.)

+5H₂O.
100 pts. H₂O dissolve pts. Ce₂(SO₄)₃ at t°.

t°	Pts. Ce ₂ (SO ₄) ₃
100	0.775
80	1.70
60	3.45
50	5.56
40	8.20

(Muthmann and Rolig, Z. anorg. 1898, 16. 456.)

100 g. H₂O dissolve at:
45° 60° 70°
8.833 3.247 1.929 g. Ce₂(SO₄)₃.
80° 90° 100.5° bpt. of sat. solution.
1.207 0.8355 0.469 g. Ce₂(SO₄)₃.

Muthmann and Rolig's determinations are inaccurate. (Koppel.)
+8H₂O. 100 pts. H₂O dissolve 14.92 pts. Ce₂(SO₄)₃ at 20° from Ce₂(SO₄)₃+8H₂O, (Jolin.)

100 pts. H₂O dissolve pts. Ce₂(SO₄)₃ at t°.

t°	Pts. Ce ₂ (SO ₄) ₃	t°	Pts. Ce ₂ (SO ₄) ₃
0	19.10	50	12.48
18	17.32	60	9.40
30	16.13	70	4.24

(Muthmann and Rolig.)

100 g. H₂O dissolve at:
0° 15° 20.4°
10.09 11.06 9.525 g. Ce₂(SO₄)₃,
30° 40° 50° 60°
7.388 5.947 4.785 4.064 g. Ce₂(SO₄)₃.

Previous determinations are inaccurate. (Koppel, Z. anorg. 1904, 41. 395.)
100 g. sat. solution at 25° contain 7.60 g. anhydrous salt. (Wirth, Z. anorg. 76. 174.)

Solubility in H₂SO₄+Aq at 25°. Solid phase Ce₂(SO₄)₃+8H₂O.

Normality H ₂ SO ₄	In 100 g. of the liquid are dissolved	
	g. Ce ₂ O ₃	g. Ce ₂ (SO ₄) ₃
0	4.604	7.60
0.1	4.615	7.618
1.1	3.64	6.00
2.16	3.04	5.018
4.32	2.0	3.301
6.685	0.9115	1.505
9.68	0.4339	0.733
15.15	0.145	0.239

(Wirth, Z. anorg. 1912, 76. 191.)

+9H₂O. 100 pts. H₂O dissolve 17.52 pts. Ce₂(SO₄)₃ from Ce₂(SO₄)₃+9H₂O. (Brauner.

100 g. H₂O dissolve at:
0° 15° 21° 30° 31.2°
20.98 11.87 9.725 7.353 7.185 g. Ce₂(SO₄)₃,
31.6° 45° 50° 60° 65°
7.164 5.13 4.673 3.88 3.595 g. Ce₂(SO₄)₃.
(Koppel.)

+12H₂O.
100 pts. H₂O dissolve pts. Ce₂(SO₄)₃ at t°.

t°	Pts. Ce ₂ (SO ₄) ₃
0	21.40
18	18.44
25	16.22

(Muthmann and Rolig, Z. anorg. 1898, 16. 457.)

100 g. H₂O dissolve at:
0° 18.8° 19.2°
16.56 17.52 17.70 g. Ce₂(SO₄)₃.
Previous determinations are inaccurate (Koppel.)

Ceroceric sulphate, Ce₂(SO₄)₃, 2Ce(SO₄)₂+24H₂O.
Decomp. by H₂O. Sol. in HCl+Aq with decomp. (Mendelejeff, A. 168. 45.)
Ce₂(SO₄)₃, 3Ce(SO₄)₂+31H₂O. (Jolin.)

Ceric sulphate, basic, CeO₂, SO₃+2H₂O.
Very sl. sol. in H₂O.
Sol. in 2500 pts. H₂O. (Mosander.)
Boiling H₂O gradually dissolves out H₂SO₄. (Erk.)
Sol. in acids.
8CeO₂, 7SO₃+12H₂O; 8CeO₂, 7SO₃+15H₂O; 6CeO₂, 5SO₃+5H₂O; 4CeO₂, 3SO₃+7H₂O; and 3Ce(SO₄)₂, 5Ce(OH)₄. All are insol. ppts.

Ceric sulphate, Ce(SO₄)₂.
Anhydrous. Very slowly sol. in cold, more rapidly in hot H₂O. When solution has once begun, almost unlimited quantities may be dissolved. Insol. in conc. H₂SO₄. (Meyer, B. 1904, 37. 144.)
+4H₂O. Sol. in H₂O with immediate decomp. (Rammelsberg.)
Decomp. by H₂O. (Muthmann, B. 1900, 33. 1764.)

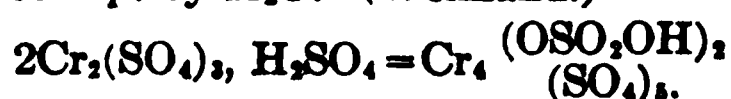
Cerous hydrogen sulphate, Ce₂(SO₄)₃, 3H₂SO₄.
Decomp. by H₂O. (Wyruboff, Bull. Soc. (3) 2. 745; Brauner, Z. anorg. 1904, 33. 329.)

Ceroceric hydrogen sulphate, Ce₂H(SO₄)₄+13H₂O.
Sol. in H₂O. Forms very supersat. solutions.
Solubility in H₂SO₄ decreases with increase in concentration of the acid. (Meyer, B. 1904, 37. 146.)

- Cerous potassium sulphate**, $\text{Ce}_2(\text{SO}_4)_3, \text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}$
 Sl. sol. in H_2O ; insol. in sat. $\text{K}_2\text{SO}_4 + \text{Aq}$. (Csudnowicz, J. pr. 80. 26.)
 $2\text{Ce}_2(\text{SO}_4)_3, 3\text{K}_2\text{SO}_4$. As above. (Hermann, J. pr. 30. 188.)
 $+ 8\text{H}_2\text{O}$. (Barre, A. ch. 1911, (8) 24. 249.)
 $\text{Ce}_2(\text{SO}_4)_3, 2\text{K}_2\text{SO}_4 + 3\text{H}_2\text{O}$. As above. (Jolin.)
 $\text{Ce}_2(\text{SO}_4)_3, 3\text{K}_2\text{SO}_4$. Sol. in about 56 pts. H_2O at $9-20^\circ$. Easily sol. in acidified H_2O . Nearly insol. in sat. $\text{K}_2\text{SO}_4 + \text{Aq}$. (Jolin.)
 $\text{Ce}_2(\text{SO}_4)_3, 5\text{K}_2\text{SO}_4$. Insol. in $\text{K}_2\text{SO}_4 + \text{Aq}$. (Barre, l.c.)
- Ceric potassium sulphate**, $\text{Ce}(\text{SO}_4)_2, 2\text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}$
 Sl. sol. in H_2O with decomp. Insol. in sat. $\text{K}_2\text{SO}_4 + \text{Aq}$.
- Ceric silver sulphate**, $10\text{Ce}(\text{SO}_4)_2, 6\text{Ag}_2\text{SO}_4$
 Only sl. sol. in cold H_2O ; decomp. by hot H_2O in which it is readily sol. (Pozzi-Escot, C. R. 1913, 156. 1074.)
- Cerous sodium sulphate**, $\text{Ce}_2(\text{SO}_4)_3, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$
 Very sl. sol. in H_2O , and still less in $\text{Na}_2\text{SO}_4 + \text{Aq}$. 100 ccm. sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ dissolve an amount corresponding to 6.2 mg. Ce_2O_3 . (Jolin.)
 Sl. sol. in $\text{HCl} + \text{Aq}$. (Csudnowicz.)
- Cerous thallous sulphate**, $\text{Ce}_2(\text{SO}_4)_3, 3\text{Tl}_2\text{SO}_4$
 Ppt.
 $\text{Ce}_2(\text{SO}_4)_3, \text{Tl}_2\text{SO}_4 + 2\text{H}_2\text{O}$. Sol. in H_2O . (Zehiesche, J. pr. 107. 98.)
 $+ 4\text{H}_2\text{O}$. Very sl. sol. in cold, somewhat more in warm H_2O . (Wyruboff, Bull. Soc. Min. 14. 83.)
- Cerous tin (stannic) hydrogen sulphate**, $\text{CeHSn}(\text{SO}_4)_4$
 Decomp. by H_2O . Sol. in very dil. HCl . (Weinland, Z. anorg. 1907, 54. 251.)
- Chromous sulphate**, $\text{CrSO}_4 + 7\text{H}_2\text{O}$
 100 pts. H_2O dissolve 12.35 pts. $\text{CrSO}_4 + 7\text{H}_2\text{O}$. Aqueous solution can be boiled without decomp. Sl. sol. in alcohol.
 $+ \text{H}_2\text{O}$. (Moissan, Bull. Soc. 37. 296.)
- Chromic sulphate, basic**, $3\text{Cr}_2\text{O}_3, 2\text{SO}_3 + 12\text{H}_2\text{O} = 2\text{Cr}_2(\text{SO}_4)(\text{OH})_4, \text{Cr}_2(\text{OH})_6 + 5\text{H}_2\text{O}$
 Insol. in H_2O . Sol. in acids. Slowly decomp. by $\text{KOH} + \text{Aq}$ or $\text{K}_2\text{CO}_3 + \text{Aq}$.
 $5\text{Cr}_2\text{O}_3, 3\text{SO}_3$. Sol. in H_2O . (Recours, C. R. 112. 1439.)
 $\text{Cr}_2\text{O}_3, \text{SO}_3 = \text{Cr}_2\text{O}_3(\text{SO}_3)$. Ppt. (Schiff, A. 124. 167.)
 $+ 10\text{H}_2\text{O}$ or $[\text{Cr}(\text{OH})_2(\text{OH}_2)_4]_2\text{SO}_4$. Nearly insol. in H_2O . (Werner, B. 1908, 41. 3451.)
 $5\text{Cr}_2\text{O}_3, 8\text{SO}_3$ (?). (Siewert, A. 126. 97.)
 $\text{Cr}_2\text{O}_3, 2\text{SO}_3 = \text{Cr}_2\text{O}(\text{SO}_3)_2$. Easily sol. in a little H_2O , but a precipitate is thrown down by further addition of H_2O , which redissolves on evaporation.
 $5\text{Cr}_2\text{O}_3, 12\text{SO}_3$ (?). (Siewert.)
 $2\text{Cr}_2\text{O}_3, 5\text{SO}_3 + 15\text{H}_2\text{O}$. Sol. in H_2O ; insol. in alcohol and acetone by which it is ppt. from aqueous solution. (Nicolardot, C. R. 1907, 145. 1338.)
- Chromic sulphate**, $\text{Cr}_2(\text{SO}_4)_3$
Anhydrous. Insol. in H_2O , HNO_3 , HCl , H_2SO_4 , aqua regia, and $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by boiling caustic alkalies, and slowly by alkali carbonates $+ \text{Aq}$. (Schrötter.) According to Traube (A. 71. 92) and Siewert (A. 126. 94), Schrötter's salt is an acid sulphate, $\text{Cr}_2(\text{SO}_4)_3(\text{OSO}_3\text{OH})_2 = 2\text{Cr}_2(\text{SO}_4)_3, \text{H}_2\text{SO}_4$. According to Etard (Bull. Soc. (2) 31. 200) both salts exist, and formula of above salt is $\text{Cr}_2(\text{SO}_4)_3\text{Cr}_2$. Formula is $2[(\text{Cr}_2\text{O}_3)_2, (\text{SO}_3)_4], 17\text{H}_2\text{SO}_4$ (?). (Cross and Higgins, Chem. Soc. 41. 118.)
 Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)
 $+ 6\text{H}_2\text{O}$ (?). *Green modification*. Readily sol. in H_2O or alcohol. Sol. in conc. H_2SO_4 . H_2O solution is converted into the violet modification by standing 3-4 weeks. (Schrötter.)
 $+ 11\text{H}_2\text{O}$ (?). Extremely deliquescent; becomes liquid in moist air in 2 minutes. Not pptd. by $\text{BaCl}_2 + \text{Aq}$. (Recours, C. R. 113. 857.)
 $+ 18\text{H}_2\text{O}$. *Violet modification*. Sol. in 0.833 pt. H_2O at 20° . When the H_2O solution is heated to $65-70^\circ$ it begins to be converted into the green modification. This conversion is also brought about by cold HNO_3 , H_2SO_4 , PCl_5 . (Etard, C. R. 84. 1090.)
 Sp. gr. of aqueous solution of violet modification of $\text{Cr}_2(\text{SO}_4)_3$ containing:
- | | | |
|--------|--------|--------|
| 5 | 10 | 20% |
| 1.0275 | 1.0560 | 1.1150 |
| 30 | 40 | 50% |
| 1.1785 | 1.2480 | 1.3250 |
- Sp. gr. of aqueous solution of green modification of $\text{Cr}_2(\text{SO}_4)_3$ containing:
- | | | |
|--------|--------|--------|
| 10 | 20 | 30% |
| 1.0510 | 1.1070 | 1.1680 |
| 40 | 50 | 60% |
| 1.2340 | 1.3055 | 1.3825 |
| 70 | 80% | |
| 1.4650 | 1.5535 | |
- (Gerlach, Z. anal. 28. 494.)
- See also Chromosulphuric acid.
- Chromic hydrogen sulphate**, $\text{Cr}_2(\text{SO}_4)_3, \text{H}_2\text{SO}_4 + 16\text{H}_2\text{O}$
Two modifications.
 a. *Violet*. Decomp. by H_2O .
 b. *Green*. Obtained from violet modification on heating. Sol. in H_2O . (Weinland, Z. anorg. 1906, 49. 157.)

+24H₂O. Decomp. by alcohol, giving the normal sulphate. (Weinland.)

Cr₂(SO₄)₃, 2H₂SO₄+18H₂O. Hygroscopic. Decomp. by H₂O. (Weinland.)



Correct composition of Cr₂(SO₄)₃ (Traube), which see.

See also Chromosulphuric acid.

Chromic cupric sulphate, Cr₂(SO₄)₃, 2CuSO₄, H₂SO₄.

Insol. in H₂O, but gradually decomp. thereby. (Étard, C. R. 87. 602.)



Insol. in H₂O. (Recoura, C. R. 1893, 117. 39.)

Chromous hydrazine sulphate, CrSO₄, 2N₂H₄, H₂SO₄.

Only sl. sol. in H₂O. Sol. in acids. (Traube B. 1913, 46. 1507.)

Chromic hydroxylamine sulphate, Cr₂(SO₄)₃, (NH₂OH)₂SO₄+24H₂O.

Sol. in H₂O. (Meyeringh.)

Chromic iron (ferrous) sulphate, Cr₂(SO₄)₃, 2FeSO₄, H₂SO₄+2H₂O.

As above. (Étard, l.c.)

Chromic iron (ferric) sulphate, Cr₂(SO₄)₃, Fe₂(SO₄)₃.

Insol. in H₂O. (Étard, C. R. 86. 1399.)

Cr₂(SO₄)₃, Fe₂(SO₄)₃, H₂SO₄. Insol. in H₂O. (Étard.)

Chromic lithium sulphate, Cr₂(SO₄)₃, 3Li₂SO₄.

Resembles the corresponding K salt. (Wernicke.)

Chromic manganous sulphate, Cr₂(SO₄)₃, 3MnSO₄.

(Étard, C. R. 86. 1402.)

Chromic manganic sulphate, Cr₂(SO₄)₃, Mn₂(SO₄)₃.

Insol. in H₂O. (Étard, C. R. 86. 1399.)

Cr₂(SO₄)₃, Mn₂(SO₄)₃, 2H₂SO₄. Sl. deliquescent. Sol. in H₂O with decomp. (Étard.)

Chromic nickel sulphate, Cr₂(SO₄)₃, NiSO₄, 2H₂SO₄+3H₂O.

Insol. in H₂O, but gradually decomp. thereby. (Étard, C. R. 87. 602.)

Chromous potassium sulphate, CrSO₄, K₂SO₄+6H₂O.

Sol. in H₂O; less sol. in alcohol. (Peligot, A. ch. (3) 12. 546.)

Chromic potassium sulphate, K₂Cr₂(SO₄)₄.

Anhydrous. a. Sol. in H₂O when not heated over 350°.

β. Insol. in cold H₂O and cold acids. When ignited is insol. in hot H₂O and acids, except slightly in boiling conc. H₂SO₄. (Fischer.)

+2H₂O (?). Insol. in cold H₂O or dil. acids. Sol. by long boiling with H₂O, and more quickly when HCl is added. (Hertwig.)

+4H₂O. Is potassium chromosulphate, which see.

+24H₂O. *Chrome-alum*. *Violet modification*. Efflorescent at 29°. Sol. in 6-7 pt. cold H₂O. When the H₂O solution is heated to 60-70° it is partially decomp. into a green modification, which is more sol. in H₂O. The green modification on standing in H₂O solution is very slowly converted back into violet modification. The green modification may also be formed by heating dry salt to 100°, at which temp. it melts in its crystal H₂O. When all crystal H₂O has been expelled at 300-350°, it still dissolves in hot H₂O, but when heated above 350° it becomes insol. in H₂O. (Löwel, A. ch. (3) 44. 313.)

125.1 g. anhydrous, or 243.9 g. hydrated salt, or 0.441 g. mols. anhydrous salt are sol. in 1 l. H₂O at 25°. (Locke, Am. Ch. J. 1901, 26. 175.)

Melts in crystal H₂O at 89°. (Tilden Chem. Soc. 45. 409.)

Sp. gr. of aqueous solution of violet modification at 15° containing:

5	10	15%	K ₂ Cr ₂ (SO ₄) ₄ +24H ₂ O.
1.02725	1.05500	1.08350	

Sp. gr. of sat. solution at 15° = 1.0965.

Sp. gr. of aqueous solution of green modification at 15° containing:

10	20	30%	K ₂ Cr ₂ (SO ₄) ₄ +24H ₂ O.
1.050	1.103	1.161	

40	50	60%	K ₂ Cr ₂ (SO ₄) ₄ +24H ₂ O.
1.225	1.295	1.371	

70	80	90%	K ₂ Cr ₂ (SO ₄) ₄ +24H ₂ O.
1.453	1.541	1.635	

(Gerlach, Z. anal. 28. 497.)

Sp. gr. of chrome-alum solutions at 15° containing:

5	10	15	20	25	% salt.
1.0174	1.0342	1.0524	1.0746	1.1004	

30	35	40	45	50	% salt.
1.1274	1.1572	1.1896	1.2352	1.2894	

55	60	65	70	% salt.
1.3704	1.4566	1.5462	1.6362	

(Franz, J. pr. (2) 5. 298.)

Insol. in alcohol.

3K₂SO₄, Cr₂(SO₄)₃. Insol. in H₂O, acids, or dil. alkalies. Decomp. by boiling with conc. KOH+Aq. (Wernicke, Pogg. 159. 576.)

Chromic rubidium sulphate, Rb₂Cr₂(SO₄)₄+24H₂O.

Sol. in H₂O. (Peterson.)

Solubility in H ₂ O.		
np.	G. anhydrous salt per l.	G. mols. of anhydrous salt per l.
0°	25.7	0.079
10°	31.7	0.093
20°	41.1	0.128
30°	59.7	0.181

sol. in crystal H₂O at 107°.
(Locke, Am. Ch. J. 1901, 26. 180.)

Calcium sodium sulphate, Na₂Cr₂(SO₄)₄ + H₂O.
Calcium chromosulphate, which see.
H₂O. More efflorescent than K or Na salt. Sol. in H₂O, and properties resemble the corresponding K salt.
Na₂Cr₂(SO₄)₄, 3Na₂SO₄. Resembles the corresponding K salt.

Calcium thallous sulphate, Tl₂Cr₂(SO₄)₄ + H₂O.
10 mols. of anhydrous salt are sol. in 100 H₂O at 25°. 1 l. H₂O dissolves 104.8 g. anhydrous or 163.8 g. hydrated salt at 107°. Melts in crystal H₂O at 92°. (Locke, Am. Ch. J. 1901, 26. 175.)

Calcium sulphate chloride, Cr₂(SO₄)₂Cl₂ + 2H₂O.
Slightly hygroscopic. Sol. in H₂O. (Schiff, 176.)
Na₂Cr₂(SO₄)₄, 5H₂O·Cl. Sol. in H₂O. (Weinmann, anorg. 1908, 58. 176.)

Chromium sulphate, (CrO₂)₂SO₄.
Decomposed by H₂O. (Pictet and Karl, Bull. Chim. 1888, (4) 3. 1114.)

Cobaltous sulphate, basic.
Insol. in H₂O. (Berzelius.)
CoSO₄·SO₃ + 10H₂O. (Athanasesco, C. R. 1891, 1.)
CoSO₄·SO₃ + 4H₂O. Ppt. Very sl. sol. in H₂O. (Habermann, M. Ch. 5. 432.)

Cobaltous sulphate, CoSO₄.
pts. H₂O dissolve at:
0° 20° 24° 29°
1.5 36.4 38.9 40 pts. anhydrous salt,
4° 50° 60° 70°
1.4 55.2 60.4 65.7 pts. anhydrous salt.
(Tobler, A. 95. 193.)

pts. H₂O at 11–14° dissolve 23.88 pts. anhydrous salt. (v. Hauer, J. pr. 103. 114.)

Solubility in 100 pts. H₂O at t°, using CoSO₄ + 7H₂O.

t°	Pts. CoSO ₄	t°	Pts. CoSO ₄	t°	Pts. CoSO ₄
0	24.6	36	43.5	72	65.0
1	25.0	37	44.0	73	65.6
2	25.5	38	44.6	74	66.2
3	26.0	39	45.2	75	66.8
4	26.5	40	45.8	76	67.4
5	27.0	41	46.4	77	68.0
6	27.5	42	47.0	78	68.6
7	28.0	43	47.6	79	69.2
8	28.5	44	48.2	80	69.8
9	29.0	45	48.8	81	70.4
10	29.5	46	49.4	82	71.0
11	30.0	47	50.0	83	71.6
12	30.5	48	50.6	84	72.2
13	31.0	49	51.2	85	72.8
14	31.5	50	51.8	86	73.4
15	32.0	51	52.4	87	74.0
16	32.5	52	53.0	88	74.6
17	33.0	53	53.6	89	75.2
18	33.5	54	54.2	90	75.9
19	34.0	55	54.8	91	76.6
20	34.5	56	55.4	92	77.2
21	35.1	57	56.0	93	77.9
22	35.6	58	56.6	94	78.6
23	36.2	59	57.2	95	79.2
24	36.8	60	57.8	96	79.9
25	37.4	61	58.4	97	80.6
26	38.0	62	59.0	98	81.3
27	38.5	63	59.6	99	81.9
28	39.1	64	60.2	100	82.6
29	39.6	65	60.8	101	83.3
30	40.2	66	61.4	102	83.9
31	40.7	67	62.0	103	84.6
32	41.3	68	62.6	104	85.3
33	41.8	69	63.2	105	86.0
34	42.4	70	63.8	106	86.7
35	42.9	71	64.4	106.4	86.9

(Mulder, calculated from his own and Tobler's determinations, Scheik. Verhandel. 1864. 68.)
100 g. H₂O dissolve 37.8 g. CoSO₄ at 25°. (Wagner, Z. phys. Ch. 1910, 71. 430.)
See also +7H₂O.

Sp. gr. of CoSO₄ + Aq at t°. S = pts. CoSO₄ in 100 pts. solution; S₁ = mols. CoSO₄ in 100 mols. of solution.

S	S ₁	Sp. gr.
6.8910	0.852	1.0765
5.8140	0.711	1.0641
4.7095	0.570	1.0517
3.5792	0.429	1.0392
2.4273	0.288	1.0263
1.2099	0.141	1.0131

(Charpy, A. ch. (6) 28. 28.)

Cobaltous magnesium potassium sulphate, $\text{CoSO}_4, \text{MgSO}_4, \text{K}_2\text{SO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Does not exist. (Aston and Pickering, Chem. Soc. 49. 123.)

Cobaltous manganous potassium sulphate, $\text{CoSO}_4, \text{MnSO}_4, 2\text{K}_2\text{SO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Cobaltous nickel potassium sulphate, $\text{CoSO}_4, \text{NiSO}_4, 2\text{K}_2\text{SO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Does not exist. (Thomson, Rep. Brit. Assn. Adv. Sci. 1877. 209.)

Cobaltous potassium sulphate, $\text{CoSO}_4, \text{K}_2\text{SO}_4 + 6\text{H}_2\text{O}$.

Less sol. in H_2O than CoSO_4 .

100 pts. H_2O dissolve at:

0° 12° 15° 20° 25°
19.1 30 32.5 39.4 45.3 pts. anhydrous salt,

30° 35° 40° 49°
51.9 55.4 64.6 81.3 pts. anhydrous salt.
(Tobler, A. 96. 126.)

100 pts. saturated solution contain at:

20° 40° 60° 80°
14 19.5 24.4 31.8 pts. anhydrous salt.
(v. Hauer, J. pr. 74. 433.)

1 l. H_2O dissolves 128.8 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Cobaltic potassium sulphate, $\text{K}_2\text{Co}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Marshall, Chem. Soc. 59. 760.)

Cobaltous potassium zinc sulphate, $\text{CoSO}_4, 2\text{K}_2\text{SO}_4, \text{ZnSO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Cobaltous rubidium sulphate, $\text{CoSO}_4, \text{Rb}_2\text{SO}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Tutton.)

1 l. H_2O dissolves 92.8 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Cobaltic rubidium sulphate, $\text{Rb}_2\text{Co}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in dil. HCl and H_2SO_4 . Decomp. by conc. HCl or H_2SO_4 . (Howe and O'Neal, J. Am. Chem. Soc. 1898, 20. 762.)

Melts in crystal H_2O at 47°. (Locke, Am. Ch. J. 1901, 26. 183.)

Cobaltous sodium sulphate, $\text{CoNa}_2(\text{SO}_4)_2 + 4\text{H}_2\text{O}$.

Solubility of $\text{CoNa}_2(\text{SO}_4)_2, 4\text{H}_2\text{O}$ in H_2O at t°. 100 g. H_2O dissolve grams CoSO_4 and grams Na_2SO_4 .

t°	g. CoSO_4	g. Na_2SO_4
20	26.65	24.91
25	25.365	23.325
30	23.13	21.61
35	22.55	20.85
40	20.975	20.055

(Koppel, Z. phys. Ch. 1905, 52. 397.)

Solubility of $\text{CoNa}_2(\text{SO}_4)_2, 4\text{H}_2\text{O} + \text{CoSO}_4, 7\text{H}_2\text{O}$ in H_2O at t°. 100 g. H_2O dissolve grams CoSO_4 and grams Na_2SO_4 .

t°	g. CoSO_4	g. Na_2SO_4	t°	g. CoSO_4	g. Na_2SO_4
18.5	28.61	23.82	30	32.695	18.17
20	29.42	23.015	35	34.065	15.61
25	30.73	20.575	40	35.01	13.715

(Koppel, Z. phys. Ch. 1905, 52. 397.)

Solubility of $\text{CoNa}_2(\text{SO}_4)_2, 4\text{H}_2\text{O} + \text{Na}_2\text{SO}_4, 10\text{H}_2\text{O}$ in H_2O at t°. 100 g. H_2O dissolve grams CoSO_4 and grams Na_2SO_4 .

t°	g. CoSO_4	g. Na_2SO_4
18.5	25.50	25.65
20	23.18	27.26
25	16.07	35.18
30	9.20	43.74

(Koppel.)

Solubility of $\text{CoNa}_2(\text{SO}_4)_2, 4\text{H}_2\text{O} + \text{Na}_2\text{SO}_4$ (anhydrous) in H_2O at t°. 100 g. H_2O dissolve grams CoSO_4 and grams Na_2SO_4 .

t°	g. CoSO_4	g. Na_2SO_4
35	7.204	50.79
40	7.456	50.095

(Koppel.)

See also $\text{CoSO}_4 + \text{Na}_2\text{SO}_4$ under CoSO_4 .

Cobaltous zinc sulphate.

Efflorescent. Decomp. on air. (Link, Crell. Ann. 1790, 1. 32.)

Cobaltous sulphate ammonia, $\text{CoSO}_4, 6\text{NH}_3$.

Sol. in H_2O with separation of ppt. (Rose, Pogg. 20. 152.) Very easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Fremy.)

Decomp. by alcohol.

Cobaltous sulphate hydrazine, $\text{CoSO}_4, 3\text{N}_2\text{H}_4$.

Insol. in H_2O . Decomp. by boiling with H_2O . Very sol. in dil. acids and $\text{NH}_4 + \text{Aq}$. (Franzen, Z. anorg. 1908, 60. 272.)

Cobaltous sulphate hydroxylamine, CoSO_4 , $\text{NH}_2\text{OH} + 2\text{H}_2\text{O}$.

Insol. in cold; sol. in hot H_2O with decomp. (Feldt, B. 1894, 27. 403.)

Columbium sulphate.

Sol. in H_2O . (Blomstrand.)

Cuprous sulphate, Cu_2SO_4 .

Decomp. by H_2O . Sol. in conc. HCl , in ammonia and sl. sol. in glacial acetic acid. (Recoura, C. R. 1909, 148. 1107.)

Cupric sulphate, basic, 10CuO , SO_3 .

(Pickering, Chem. Soc. 1907, 91. 1984.)

8CuO , $\text{SO}_3 + 12\text{H}_2\text{O}$. Ppt. (Kane, A. ch. 72. 269.)

5CuO , $\text{SO}_3 + 6\text{H}_2\text{O}$. Ppt. (Smith, Phil. Mag. J. 23. 196.)

4CuO , $\text{SO}_3 + 3\text{H}_2\text{O}$. Insol. in H_2O . (Roucher, J. Pharm. (3) 37. 50.)

Min. *Brochantite*. Sol. in acids and $\text{NH}_4\text{OH} + \text{Aq}$.

$+3\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O . Easily sol. in dil. acids, even HCl , $\text{H}_2\text{O}_2 + \text{Aq}$. Sl. sol. in $\text{CuSO}_4 + \text{Aq}$. Insol. in $\text{NaC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Casselmann, Z. anal. 4. 24.)

$+4\text{H}_2\text{O}$. Insol. in H_2O . (Proust.) Sol. in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$, and more easily in NH_4Cl , and $\text{NH}_4\text{NO}_3 + \text{Aq}$. (Lea.)

1 l. cold H_2O dissolves 0.017 g. (Pickering, C. N. 1883, 47. 182.)

$+5\text{H}_2\text{O}$. Min. *Langite*.

$+16\text{H}_2\text{O}$. (André, C. R. 100. 1138.)

7CuO , $2\text{SO}_3 + 5\text{H}_2\text{O}$. (Reindel, J. pr. 100. 1.)

$+6\text{H}_2\text{O}$. Wholly insol. in cold or hot H_2O . (Habermann, M. Ch. 5. 432.)

$+7\text{H}_2\text{O}$. Insol. in H_2O ; easily sol. in acids. Insol. in boiling $\text{CuSO}_4 + \text{Aq}$. (Reindel.)

3CuO , $\text{SO}_3 + 1\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O ; easily sol. in acids. (Steinmann, B. 15. 1412.)

$+2\text{H}_2\text{O}$. Insol. in H_2O ; sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Shenstone, Chem. Soc. 47. 375.)

$+2\frac{1}{2}\text{H}_2\text{O}$. (Reindel, J. pr. 102. 204.)

$+4\text{H}_2\text{O}$. Insol. in H_2O . (Grimbert and Barré, J. Pharm. (5) 21. 414.)

5CuO , $2\text{SO}_3 + 3\text{H}_2\text{O}$. (Wibel, Dissert. 1864.)

11CuO , $4\text{SO}_3 + 8\text{H}_2\text{O}$. (Clowes, C. N. 1898, 78. 155.)

8CuO , $3\text{SO}_3 + 10\text{H}_2\text{O}$. (Marchlewski and Sachs, Z. anorg. 1892, 1. 405.)

7CuO , $3\text{SO}_3 + 12\text{H}_2\text{O}$. (Étard, C. R. 1887, 104. 1615.)

5CuO , $2\text{SO}_3 + 5\text{H}_2\text{O}$. (Sabatier, Gm. K. 5. 1, 839.)

$6\text{H}_2\text{O}$. Min. *Arnimite*. (Weisbach, J. B. 1886. 2253.)

2CuO , SO_3 . Decomp. by cold H_2O into CuSO_4 and 4CuO , SO_3 . (Roucher.)

Insol. in H_2O . Decomp. by hot H_2O . Sol. in dil. acids. (Pozzi-Escot, Bull. Soc. 1913 (4) 13. 816.)

According to Pickering (C. N. 47. 181) only

3CuO , $\text{SO}_3 + 2\frac{1}{2}\text{H}_2\text{O}$ and 4CuO , $\text{SO}_3 + 4\text{H}_2\text{O}$ are true chemical compounds.

There is at 25° no definite basic sulphate of copper, all the basic sulphates being solid solutions. The solutions in contact with these basic sulphates contain SO_3 and CuO in equivalent quantities and are all acid in reaction. (Bell, J. phys. Chem. 1906, 12. 179.)

Cupric sulphate, CuSO_4 .

Anhydrous. Absorbs H_2O from the air. Combines with, and dissolves in H_2O with great evolution of heat.

$+ \text{H}_2\text{O}$. Permanent. Sol. in H_2O . (Étard, C. R. 87. 602.)

$+ 2\text{H}_2\text{O}$ (?). (Storer's Dict.)

$+ 3\text{H}_2\text{O}$. (Étard, C. R. 104. 1614.)

Does not exist. (Cross, C. N. 49. 229)

See Foote, p. 965.

$+ 5\text{H}_2\text{O}$. Superficially efflorescent in dry air.

Sol. in 2.34 pts. H_2O at 18° , and sat. solution has sp. gr. 1.2147. (Schiff, A. 109. 326.)

100 pts. $\text{CuSO}_4 + \text{Aq}$ sat. at b. pt., 102.2° , contains 45 pts. of the dry salt, or 100 pts. H_2O at 102.2° contain 81.82 pts. CuSO_4 . (Griffiths, Q. J. Sci. 18. 90.)

Sol. in less than 4 pts. H_2O at ord. temp., and even more sol. in boiling H_2O . (Bergmann.)

Sol. in 4 pts. cold, and 2 pts. hot H_2O . (See also: 100 pts. H_2O dissolve 33.103 pts. $\text{CuSO}_4 + 5\text{H}_2\text{O}$ at 15° , and solution has sp. gr. = 1.1859. (Mittler and Kraft, A. ch. (3) 41. 478.)

$\text{CuSO}_4 + \text{Aq}$ sat. at 8° has 1.17 sp. gr. (André, C. R. 24. 210.)

1 pt. $\text{CuSO}_4 + 5\text{H}_2\text{O}$ dissolves at:

4°	19°	31°	37.5°	54°
in 3.32	2.71	1.84	1.7	1.14 pts. H_2O

62.5°	75°	87.5°	100°	104°
in 1.27	1.07	0.75	0.55	0.47 pts. H_2O

(Brandes and Gruner, 1888.)

Sol. at 17.5 in 2.412 pts. H_2O . (Karrer.)

100 pts. H_2O dissolve at:

9°	10°	20°	30°
31.61	36.95	42.31	48.81 pts. $\text{CuSO}_4 + 5\text{H}_2\text{O}$

40°	50°	60°	70°
56.90	65.83	77.39	94.60 pts. $\text{CuSO}_4 + 5\text{H}_2\text{O}$

80°	90°	100°
118.03	156.44	203.32 pts. $\text{CuSO}_4 + 5\text{H}_2\text{O}$

(Poggiale, A. ch. (3) 8. 463.)

100 pts. H_2O dissolve at:

0°	20°	35°	54°
17	24.3	28.6	36.1 pts. anhydrous CuSO_4

(Tobler, A. 95. 193.)

100 pts. $\text{CuSO}_4 + \text{Aq}$ sat. at $11-14^\circ$ contains 16.23 pts. anhydrous CuSO_4 . (v. Hauer, J. pr. 103. 114.)

100 pts. H_2O dissolve 15.107 pts. CuSO_4 at 0° . (Piaff, A. 99. 224.)

100 pts. H₂O dissolve pts. CuSO₄ at t°.

t°	Pts. CuSO ₄
0	14.99
17.9	20.16
24.1	22.37

(Dracon, J. B. 1866. 61.)

100 pts. H₂O dissolve pts. CuSO₄ at t°.

t°	Pts. CuSO ₄	t°	Pts. CuSO ₄	t°	Pts. CuSO ₄
0	14.15	40	28.50	80	54.53
10	17.50	50	33.31	90	64.35
20	20.53	60	39.01	100	75.22
30	24.34	70	45.74		

(Patrick and Aubert, Transactions of Kansas Acad. of Sci. 1874. 19.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. CuSO ₄	t°	Pts. CuSO ₄	t°	Pts. CuSO ₄
0	15.5	15	27.5	70	45.7
1	16.3	16	27.9	71	46.4
2	16.6	17	28.3	72	47.2
3	16.9	18	28.7	73	47.9
4	17.2	19	29.1	74	48.7
5	17.5	20	29.5	75	49.5
6	17.8	21	29.9	76	50.3
7	18.1	22	30.3	77	51.1
8	18.4	23	30.7	78	51.9
9	18.7	24	31.1	79	52.7
10	19.1	25	31.5	80	53.5
11	19.3	26	31.9	81	54.3
12	19.6	27	32.3	82	55.1
13	19.9	28	32.7	83	55.9
14	20.2	29	33.2	84	56.8
15	20.5	30	33.6	85	57.8
16	20.8	31	34.1	86	58.7
17	21.1	32	34.5	87	59.7
18	21.4	33	35.0	88	60.7
19	21.7	34	35.5	89	61.7
20	22.0	35	36.0	90	62.7
21	22.3	36	36.6	91	63.7
22	22.6	37	37.2	92	64.8
23	23.0	38	37.8	93	65.8
24	23.3	39	38.4	94	66.9
25	23.7	40	39.0	95	68.0
26	24.0	41	39.6	96	69.1
27	24.4	42	40.2	97	70.2
28	24.7	43	40.9	98	71.3
29	25.1	44	41.5	99	72.4
30	25.5	45	42.2	100	73.5
31	25.9			101	74.6
32	26.3			102	75.7
33	26.7			103	76.8
34	27.1			104	77.95

(Mulder, Scheik. Verhandel. 1864. 79.)

If solubility S = pts. anhydrous CuSO₄ in 100 pts. solution, $S = 11.6 + 0.2614t$ from -2° to 55°; $S = 28.5 + 0.3700t$ from 55° to 105°; $S = 45.0 - 0.0293t$ from 105° to 190°. (Étard, C. R. 104. 1614.)

Solubility decreases above 120°, owing to formation of basic salt. (Tilden and Shenstone, Phil. Trans. 1884. 23.)

100 ccm. H₂O dissolve 14.92 g. CuSO₄ at 0°. (Engel, C. R. 102. 113.)

100 ccm. H₂O dissolve 22.28-22.30 g. CuSO₄ at 20°. (Trevor, Z. phys. Ch. 7. 468.)

Sat. CuSO₄ + Aq contains % CuSO₄ at t°.

t°	% CuSO ₄	t°	% CuSO ₄
-1	12.1	111	38.8
+7	14.1	112	38.9
9	14.5	94	41.8
18	16.9	96	41.9
20	17.2	97	42.0
20	17.4	100	43.6
35	21.3	108	43.8
39	21.8	110	43.4
45	23.9	116	43.8
54	26.9	116	44.0
54	26.6	120	44.8
61	28.8	132	44.8
63	29.1	133	44.7
65	30.0	143	45.0
70	31.6	150	44.2
71.7	32.6	165	44.5
76	34.5	179	42.9
80	36.6	189	42.2
86	37.8		

(Étard, A. ch. 1894, (7) 2. 554.)

Solubility in H₂O at t°.

t°	g CuSO ₄ per 100 g H ₂ O
0	14.15
10	17.68
15	19.25
20	20.78
25	22.29 (by interpolation)

(Cohen, Z. phys. Ch. 1907, 60. 713.)

1.399 mol. are sol. in 1 l. H₂O at 25°. (Herz, Z. anorg. 1910, 67. 366.)

100 g. CuSO₄ + Aq sat. at 30° contain 20.32 anhyd. CuSO₄. (Schreinemakers, Z. phys. Ch. 1910, 71. 110.)

+7H₂O. (Houbaudran, C. R. 65. 1249.)

+6H₂O. (Houbaudran, C. R. 66. 487.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at 18° . $\% = \%$
 $\text{CuSO}_4 + 5\text{H}_2\text{O}$.

%	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.0063	11	1.0716	21	1.1427
2	1.0126	12	1.0785	22	1.1501
3	1.0190	13	1.0854	23	1.1585
4	1.0254	14	1.0923	24	1.1659
5	1.0319	15	1.0993	25	1.1738
6	1.0384	16	1.1063	26	1.1817
7	1.0450	17	1.1135	27	1.1898
8	1.0516	18	1.1208	28	1.1980
9	1.0582	19	1.1281	29	1.2063
10	1.0649	20	1.1354	30	1.2146

(Schiff, calculated by Gerlach, Z. anal. 8. 288.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at 23.9° . a=no. of $\frac{1}{2}$ mols. in grms. dissolved in 1000 grms. H_2O ; b=sp. gr. if a is $\text{CuSO}_4 = 5\text{H}_2\text{O}$ ($\frac{1}{2}$ mol. wt. = 125); c=sp. gr. if a is CuSO_4 ($\frac{1}{2}$ mol. wt. = 80).

a	b	c
1	1.076	1.080
2	1.142	1.154
3	1.200	1.225

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at 15° .
 $\% = \%$ $\text{CuSO}_4 + 5\text{H}_2\text{O}$.

%	Sp. gr.	%	Sp. gr.
5	1.0335	20	1.1443
10	1.0688	25	1.1848
15	1.1060	mother liquor	1.185

(Gerlach, Dingl. 181. 131.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at 18° .

% CuSO_4	Sp. gr.	% CuSO_4	Sp. gr.
5	1.0513	15	1.1675
10	1.1073	17.5	1.2003

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at 0° . S=pts. CuSO_4 in 100 pts. solution.

S	Sp. gr.	S	Sp. gr.
11.9315	1.1371	5.2181	1.0578
9.8159	1.1108	2.6460	1.0290
7.5474	1.0833

(Charpy, A. ch. (6) 29. 26.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at room temp

% CuSO_4	Sp. gr.
6.79	1.055
12.57	1.1151
17.49	1.1635

(Wagner, W. Ann. 1883, 18. 265.)

Sp. gr. of $\text{CuSO}_4 + \text{Aq}$ at 25° .

Concentration of $\text{CuSO}_4 + \text{Aq}$	Sp. gr.
1-normal	1.0790
$\frac{1}{5}$ " "	1.0402
$\frac{1}{10}$ " "	1.0205
$\frac{1}{20}$ " "	1.0103
$\frac{1}{40}$ " "	1.0050

(Wagner, Z. phys. Ch. 1890, 5. 38.)

B.-pt. $\text{CuSO}_4 + \text{Aq}$ containing pts. CuSO_4 100 pts. H_2O .

B.-pt.	Pts. CuSO_4	B.-pt.	Pts. C
100.5°	21.3	103.0°	69
101.0	36.9	103.5	74
101.5	48.0	104.0	80
102.0	56.2	104.2	82
102.5	63.0

(Gerlach, Z. anal. 28. 434.)

Sat. $\text{CuSO}_4 + \text{Aq}$ boils at 102.2° , and tains 81.8 pts. CuSO_4 to 100 pts. (Griffiths.)

Crust forms at 102.3° , and solution con 60.3 pts. CuSO_4 to 100 pts. H_2O ; highest observed, 104.8° . (Gerlach, Z. anal. 28.

Sol. in $\text{HCl} + \text{Aq}$, causing a₂ reducti temperature of about 17° .

Very sl. sol. in conc. H_2SO_4 . (Schub

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 0° .

G. per 100 g. H_2O		Sp. gr.
H_2SO_4	CuSO_4	
0.00	14.85	1.144
2.03	14.29	1.142
7.16	15.65	1.156
15.20	9.90	1.170
26.57	6.43	1.190
27.57	6.19	1.211
35.2	3.99	1.224

(Engel, C. R. 1887, 104. 507.)

Solubility in H₂SO₄+Aq at 25°.

Solution		Solid phase
% CuO	% SO ₃	
9.17	9.26	CuSO ₄ +5H ₂ O
5.91	15.90	
3.39	23.09	
1.82	28.75	
1.32	39.74	
...	41.29	CuSO ₄ +5H ₂ O and CuSO ₄ +3H ₂ O
...	41.04	
1.38	43.63	CuSO ₄ +3H ₂ O
1.02	47.82	
...	49.07	
0.38	51.46	CuSO ₄ +H ₂ O
0.368	53.51	
0.109	62.14	
0.105	68.34	
0.15	72.41	CuSO ₄
0.07	74.26	

e results show that the hydrates of which are stable at 25° are CuSO₄+3H₂O and +H₂O.
nd Taber, J. phys. Chem. 1908, 12. 175.)

Solubility in H₂SO₄+Aq at 25°.

tion contains		Solid phase
10	% H ₂ SO ₄	
7	none	CuSO ₄ +5H ₂ O
2	11.14	
2	25.53	
5	36.77	
3	42.15	
9	47.66	CuSO ₄ +5H ₂ O and CuSO ₄ +3H ₂ O
3	49.00	
3	49.20	
4	49.29	CuSO ₄ +3H ₂ O
0	50.23	
9	54.78	CuSO ₄ +3H ₂ O and CuSO ₄ +H ₂ O
1	55.84	
5	55.60	CuSO ₄ +H ₂ O
5	61.79	
7	77.93	
5	83.29	CuSO ₄ +H ₂ O and CuSO ₄
9	85.46	
4	85.72	CuSO ₄
2	85.81	
0	86.04	CuSO ₄
9	92.70	

e results show that the hydrates of which are stable at 25° are CuSO₄+3H₂O and +H₂O.
J. Am. Chem. Soc. 1915, 37. 290.)

ol. in sat. NH₄Cl+Aq, with separation
uble sulphate.

Solubility of CuSO₄ in CuCl₂+Aq at 30°.

% CuCl ₂	% CuSO ₄	Solid phase
0	20.32	CuSO ₄ , 5H ₂ O
6.58	13.62	"
15.68	8.93	"
25.67	4.77	"
39.48	3.21	"
42.47	2.90	CuSO ₄ , 5H ₂ O+CuCl ₂ , 2H ₂ O
43.25	1.14	CuCl ₂ , 2H ₂ O
43.95	0	"

(Schreinemakers, Arch. Néer. Sci. 1910, (2) 15. 117.)

Solubility of CuSO₄ in LiCl+Aq at 25°.
Solid phase, CuSO₄, 5H₂O.
(G. mols. per l. of solution.)

LiCl	CuSO ₄
0	1.399
0.73	1.257
1.40	1.176
2.83	1.067

(Herz, Z. anorg. 1910, 67. 366.)

Solubility of CuSO₄ in KCl+Aq at 25°.
Solid phase, CuSO₄+5H₂O.
(G. mols. per l. of solution.)

KCl	CuSO ₄
0.56	1.496
1.17	1.561
2.34	1.819

(Herz.)

Solubility of CuSO₄ in NaCl+Aq at 25°.
Solid phase, CuSO₄+5H₂O.
(G. mols. per l. of solution.)

NaCl	CuSO ₄
0	1.399
0.36	1.404
1.32	1.426
2.53	1.507

(Herz.)

Solubility of CuSO₄ in RbCl+Aq at 25°, containing 1.094 g. mols. per l.=1.568 g. mols. (Herz.)

Slowly sol. in sat. KNO₃+Aq, with separation of a double sulphate.

Very slowly sol. in sat. NaNO₃+Aq, with separation of a double sulphate. (Karsten, Berl. Abhandl. 1840. 10.)

Solubility of CuSO₄ in (NH₄)₂SO₄+Aq at 0°.

g. per 100 cc. solution		Sp. gr.	g. per 100 cc. solution		Sp. gr.
(NH ₄) ₂ SO ₄	CuSO ₄		(NH ₄) ₂ SO ₄	CuSO ₄	
0	14.79	1.144	5.59	5.13	1.081
3.61	16.09	1.190	7.51	2.95	1.071
4.63	8.38	1.108	12.31	0.94	1.082
4.90	7.26	1.099	20.65	0.80	1.116

(Engel, C. R. 1886, 102. 114.)

See also under (NH₄)₂SO₄.

Solubility of CuSO₄ in Li₂SO₄+Aq at 30°.

Composition of the solution		Solid phase
% by wt. CuSO ₄	% by wt. Li ₂ SO ₄	
20.32	0	CuSO ₄ ·5H ₂ O
17.50	3.54	"
16.10	6.08	"
13.55	11.94	"
12.14	15.72	"
11.04	17.92	"
10.05	20.55	CuSO ₄ ·5H ₂ O + Li ₂ SO ₄ ·H ₂ O
10.08	20.51	"
10.07	20.49	"
6.41	22.23	Li ₂ SO ₄ ·H ₂ O
3.39	23.59	"
0	25.24	"

(Schreinemakers, Z. phys. Ch. 1909, 66. 692.)

Sol. in CuCl₂, (NH₄)₂SO₄, NH₄Cl+Aq at 30°. (Schreinemakers, Z. phys. Ch. 1909, 69. 565.)

Sol. in (NH₄)₂SO₄, Li₂SO₄+Aq. at 30°. (Schreinemakers, Z. phys. Ch. 1909, 66. 694.)

100 pts. sat. solution of CuSO₄ and FeSO₄ contain 17.43 pts. of the salts at 11-14°. (v. Hauer, J. pr. 103. 114.)

100 pts. H₂O dissolve 10.85 pts. CuSO₄, 17.47 pts. MgSO₄, and 5.78 pts. Na₂SO₄ at 0°. (Diacon.)

100 pts. H₂O dissolve 7.169 pts. CuSO₄, 21.319 pts. MgSO₄, and 6.830 pts. Na₂SO₄ at 0°. (Plaff.)

Slowly and sl. sol. in sat. MgSO₄+Aq. (Karsten.)

Solubility of CuSO₄ in H₂O in presence of MgSO₄. 100 pts. H₂O dissolve—

No.	CuSO ₄	MgSO ₄	No.	CuSO ₄	MgSO ₄
1	0	26.37	5	12.03	15.67
2	2.64	25.91	6	13.61	8.64
3	4.75	25.30	7	14.99	0
4	9.01	23.54

In 1, 2, and 3, MgSO₄ was in excess and given amt. CuSO₄ added; in 4, both CuSO₄ and

MgSO₄ were in excess; in 5, 6, and 7, CuSO₄ was in excess. (Diacon, l. c.)

100 pts. sat. solution of CuSO₄ and MgSO₄ contain 28.58 pts. of the salts at 11-14°. (v. Hauer, J. pr. 103. 114.)

100 pts. sat. solution of CuSO₄ and MnSO₄ contain 37.90 pts. of the salts at 11-14°. (v. Hauer.)

Solubility of CuSO₄+MnSO₄ in H₂O at 25°.

G. per 100 g. H ₂ O		G. per 100 g. H ₂ O	
CuSO ₄	MnSO ₄	CuSO ₄	MnSO ₄
20.2	0	9.39	46.77
19.76	3.69	6.47	53.39
13.65	31.52	3.01	58.93
11.61	39.41	0.0	61.83

(Stortenbecker, Z. phys. Ch. 1900, 34. 112)

100 pts. sat. solution of CuSO₄ and NiSO₄ contain 31.03 pts. of the salts at 11-14°. (v. Hauer.)

Solubility of CuSO₄+NiSO₄ in H₂O.

g. per 100 g. H ₂ O			Mol. % CuSO ₄	
t°	CuSO ₄	NiSO ₄	Solution	Solid phase
35	9.62	583.9	1.57	0.35
	41.66	484.4	7.69	2.12
	75.39	553.5	11.66	4.77
	106.40	506.5	16.92	6.52
	172.0	483.8	25.6	13.86
	186.9	468.8	27.90	{ 18.77 94.91
67	20.04	729.3	2.65	0.93
	66.01	706.2	8.31	2.46
	88.08	501.6	13.55	3.92
	147.94	675.0	16.39	6.66
	249.9	747.8	24.46	23.32

(Fock, Z. Kryst. Min. 1897, 28. 387.)

More easily sol. in sat. K₂SO₄+Aq than in Na₂SO₄ or MgSO₄+Aq, forming a double sulphate, which separates out. (Karsten.)

K₂SO₄ and CuSO₄ mutually displace each other in saturated solutions. (Rüdorff, Pogg. 148. 555.)

When K₂SO₄ and CuSO₄ both in excess, are dissolved in H₂O, a maximum of solubility of 15.61 pts. of the two salts in 100 pts. H₂O at 25° is reached in 30 minutes, after which the solubility decreases. This result is obtained either by treating excess of the two salts with H₂O at 25°, or cooling solutions of the two salts sat. at higher temp. to 25°. The salts are in the proportion of 5.2 pts. K₂SO₄ to 10.4 pts. CuSO₄. If present in the same proportion as in their saturated solutions, 5.41 pts. K₂SO₄ to 10.13 pts. CuSO₄ would be required.

If sat. solution of one salt is added to sat. solution of the other, $K_2Cu(SO_4)_2 + 6H_2O$ separates, as it is less sol. than either simple salt, until a state of equilibrium is reached, after which there is no separation, contrary to Rüdorff (see above). (Trevor, Z. phys. Chem. 7. 486.)

$CuSO_4 + Na_2SO_4$.
Solubility of $CuSO_4$ in presence of Na_2SO_4 at 0°. 100 pts. H_2O dissolve—

No.	$CuSO_4$	Na_2SO_4	No.	$CuSO_4$	Na_2SO_4
1	0	4.53	5	15.84	3.55
2	6.01	5.34	6	15.33	1.98
3	9.81	5.73	7	14.99	0
4	16.67	6.48

In 1, 2, and 3, Na_2SO_4 was in excess and given amt. $CuSO_4$ added; in 4, both $CuSO_4$ and Na_2SO_4 were in excess; in 5, 6, and 7, $CuSO_4$ was in excess and Na_2SO_4 added. (Piacon, J. B. 1866. 61.)
100 pts. H_2O dissolve 8.038 pts. $CuSO_4$ and 31 pts. Na_2SO_4 at 0°. (Pfaff, A. 99. 224.)
100 pts. H_2O dissolve 20.7 pts. $CuSO_4$ and 9.9 pts. Na_2SO_4 at 15°. (Rüdorff, B. 6. 484.)

Solubility of $CuSO_4 + Na_2SO_4$.
1. Solid phase, 3 mol. $CuSO_4 + 1$ mol. Na_2SO_4 .
2. Solid phase, 1 mol. $CuSO_4 + 1$ mol. Na_2SO_4 .
3. Solid phase, 1 mol. $CuSO_4 + 3$ mol. Na_2SO_4 .
(G. in 100 g. H_2O .)

°	1		2		3	
	$CuSO_4$	Na_2SO_4	$CuSO_4$	Na_2SO_4	$CuSO_4$	Na_2SO_4
	19.75	12.49	19.70	12.50	19.69	12.55
	20.69	15.88	20.75	15.90	20.70	15.92
	22.03	16.36	21.00	20.14	15.28	22.70
	32.37	11.75	31.45	13.41	28.76	15.25

Massol and Males, C. R. 1901, 133. 287.)

Solubility of $CuSO_4 \cdot 5H_2O + Na_2SO_4 \cdot 10H_2O$.

t°	% $CuSO_4$	% Na_2SO_4
0	13.40	6.23
12	14.83	9.82
15	15.00

(Koppel, Z. phys. Chem. 1903, 42. 8.)
See also under $CuNa_2(SO_4)_2 + 6H_2O$.

$CuSO_4 + ZnSO_4$.
Very slowly sol. in sat. $ZnSO_4 + Aq$, forming a double salt which separates. (Karsten.)
100 pts. sat. solution of $CuSO_4$ and $ZnSO_4$ contain 32.70 pts. of the salts at 11–14°. (Hauer.)

Solubility of $CuSO_4 + ZnSO_4$ in H_2O at 25°.

Mols. per 100 mols. H_2O		Mols. per 100 mols. H_2O	
Cu	Zn	Cu	Zn
2.28	0	0.82	5.03
1.83	2.08	0.51	5.59
1.41	3.60	0.30	5.56
1.19	5.01	0.00	6.42
1.86	3.36	1.19	5.01
1.22	4.45	0.51	5.59
1.01	4.72	0.267	5.77
...	...	0.00	5.94

(Stortenbecker, Z. phys. Chem. 1897, 22. 62.)

Insol. in liquid NH_3 . (Franklin, Am. Chem. J. 1898, 20. 827.)

Insol. in liquid CO_2 . (Büchner, Z. phys. Chem. 1906, 54. 674.)

100 pts. of a sat. solution in 40% alcohol contains 0.25 pt. $CuSO_4 + 5H_2O$; 20% alcohol, 3.1 pts.; 10% alcohol, 13.3 pts. (Schiff, A. 118. 362.)

Anhydrous $CuSO_4$ is sol. in absolute methyl alcohol, but insol. in absolute ethyl alcohol. $CuSO_4 + xH_2O$ is insol. in methyl or ethyl alcohol. (Klepl, J. pr. (2) 25. 526.)

100 pts. absolute methyl alcohol dissolve 1.05 pts. anhydrous $CuSO_4$ at 18°.

100 pts. absolute methyl alcohol dissolve 15.6 pts. $CuSO_4 + 5H_2O$ at 18°; 100 pts. 93.5% methyl alcohol dissolve 0.93 pt. $CuSO_4 + 5H_2O$ at 18°; 100 pts. 50% methyl alcohol dissolve 0.4 pt. $CuSO_4 + 5H_2O$ at 18°; 100 pts. absolute methyl alcohol dissolve 13.4 pts. $CuSO_4 + 5H_2O$ at 3°.

100 pts. absolute ethyl alcohol dissolve 1.1 pts. $CuSO_4 + 5H_2O$ at 3°. (de Bruyn, Z. phys. Chem. 10. 786.)

Methyl alcohol dissolves 11.5% $CuSO_4 + 5H_2O$ at 0°. (Auger, C. R. 1906, 142. 1272.)

Glacial acetic acid precipitates $CuSO_4$ completely from $CuSO_4 + Aq$.

100 g. 95% formic acid dissolve 0.05 g. $CuSO_4 + 5H_2O$ at 18.5°. (Aschan, Chem. Ztg. 1913, 37. 1117.)

Sol. in glycerine (Pelouze), picoline (Unverdorben).

100 g. glycerine dissolve 36.3 g. $CuSO_4 + 5H_2O$ at 15–16°. (Ossendowski, Pharm. J. 1907, 79. 575.)

100 g. glycerine dissolve 30.0 g. $CuSO_4$ at 15.5°.

100 g. solution of $CuSO_4$ in glycol contain 7.6 g. at 14.6°. (de Coninck, Bull. Ac. Belg. 1905. 257.)

Anhydrous $CuSO_4$ is insol. in acetone. (Krug and M'Elroy, J. Anal. Chem. 6. 184.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014); methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314); benzonitrile. (Naumann, B. 1914, 47. 1370.)

Min. Chalcanthite.

Cupric sodium sulphate, $\text{CuSO}_4, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$.
Solubility of the pure double salt $\text{CuSO}_4, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$.

t°	100 grams solution contain	
	CuSO ₄ grams	Na ₂ SO ₄ grams
17.7	14.34	13.34
19.5	14.54	12.90
23	14.36	12.76
30	14.07	12.37
40.15	13.73	12.26

(Koppel, Z. phys. Ch. 1903, 42. 8.)

Solubility of the mixture of $\text{CuSO}_4, \text{Na}_2\text{SO}_4 + \text{H}_2\text{O}$ and CuSO_4 .

t°	100 g. solution contain	
	CuSO ₄ grams	Na ₂ SO ₄ grams
17.7	14.99	13.48?
19.5	15.62	12.06
23	16.41	11.35
30	17.97	9.95
40.15	20.56	8.00

(Koppel.)

Solubility of the mixture of $\text{CuSO}_4, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$ and Na_2SO_4 .

t°	100 g. solution contain	
	CuSO ₄ grams	Na ₂ SO ₄ grams
18	13.53	13.844
19.5	11.847	15.116
20	11.339	15.697
23	8.185	18.723
25	6.284	21.198
26	5.507	22.44
28	3.746	24.963
28.3	3.661	...
30	2.607	28.383
30.2	2.422	...
32.2	1.465	32.442
33.9	1.475	32.299
35.3	1.471	32.072
37.2	1.494	31.96

(Koppel.)

Solubility of $\text{CuSO}_4, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$ in presence of varying amounts of Na_2SO_4 .

t°	100 g. solution contain	
	CuSO ₄ grams	Na ₂ SO ₄ grams
30	5.38	22.17
30	5.41	21.92
30.1	3.69	25.37
40.15	3.97	23.90
30	1.57	32.09

(Koppel.)

Copper sulphate and sodium sulphate unite to form a double salt, $\text{CuSO}_4, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$, which is stable in the presence of the solution above 16.7°. In the presence of copper sulphate the solubility of $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ is greater than that of the pure salt. (Koppel.)

Cupric thalious sulphate, $\text{CuSO}_4, \text{Ti}_2\text{SO}_4 + 6\text{H}_2\text{O}$.

Decomp. by recrystallising from H_2O . (Willm, A. ch. (4) 5. 55.)
1 l. H_2O dissolves 81 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Cupric zinc sulphate, $\text{CuSO}_4, 3\text{ZnSO}_4 + 28\text{H}_2\text{O}$
Efflorescent. 100 pts. H_2O dissolve 80 pts. salt at 8°. Sol. in all proportions in boiling H_2O . (Lefort.)
 $\text{CuSO}_4, 2\text{ZnSO}_4 + 21\text{H}_2\text{O}$. (v. Hauer, Pogg. 125. 637.)
 $\text{CuSO}_4, \text{ZnSO}_4 + 12\text{H}_2\text{O}$. (Boisbaudran.)
 $2\text{CuSO}_4, 2\text{ZnSO}_4, \text{H}_2\text{SO}_4$. (Étard.)

Cuprous sulphate ammonia, $\text{Cu}_2\text{SO}_4, 4\text{NH}_3$.
Sol. in $\text{NH}_4\text{OH} + \text{Aq}$; decomp. by pure H_2O . (Péchar, C. R. 1903, 136. 504.)
+ H_2O . Decomp. by H_2O . (Foerster and Blankenburg, B. 1906, 39. 4434.)

Cupric sulphate ammonia, basic, $\text{CuSO}_4, 3\text{CuO}, 2\text{NH}_3 + 5\text{H}_2\text{O}$.
Decomp. by hot H_2O . (Pickering, Chem. Soc. 43. 336.)

Cupric sulphate ammonia (Cuprammonium sulphate), $\text{CuSO}_4, \text{NH}_3$.
Decomp. by H_2O . (Kane.)
Decomp. by H_2O giving a basic sulphate. (Bouzat, C. R. 1902, 135. 535.)
 $\text{CuSO}_4, 2\text{NH}_3, [\text{CuSO}_4, 2\text{NH}_3 + 3\text{H}_2\text{O}]$. (Mendelejeff, B. 3. 422.). Decomp. by excess of H_2O into—
 $\text{CuSO}_4, 4\text{NH}_3 + \text{H}_2\text{O}$. Sol. in 1.5 pts. H_2O , but decomp. by much H_2O . Insol. in alcohol. Insol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. (André, C. R. 100. 1138.)

Sol. in small quantity of H_2O ; decomp. in dil. solution. (Bouzat.)
100 g. H_2O dissolve 44.56 g. anhydrous comp. at 25°. (Pudschies, Dissert.)
100 g. H_2O dissolve 18.05 g. at 21–22°. (Horn and Taylor, Am. Ch. J. 1904, 32. 268.)
 $\text{CuSO}_4, 5\text{NH}_3$. Completely sol. in H_2O . (Rose, Pogg. 20. 150.)
Sol. in small amt. of H_2O ; decomp. in dil. solution. Insol. in liquid NH_3 . (Bouzat, C. R. 1902, 135. 535.)

Cuprous sulphate carbon monoxide, $\text{Cu}_2\text{SO}_4, 2\text{CO} + \text{H}_2\text{O}$.
Very unstable. (Joannis, C. R. 1903, 138. 615.)

Cupric sulphate zinc oxide, $\text{CuSO}_4, 2\text{ZnO} + 21\text{H}_2\text{O}$.

(Larsen, Ch. Z. Repert. 1896, 20. 317.)
 $2\text{CuSO}_4, 3\text{ZnO} + 12\text{H}_2\text{O}$. (Mailhe, A. ch. 1902, (7) 27. 169.)

Didymium sulphate, basic, $\text{Di}_2\text{O}_3, \text{SO}_3 = (\text{DiO})_2\text{SO}_4$.

Insol. in cold or boiling H_2O . (Marignac.)
 Slowly sol. in hot dil. $\text{HCl} + \text{Aq}$. Easily sol. in conc. acids.

+ $8\text{H}_2\text{O}$. Precipitate. (Hermann.)
 Composition is $2\text{Di}_2\text{O}_3, 3\text{SO}_3 + 3\text{H}_2\text{O}$ or $\text{Di}_2(\text{SO}_4)_3 + \text{Di}_2\text{O}_3 \cdot \text{H}_2\text{O}$. (Frerichs and Smith.)
 Composition is $5\text{Di}_2\text{O}_3, 3\text{SO}_3 + x\text{H}_2\text{O}$. (Cleve, B. 11. 910.)

Didymium sulphate, $\text{Di}_2(\text{SO}_4)_3$.

Anhydrous. By saturating cold H_2O and warming the solution, the following results were obtained—100 pts. H_2O dissolve at:

12° 18° 25° 38° 50°
 43.1 25.8 20.6 13.0 11.0 pts. $\text{Di}_2(\text{SO}_4)_3$.

+ $6\text{H}_2\text{O}$. H_2O dissolves this salt very slowly; 100 pts. H_2O dissolve 13 pts. $\text{Di}_2(\text{SO}_4)_3$ in 24 hours, and 16.4 pts. in 2 days. If solution is evap. in vacuo until $\text{Di}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ separates out, 34 pts. $\text{Di}_2(\text{SO}_4)_3$ remain dissolved in 100 pts. H_2O .

+ $5\text{H}_2\text{O}$. (Cleve.)

+ $8\text{H}_2\text{O}$. Solutions of this salt contain at:

19° 40° 50° 100°
 11.7 8.8 6.5 1.6 pts. $\text{Di}_2(\text{SO}_4)_3$.

(Marignac, A. ch. (3) 38. 170.)

+ $9\text{H}_2\text{O}$. (Zschiesche, J. Pr. 107. 75.)

Didymium potassium sulphate, $\text{K}_2\text{SO}_4, \text{Di}_2(\text{SO}_4)_3 + 2\text{H}_2\text{O}$.

Sol. in 63 pts. H_2O . Insol. in sat. $\text{K}_2\text{SO}_4 + \text{Aq}$. (Marignac.)

$3\text{K}_2\text{SO}_4, \text{Di}_2(\text{SO}_4)_3$. Sol. in 83 pts. H_2O at 18°. Insol. in cold, sl. sol. in boiling sat. $\text{K}_2\text{SO}_4 + \text{Aq}$, 100 ccm. of which retain 55 mg. Di_2O_3 in solution. (Cleve.)

$4\text{K}_2\text{SO}_4, \text{Di}_2(\text{SO}_4)_3$. (Cleve.)

$9\text{K}_2\text{SO}_4, 2\text{Di}_2(\text{SO}_4)_3 + 3\text{H}_2\text{O}$. (Cleve.)

Didymium sodium sulphate, $\text{Di}_2(\text{SO}_4)_3, \text{Na}_2\text{SO}_4$, and $+2\text{H}_2\text{O}$.

Sol. in 200 pts. H_2O (Marignac), and still less in sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$, 100 ccm. of which dissolve only 70 mg. Di_2O_3 at ord. temp. (Cleve.)

Didymium thalious sulphate, $(\text{Di}_2\text{SO}_4)_3, 3\text{Ti}_2\text{SO}_4$.

Ppt.

$\text{Di}_2(\text{SO}_4)_3, \text{Ti}_2\text{SO}_4 + 2\text{H}_2\text{O}$. Sol. in H_2O . (Zschiesche, J. pr. 107. 98.)

Erbium sulphate, $\text{Er}_2(\text{SO}_4)_3$.

Anhydrous. Easily and rapidly sol. in H_2O . 100 pts. H_2O dissolve 43 pts. anhydrous salt at 0°.

+ $8\text{H}_2\text{O}$. Less sol. in H_2O than anhydrous salt. 100 pts. H_2O dissolve 30 pts. $\text{Er}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ (=23 pts. $\text{Er}_2(\text{SO}_4)_3$) at about 20°; at 100°, 100 pts. $\text{Er}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ remain dissolved. Sat. solution deposits crystals when heated to 55°. (Höglund.)

100 g. of sat. solution of $\text{Er}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ at 25° in H_2O contains 11.94 g. anhyd. $\text{Er}_2(\text{SO}_4)_3$. (Wirth, Z. anorg. 1912, 76. 174.)

Solubility of $\text{Er}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ in H_2SO_4 at 25°.

N = equiv. g. H_2SO_4 in 1 l. of solvent.

C = g. oxide in 100 g. of solution.

C' = g. anhyd. salt in 100 g. of solution.

N	C	C'	N	C	C'
0	4.604	7.60	4.32	2.00	3.301
0.1	4.615	7.618	6.685	0.9115	1.505
1.1	3.64	6.00	9.68	0.4439	0.73
2.16	3.04	5.018	15.15	0.145	0.239

(Wirth, Z. anorg. 1912, 76. 174.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Erbium potassium sulphate, $\text{Er}_2(\text{SO}_4)_3, 3\text{K}_2\text{SO}_4$.

Slowly sol. in H_2O . (Höglund.)

Erbium sodium sulphate, $\text{Er}_2(\text{SO}_4)_3, 5\text{Na}_2\text{SO}_4 + 7\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve.)

Europium sulphate, $\text{Eu}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$.

Stable in the air. (Urbain and Lecomte, C. R. 1904, 138. 628.)

Gadolinium sulphate, $\text{Gd}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$.

Solubility in H_2O .

Temp.	Pts. of $\text{Gd}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ in 100 pts. H_2O
0°	3.98
9.3–10.6°	3.33
14.0°	2.80
25.0°	2.40
34.4°	2.26

(Benedicks, Z. anorg. 1899, 22. 409–410.)

100 g. sat. solution of $\text{Gd}(\text{SO}_4)_3 + 8\text{H}_2\text{O}$ at 25° in H_2O contain 2.981 g. anhyd. $\text{Gd}_2(\text{SO}_4)_3$. (Wirth, Z. anorg. 1912, 76. 174.)

Solubility in H_2SO_4 . Solid phase, $\text{Gd}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$.

N = equiv. g. H_2SO_4 in 1 l. of solvent.

C = g. oxide in 100 g. of solution.

C_1 = g. anhyd. $\text{Gd}_2(\text{SO}_4)_3$ in 100 g. of solution.

N	C	C_1	N	C	C_1
0	1.793	2.981	2.16	1.789	2.974
0.1	1.98	3.291	6.175	0.528	0.8777
0.505	2.365	3.931	12.6	0.0521	0.0867
1.1	2.29	3.807

(Wirth.)

Sol. in sat. $\text{K}_2\text{SO}_4 + \text{Aq}$.

Gadolinium potassium sulphate, $\text{Gd}_2(\text{SO}_4)_3 \cdot \text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}$.

100 grams. sat. solution in $\text{K}_2\text{SO}_4 + \text{Aq}$ contains 0.87—0.77 grams Gd_2O_3 . (Benedicks, Z. anorg. 1900, 22. 410.)

Gallium sulphate, $\text{Ga}_2(\text{SO}_4)_3$.

Not deliquescent, but very sol. in H_2O . Sol. in 60% alcohol; insol. in ether. (Boisbaudran.)

Aqueous solution decomp. into basic salt by boiling, which redissolves, however, on cooling.

Gallium potassium sulphate, $\text{Ga}_2\text{K}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

(Soret, Arch. sc. phys. nat. 1885, 14. 96.)

Gallium rubidium sulphate, $\text{Rb}_2\text{Ga}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

(Soret, Arch. sc. phys. nat. 1885, 14. 96.)

Glucinum sulphate, basic, $3\text{GdO} \cdot \text{SO}_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O , but decomp. by heating or dilution. (Berzelius.)

$2\text{GdO} \cdot \text{SO}_3 + 3\text{H}_2\text{O}$. Sol. in H_2O .

$9\text{GdO} \cdot \text{SO}_3 + 14\text{H}_2\text{O}$ (?). Precipitate. Insol. in H_2O . (Berzelius.)

According to Debray, this salt when carefully washed is GdO_2H_2 .

Glucinum sulphate, GdSO_4 .

Anhydrous.

Nearly insol. in H_2O , but slowly attacked by cold, rapidly by hot H_2O , and is converted into $\text{GdSO}_4 + 4\text{H}_2\text{O}$ before dissolving. (Parsons, Z. anorg. 1904, 42. 253.)

Sp. gr. of $\text{GdSO}_4 + \text{Aq}$ at 25° .

Concentration of $\text{GdSO}_4 + \text{Aq}$	Sp. gr.
1—normal	1.0451
$\frac{1}{2}$ —"	1.0229
$\frac{1}{4}$ —"	1.0114
$\frac{1}{16}$ —"	1.0027

(Wagner, Z. phys. Ch. 1890, 5. 35.)

Solubility of GdSO_4 in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

$\text{H}_2\text{SO}_4 + \text{Aq}$ % H_2SO_4	100 g. of the solution contain g. GdSO_4	Solid phase
...	8.212	$\text{GdSO}_4 + 6\text{H}_2\text{O}$
5.23	8.429	
9.61	7.944	
18.70	6.603	
34.00	5.631	
40.35	5.773	$\text{GdSO}_4 + 4\text{H}_2\text{O}$
45.51	6.628	
50.63	5.438	
56.59	3.640	
63.24	2.244	
65.24	2.128	
73.64	2.185	

(Wirth, Z. anorg. 1913, 79. 359.)

See also under +2, 4, and $6\text{H}_2\text{O}$.

Insol. in liquid NH_3 . (Gore, Am. Ch. J., 1898, 20. 828.)

+ H_2O . (Levi-Malvano, Z. anorg. 1906, 48. 447.)

+ $2\text{H}_2\text{O}$.

Solubility in H_2O at t° .

t°	G. GdSO_4 per 100 g.	
	H_2O	Solution
80	84.76	45.87
91.4	97.77	49.42
105	118.4	54.21
119	149.3	59.88

(Levi-Malvano.)

+ $4\text{H}_2\text{O}$. Very sol. in H_2O .

Sol. in its own weight of H_2O at 14° , and in every proportion of boiling H_2O . Less sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ than in water. (Debray, A. ch. (3) 44. 25.)

Solubility in H_2O at t° .

t°	g. GdSO_4 per 100 g.		t°	g. GdSO_4 per 100 g.	
	H_2O	solution		H_2O	solution
30	43.78	30.45	95.4	90.63	47.55
40	46.74	31.85	107.2	115.3	53.58
68	61.95	38.27	111	128.3	56.19
85	76.30	43.28

(Levi-Malvano.)

Sl. sol. in dilute, insol. in absolute alcohol. Can be completely pptd. from $\text{GdSO}_4 + \text{Aq}$ by $\text{HC}_2\text{H}_3\text{O}_2$. (Persoz.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+6H₂O.Solubility in H₂O at t°.

t°	g. GlSO ₄ per 100 g.	
	H ₂ O	solution
31	52.23	34.32
50	60.67	37.77
72.2	74.94	42.85
77.4	81.87	45.01

(Levi-Malvano.)

100 g. of the aqueous solution contain at 25°, 8.212 g. GlSO₄. (Wirth, Z. anorg. 1913, 79. 358.)

Glucinum iron (ferrous) sulphate, GlSO₄, FeSO₄ + 17½H₂O.

Sol. in H₂O. (Klatzo, J. B. 1868. 204.)3GlSO₄, FeSO₄ + 28H₂O. Sol. in H₂O. (Klatzo.)

Do not exist. (Marignac, A. ch. (4) 30. 45.)

Glucinum nickel sulphate, (Gl,Ni)SO₄ + 4H₂O, or 7H₂O.

(Klatzo, J. B. 1868. 205.)

Does not exist. (Atterberg, Sv. V. A. F. 1873, 4. 81.)

Glucinum potassium sulphate, GlSO₄, K₂SO₄ + 2H₂O.

Sl. sol. in cold, slowly but more sol. in hot H₂O. (Debray.)+3H₂O. (Klatzo.)

Glucinum potassium hydrogen sulphate, GlH₂(SO₄)₂, 2K₂SO₄ + 4H₂O.

Easily sol. in H₂O. Partly decomp. by recrystallisation. (Atterberg.)

Glucinum sodium sulphate, 2GlSO₄, 3Na₂SO₄ + 18H₂O.

Sol. in H₂O. (Atterberg.)

Glucinum zinc sulphate, 2GlSO₄, 3ZnSO₄ + 35H₂O.

Sol. in H₂O. (Klatzo, J. B. 1868. 205.)

Does not exist. (Atterberg.)

Gold (auroauric) sulphate, Au₂(SO₄)₃.

Decomp. by moist air, H₂O, glacial acetic acid, or HNO₃ + Aq (1.42 sp. gr.) Insol. in conc. H₂SO₄. (Schottländer, A. 217. 375.)

Gold (auric) sulphate, Au₂O₃, 2SO₃ + H₂O, or Auryl hydrogen sulphate, (AuO)HSO₄.

Deliquescent. Decomp. by H₂O. Sol. in HCl + Aq; not attacked by conc. HNO₃ + Aq. Sol. in 6 pts. conc. H₂SO₄. (Schottländer.)

Gold (auric) potassium sulphate, Au₂(SO₄)₃, K₂SO₄.

Not decomp. immediately by cold H₂O. (Schottländer.)

Hydrazine mercuric sulphate hydrazine, (N₂H₄)₂SO₄, 3HgSO₄, 2N₂H₄.

Insol. in H₂O. Sol. in H₂O containing HCl. (Ferratini, Gazz. ch. it. 1912, 42 (1) 142.)

Hydroxylamine uranyl sulphate,

(NH₂OH)₂, H₂SO₄, 2(UO₂)SO₄ + 5H₂O.Extremely sol. in H₂O from which it can be cryst. (Rimbach, Dissert. 1904.)

Indium sulphate, In₂(SO₄)₃.

Easily sol. in H₂O.+9H₂O. Easily sol. in H₂O.

Indium hydrogen sulphate, InH(SO₄)₂ + 4HO.

Very deliquescent. (Meyer.)

Indium potassium sulphate, InK(SO₄)₂ + 4H₂O.

Sol. in H₂O, but decomp. by boiling. (Rössler, J. pr. (2) 7. 14.)(InO)₂K(SO₄)₂ + 3H₂O. Insol. in H₂O. (Rössler.)

Indium rubidium sulphate,

In₂(SO₄)₃, Rb₂SO₄ + 24H₂O.44.28 pts. are sol. in 100 pts. H₂O at 15°. (Chabrié, C. R. 1901, 132. 473.)Melts in crystal H₂O at 42°. (Locke, Am. Ch. J. 1901, 26. 183.)

Indium sodium sulphate, InNa(SO₄)₂ + 4H₂O.

Sol. in H₂O. (Rössler, J. pr. (2) 7. 14.)

Iodine sulphate, I₂(SO₄)₂.

Ppt. Decomp. by H₂O. (Fichter, Z. anorg. 1915, 91. 140.)

Iodyl sulphate, (IO)₂(SO₄)₂.

Possible composition of Weber's (B. 22. 86) I₂O₅, 3SO₃.

Iridium sulphate.

Sol. in H₂O or alcohol. (Berzelius.)
 Ir(SO₄)₃. Sol. in H₂O. (Rimbach, Z anorg. 1907, 52. 409.)

Iridium potassium sulphate, Ir₂(SO₄)₃, K₂SO₄+24H₂O.

Mpt. 102–103°. Easily sol. in H₂O. (Marino, Z. anorg. 1904, 42. 220.)
 Ir₂(SO₄)₃, 3K₂SO₄. Sol. in H₂O or dil. H₂SO₄+Aq; nearly insol. in sat. K₂SO₄+Aq. (Boisbaudran, C. R. 96. 1406.)

Iridium rubidium sulphate, Ir₂(SO₄)₃.Rb₂.

Sol. in cold; very sol. in hot H₂O. (Marino, Gazz. ch. it. 1903, 32, (2) 511.)
 Mpt. 108–109°. (Marino, Z. anorg. 1904, 42. 219.)

Iridium thallium sulphate, Ir₂(SO₄)₃, Tl₂SO₄+24H₂O.

Very sol. in H₂O. (Marino, Z. anorg. 1904, 42. 222.)

Iron (ferrous) sulphate, FeSO₄.

+H₂O.
 +2H₂O. Not more sol. in H₂O than gypsum. (Mitscherlich.)
 +3H₂O. Sol. in H₂O. (Kane.)
 +4H₂O. Separates from conc. FeSO₄+Aq at 80°.
 +7H₂O. Efflorescent at 33°.

1 pt. FeSO₄+7H₂O dissolves in 1.6 pts. cold, and 0.3 pt. boiling H₂O.

1 pt. FeSO₄+7H₂O dissolves at:
 10° 15° 25° 33° 46° 60° 81° 90° 100°
 in 1.64 1.43 0.87 0.66 0.44 0.38 0.37 0.27 0.3 pts. H₂O
 (Brandes and Firnhaber, Br. Arch. 7. 83.)

When boiled with insufficient H₂O for solution a white hydrate is formed which separates out. Solubility increases up to 87.5°, and then diminishes, owing to the above separation. (Brandes, Pogg. 20. 581.)

Sol. in 2 pts. cold, and 1 pt. boiling H₂O (Fourcroy); sol. in 2 pts. cold H₂O at 18.75° (Abl); sol. in 6 pts. H₂O at moderate heat, and 0.75 pt. at 100°. (Bergmann.)

100 pts. H₂O at 15.5° dissolve 45–50 pts. (Ure's Dict.)

100 pts. H₂O dissolve pts. FeSO₄ at t°.

t°	Pts. FeSO ₄	t°	Pts. FeSO ₄	t°	Pts. FeSO ₄
0	15.8	21	27.4	45	42.9
10	19.9	30	32.6	55	47.0
12	21.3	37	36.5	70	56.5
20	26.0

(Tobler, A. 95. 198.)

100 pts. FeSO₄+Aq sat. at 11–14° contain 17.02% FeSO₄. (v. Hauer, J. pr. 103. 114.)

100 pts. FeSO₄+Aq. sat at 15° contain 37.2% FeSO₄+7H₂O; solution has sp. gr. 1.2232. (Schiff, A. 118. 362.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. FeSO ₄	t°	Pts. FeSO ₄	t°	Pts. FeSO ₄
0	7.9	34	37.1	67	65.1
1	8.7	35	38.0	68	65.0
2	9.5	36	38.9	69	64.9
3	10.4	37	39.8	70	64.8
4	11.2	38	40.7	71	64.7
5	12.0	39	41.7	72	64.5
6	12.9	40	42.6	73	64.4
7	13.7	41	43.5	74	64.2
8	14.5	42	44.4	75	64.0
9	15.3	43	45.3	76	63.7
10	16.2	44	46.2	77	63.4
11	17.0	45	47.1	78	63.1
12	17.9	46	48.1	79	62.7
13	18.7	47	49.0	80	62.3
14	19.5	48	50.0	81	61.9
15	20.4	49	51.0	82	61.5
16	21.2	50	51.9	83	61.0
17	22.1	51	52.9	84	60.4
18	23.0	52	53.8	85	59.8
19	23.8	53	54.8	86	59.2
20	24.7	54	55.7	87	58.5
21	25.6	55	56.7	88	57.7
22	26.4	56	57.7	89	57.0
23	27.3	57	58.7	90	56.2
24	28.1	58	59.7	91	55.3
25	29.0	59	60.7	92	54.3
26	29.9	60	61.7	93	53.3
27	30.8	61	62.7	94	52.2
28	31.7	62	63.7	95	51.0
29	32.6	63	64.8	96	49.6
30	33.5	63.5	65.4	97	48.0
31	34.4	64	65.4	98	46.3
32	35.3	65	65.3	99	44.5
33	36.2	66	65.2	100	42.6

(Mulder, Scheik. Verhandel. 1864. 141.)

If solubility S=pts. anhydrous FeSO₄ in 100 pts. solution, S=13.5+0.3788t from –2° to +65°; S=37.5 constant from 65° to 98°; S=37.5–0.6685t from 98° to 156°. Practically insol. at 156°. (Étard, C. R. 106. 740.)

Sat. FeSO₄+Aq contains at:

–1°	+5°	24°	34°	52°
13.0	15.1	22.7	26.3	32.5% FeSO ₄ ,
60°	67°	77°	86°	94°
36.4	37.7	37.8	37.8	36.7% FeSO ₄ ,
102°	112°	130°	152°	
34.7	28.0	17.3	2.5% FeSO ₄ .	

(Étard, A. ch. 1894, (7) 2. 553.)

100 g. H₂O dissolve 26.69 g. FeSO₄ at 25°. (Stortenbecker, Z. phys. Ch. 1900, 34. 109.)

Solubility of FeSO_4 in H_2O at t° .
100 g. H_2O dissolve g. FeSO_4 .

t°	G. FeSO_4	t°	G. FeSO_4
0.00	15.65	52.00	50.20
10.00	20.51	54.03	52.07
15.25	23.86	60.01	54.95
20.13	26.58	65.00	55.59
25.02	29.60	68.02	52.31
30.03	32.93	70.04	56.08
35.07	36.87	77.00	45.90
40.05	40.20	80.41	43.58
45.18	44.32	85.02	40.46
50.21	48.60	90.13	37.27

$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ is stable from -1.82° to $+56.6^\circ$; $\text{FeSO}_4 \cdot 4\text{H}_2\text{O}$ from 56.6° to 64.4° ; $\text{FeSO}_4 \cdot \text{H}_2\text{O}$ above this point

(Fraenckel, Z. anorg. 1907, 55, 228.)

$\text{FeSO}_4 + \text{Aq}$ sat. at 30° contains 24.9 g. FeSO_4 in 100 g. of solution. (Schreinemakers, Z. phys. Ch. 1912, 71, 110.)

Sp. gr. of $\text{FeSO}_4 + \text{Aq}$ at 15° .

$c_c = \%$ $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$.

c_c	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.005	15	1.082	28	1.161
2	1.011	16	1.088	29	1.168
3	1.016	17	1.094	30	1.174
4	1.021	18	1.100	31	1.180
5	1.027	19	1.106	32	1.187
6	1.032	20	1.112	33	1.193
7	1.037	21	1.118	34	1.200
8	1.043	22	1.125	35	1.206
9	1.048	23	1.131	36	1.213
10	1.054	24	1.137	37	1.219
11	1.059	25	1.143	38	1.226
12	1.065	26	1.149	39	1.232
13	1.071	27	1.155	40	1.239
14	1.077				

(Gerlach, Z. anal. 8, 287.)

Sp. gr. 16.6° of sat. solution = 1.219. (Greenish and Smith, Pharm. J. 1903, 71, 881.)

Sat. $\text{FeSO}_4 + \text{Aq}$ boils at 102.2° (Griffiths), and solution contains 64% FeSO_4 . Crust forms at 102.3° ; highest temp. observed, 104.8° . (Gerlach, Z. anal. 26, 426.)

B-pt. of $\text{FeSO}_4 + \text{Aq}$ containing pts. FeSO_4 to 100 pts. H_2O

B-pt.	Pts. FeSO_4	B-pt.	Pts. FeSO_4
100.5°	17.7	101.5°	50.4
101.0	34.4	101.6	53.2

(Gerlach, Z. anal. 26, 433.)

100 g. of the sat. solution contain 22.84 g. FeSO_4 at 25° (Wirth, Z. anorg. 1913, 79, 364.)

Sol. in hot $\text{HCl} + \text{Aq}$. (Kane.)
Somewhat sol. in conc. H_2SO_4 . (Bussy and Lecann.)

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

$\text{H}_2\text{SO}_4 + \text{Aq}$ Normality	100 g. of the solution contain g. FeSO_4	Solid phase
0	22.84	$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$
2.25	19.03	
6.685	13.40	
10.2	10.30	
12.46	7.26	$\text{FeSO}_4 + \text{H}_2\text{O}$
15.15	4.015	
19.84	0.1522	

(Wirth, Z. anorg. 1913, 79, 364.)

More sol. in water containing NO than in pure H_2O . (Gay, Bull. Soc. (2) 44, 175)

Completely pptd. from $\text{FeSO}_4 + \text{Aq}$ by glacial $\text{HC}_2\text{H}_3\text{O}_2$. (Persoz.)

Solubility in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$. See under $(\text{NH}_4)_2\text{SO}_4$.

Solubility in $\text{Li}_2\text{SO}_4 + \text{Aq}$ at 30°

Composition of the solution		Solid phase
% by wt. FeSO_4	% by wt. Li_2SO_4	
24.87	0	$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$
22.45	4.00	"
21.15	5.58	"
18.79	11.16	"
16.51	15.81	"
16.18	16.52	$\text{FeSO}_4 \cdot 7\text{H}_2\text{O} + \text{Li}_2\text{SO}_4 \cdot \text{H}_2\text{O}$
16.04	16.49	"
15.39	16.80	$\text{Li}_2\text{SO}_4 \cdot \text{H}_2\text{O}$
12.68	18.31	"
5.32	22.15	"
3.74	23.15	"
0	25.1	"

(Schreinemakers, Z. phys. Ch. 1910, 71, 110)

Solubility of $\text{FeSO}_4 \cdot \text{H}_2\text{O} + \text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$ in 100 g. H_2O at t°

t°	Grams FeSO_4	Grams Na_2SO_4
0	18.05	6.13
15.5	25.05	15.97

(Koppel, Z. phys. Ch. 1905, 52, 405)

See also under $\text{FeNa}_2(\text{SO}_4)_2$

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20, 828.)

Insol. in liquid CO_2 . (Büchner, Z. phys. Ch. 1906, 54, 874.)

100 pts. sat. solution of FeSO_4 in 40° alcohol contains 0.3% FeSO_4 . (Schiff.)

Insol. in alcohol of 0.905 sp. gr. or less. (Anthon, J. pr. 14. 125.)

Alcohol and H_2SO_4 precipitate FeSO_4 from $\text{FeSO}_4 + \text{Aq}$, also glacial acetic acid.

Anhydrous FeSO_4 is insol. in acetone. (Krug and M'Elroy, 1893.)

Insol. in acetone. (Fidmann, C. C. 1899, II. 1014.)

100 g. sat. solution in glycol contain 6.0 g. FeSO_4 at ord. temp. (de Coninck.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1904, 37. 3601.)

Iron (ferrous) sulphate, acid, 2FeO , 3SO_3 , $2\text{H}_2\text{O}$.

This salt exists in contact with solutions containing $\text{SO}_3 + 1.637 \text{ H}_2\text{O}$ to about $\text{SO}_3 + 2.186 \text{ H}_2\text{O}$. (Kenrick, J. phys. Chem. 1908, 12. 704.)

FeO , $2\text{SO}_3 + \text{H}_2\text{O}$. This compd. exists with solutions containing $\text{SO}_3 + 1.342 \text{ H}_2\text{O}$ to (about) $\text{SO}_3 + 1.595 \text{ H}_2\text{O}$. (Kenrick.)

FeO , $4\text{SO}_3 + 3\text{H}_2\text{O}$. This compd. is stable with solutions containing from $\text{SO}_3 + 1.122 \text{ H}_2\text{O}$ to (about) $\text{SO}_3 + 1.342 \text{ H}_2\text{O}$. Rapidly sol. in H_2O with ppt. of $\text{FeSO}_4 + \text{H}_2\text{O}$. (Kenrick.)

Min. *Melanterite*.

Iron (ferric) sulphate, basic, $10\text{Fe}_2\text{O}_3$, $\text{SO}_3 + \text{H}_2\text{O}$.

(Athanasesco, C. R. 103. 27.)

$6\text{Fe}_2\text{O}_3$, $\text{SO}_3 + 10\text{H}_2\text{O}$. Insol. in H_2O . Sl. sol. in warm $\text{HCl} + \text{Aq}$. (Scheerer, Pogg. 45. 188.)

$4\text{Fe}_2\text{O}_3$, $\text{SO}_3 + 11\text{H}_2\text{O}$. (Anthon, Repert. 81. 237.)

$3\text{Fe}_2\text{O}_3$, $\text{SO}_3 + 4\text{H}_2\text{O}$. Insol. in H_2O . Rather easily sol. in acids. (Scheerer, Pogg. 44. 453; Meister, B. 8. 771.)

$2\text{Fe}_2\text{O}_3$, $\text{SO}_3 + 6\text{H}_2\text{O}$. When pptd. from cold solutions, is sol. in $\text{Fe}_2(\text{SO}_4)_3 + \text{Aq}$, but insol. therein when pptd. from hot solutions. (Maus.)

Only basic sulphate which is a true chemical compound. (Pickering, Chem. Soc. 37. 807.)

Min. *Glockerite*. Insol. in H_2O . Sol. in conc. H_2SO_4 .

+ $7\text{H}_2\text{O}$. (Meister.)

+ $8\text{H}_2\text{O}$. (Mühlhauser.)

+ $15\text{H}_2\text{O}$. Min. *Pissophanite*.

Fe_2O_3 , $\text{SO}_3 = (\text{FeO})_2\text{SO}_4 + 3\text{H}_2\text{O}$. Insol. in H_2O . (Soubeiran, A. ch. 44. 329.)

$3\text{Fe}_2\text{O}_3$, $4\text{SO}_3 + 9\text{H}_2\text{O}$. (Athanasesco.)

$2\text{Fe}_2\text{O}_3$, $3\text{SO}_3 + 8\text{H}_2\text{O}$. Insol. in H_2O . (Wittstein.)

+ $18\text{H}_2\text{O}$. Min. *Fibroferrite*. Sl. sol. in cold, more easily in hot H_2O .

Fe_2O_3 , $2\text{SO}_3 + 10\text{H}_2\text{O}$. Min. *Stypticite*. + $15\text{H}_2\text{O}$. Sol. in H_2O ; decomp. by heat or evaporation. (Muck, J. pr. 99. 103.)

$2\text{Fe}_2\text{O}_3$, $5\text{SO}_3 + 13\text{H}_2\text{O}$. Min. *Copiapite*.

$\text{Fe}_4\text{S}_2\text{O}_3 + 18\text{H}_2\text{O} = 2\text{Fe}_2\text{O}_3$, $5\text{SO}_3 + 18\text{H}_2\text{O}$. *a-Copiapit*. This salt is in equilibrium at 25°

with solutions in which the molecular ratio $\text{Fe}_2\text{O}_3 : \text{SO}_3$ lies between 1 : 2.889 and 1 : 2.614. (Wirth, Z. anorg. 1914, 87. 37.)

$(\text{OH})\text{Fe}_2(\text{SO}_4)_4 + 13\text{H}_2\text{O}$. *β -Copiapit*. This salt is in equilibrium at 25° with solutions in which the molecular ratio $\text{Fe}_2\text{O}_3 : \text{SO}_3$ lies between 1 : 3.472 and 1 : 2.889. (Wirth, Z. anorg. 1914, 87. 37.)

According to Pickering (Chem. Soc. 37. 807), all basic ferric sulphates are mixtures excepting $2\text{Fe}_2\text{O}_3$, SO_3 .

Iron (ferric) sulphate, $\text{Fe}_2(\text{SO}_4)_3$.

Anhydrous. Slowly deliquescent. Nearly insol. in H_2O , and $\text{HCl} + \text{Aq}$. Insol. in conc. H_2SO_4 . Very rapidly sol. in $\text{FeSO}_4 + \text{Aq}$, even when very dil. (Barreswil, C. R. 20. 1366.)

Sp. gr. of $\text{Fe}_2(\text{SO}_4)_3 + \text{Aq}$. According to F = Franz at 17.5° (J. pr. (2) 5. 280); G = Gerlach at 15° (Z. anal. 28. 494); H = Hager at 18° (Z. anal. 27. 280).

	5	10	15	20 % $\text{Fe}_2(\text{SO}_4)_3$
F	1.0426	1.0854	1.1324	1.1826
G	...	1.096	...	1.205
H	1.046	1.097	1.151	1.208
	25	30	35	40 % $\text{Fe}_2(\text{SO}_4)_3$
F	1.2426	1.3090	1.3782	1.4506
G	...	1.331	...	1.478
H	1.271	1.337	1.411	1.490
	45	50	55	60 % $\text{Fe}_2(\text{SO}_4)_3$
F	1.5298	1.6148	1.7050	1.8006
G	...	1.650

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

$\text{H}_2\text{SO}_4 + \text{Aq}$ Normality	100 g. of the solution contain g. $\text{Fe}_2(\text{SO}_4)_3$
2.25	25.02
6.685	14.58
19.84	0.05

(Wirth, Z. anorg. 1913, 79. 364.)

Solubility in $\text{Al}_2(\text{SO}_4)_3 + \text{Aq}$ at 25° .

100 g. of the solution contain	
G. $\text{Al}_2(\text{SO}_4)_3$	G. $\text{Fe}_2(\text{SO}_4)_3$
...	44.97
2.342	42.44
5.200	38.83
6.626	35.82
8.819	34.02
10.03	32.42
*10.23	31.90
10.70	31.91

*Solution sat. with respect to both salts. (Wirth and Bakke, Z. anorg. 1914, 87. 48.)
See also under $\text{Al}_2(\text{SO}_4)_3$.

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 828.)

Completely pptd. from $\text{Fe}_2(\text{SO}_4)_3 + \text{Aq}$ by $\text{HC}_2\text{H}_3\text{O}_2$. Sol. to large extent in alcohol.

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in acetone.

$+x\text{H}_2\text{O}$. Very deliquescent, and sol. in H_2O . Conc. $\text{Fe}_2(\text{SO}_4)_3 + \text{Aq}$ may be boiled without decomp., but dil. solutions are decomp. on heating. A solution containing 1 pt. salt to 100 pts. H_2O becomes turbid at 76° ; 1 pt. to 200 pts., at 56° ; 1 pt. to 400 pts., at 47° ; 1 pt. to 800 pts., at 40° ; 1 pt. to 1000 pts., at 38° ; 1 pt. to 10,000 pts., at 14° . (Scheerer.)

$+9\text{H}_2\text{O}$. Min. *Coquimbite*.

81.43 pts. are sol. in 18.57 pts. H_2O . (Wirth, Z. anorg. 1914, 87. 23.)

This salt is stable at 25° only in contact with solutions in which the molecular ratio $\text{Fe}_2\text{O}_3 : \text{SO}_3$ lies between 1 : 3.472 and 1 : 6.699. (Wirth, Z. anorg. 1914, 87. 35.)

$+10\text{H}_2\text{O}$. Slowly sol. in H_2O . (Oudemans, R. t. c. 3. 331.)

Iron (ferroferric) sulphate, 6FeSO_4 , $\text{Fe}_2(\text{SO}_4)_3 + 60\text{H}_2\text{O}$.

Sol. in all proportions in H_2O . (Poumarède, C. R. 18. 854.)

3FeSO_4 , $2\text{Fe}_2(\text{SO}_4)_3 + 12\text{H}_2\text{O}$. Decomp. by H_2O . Easily sol. in dil. $\text{HCl} + \text{Aq}$. Insol. in alcohol. (Abich, 1842.)

FeSO_4 , $\text{Fe}_2(\text{SO}_4)_3 + 12\text{H}_2\text{O}$. Min. *Voltaite*. Difficultly sol. in H_2O .

FeO , Fe_2O_3 , $6\text{SO}_3 + 15\text{H}_2\text{O}$. Deliquescent. (Lefort, J. Pharm. (4) 10. 87.)

Iron (ferrous) pyrosulphate, FeS_2O_7 .

Deliquescent. Decomp. by H_2O . (Bolas, Chem. Soc. (2) 12. 212.)

Iron (ferric) hydrogen sulphate, $\text{Fe}_2(\text{SO}_4)_3$, $\text{H}_2\text{SO}_4 + 8\text{H}_2\text{O}$.

This salt is in equilibrium at 25° only with solutions where the molecular ratio $\text{Fe}_2\text{O}_3 : \text{SO}_3$ is more acid than 1 : 6.699. (Wirth and Bakke, Z. anorg. 1914, 87. 34.)

Sat. solution of $\text{Fe}_2(\text{SO}_4)_3$, $\text{H}_2\text{SO}_4 + 8\text{H}_2\text{O}$ in abs. alcohol at 25° contains 8 g. $\text{Fe}_2\text{O}_3 + 17.18$ g. SO_3 per 100 g. sat. solution. (Wirth and Bakke.)

Decomp. by H_2O . (Recoura, C. R. 1903, 137. 118.)

In contact with solutions containing 25% to 28% SO_3 at 25° , the stable solid is Fe_2O_3 , $3\text{SO}_3 + 10\text{H}_2\text{O}$. In contact with solutions containing more than 28%, the stable solid is Fe_2O_3 , $4\text{SO}_3 + 10\text{H}_2\text{O}$. (Cameron and Robinson, J. phys. Chem. 1907, 11. 650.)

Iron (ferroferric) hydrogen sulphate, $\text{Fe}_2(\text{SO}_4)_3$, FeSO_4 , $2\text{H}_2\text{SO}_4$.

Insol. in H_2O , but slowly decomp. thereby. Sol. in H_2SO_4 . (Étard, C. R. 87. 602.)

Iron (ferrous) hydrazine sulphate, $\text{FeH}_2(\text{SO}_4)_2$, $2\text{N}_2\text{H}_4$.

1 pt. is sol. in 325 pts. H_2O at 12° . (Curtis, J. pr. 1894, (2) 50. 331.)

Iron (ferrous) magnesium sulphate, FeSO_4 , $\text{MgSO}_4 + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Schiff.)

Iron (ferric) magnesium sulphate, $\text{Fe}_2(\text{SO}_4)_3$, $\text{MgSO}_4 + 24\text{H}_2\text{O}$.

(Bastick.)

Iron (ferrous) magnesium potassium sulphate, $2\text{K}_2\text{SO}_4$, FeSO_4 , $\text{MgSO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Iron (ferric) manganous hydrogen sulphate, $\text{Fe}_2(\text{SO}_4)_3$, 2MnSO_4 , H_2SO_4 .

Insol. in cold H_2O . (Étard.)

$\text{Fe}_2(\text{SO}_4)_3$, 2MnSO_4 , $3\text{H}_2\text{SO}_4$. Sol. in H_2O . (Étard, C. R. 86. 1399.)

Iron (ferric) manganic sulphate, $\text{Fe}_2(\text{SO}_4)_3$, $\text{Mn}_2(\text{SO}_4)_3$.

Insol. in cold H_2O ; decomp. by hot H_2O and $\text{HCl} + \text{Aq}$. (Étard.)

Iron (ferrous) manganous potassium sulphate, FeSO_4 , MnSO_4 , $2\text{K}_2\text{SO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Iron (ferrous) nickel sulphate, 2FeSO_4 , 2NiSO_4 , H_2SO_4 .

(Étard, C. R. 87. 602.)

Iron (ferric) nickel sulphate, $\text{Fe}_2(\text{SO}_4)_3$, NiSO_4 , $2\text{H}_2\text{SO}_4$.

Insol. in H_2O , but gradually decomp. thereby. (Étard, C. R. 87. 602.)

Iron (ferrous) nickel potassium sulphate, FeSO_4 , NiSO_4 , $2\text{K}_2\text{SO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl, A. 94. 57.)

Iron (ferrous) potassium sulphate, FeSO_4 , K_2SO_4 .

$+2\text{H}_2\text{O}$. (Marignac, Ann. Min. (5) 2. 19.)

Solubility in H_2O at t° .

t°	% $\text{FeK}_2(\text{SO}_4)_2$	t°	% FeK_2SO_4
0.5	22.79	80	42.34
17.2	31.98	90	42.73
40.1	40.86	95	41.01
60	42.63

(Küster and Thiel, Z. anorg. 1899, 21. 116.)

+4H₂O.Solubility in H₂O at t°.

t°	% FeK ₂ (SO ₄) ₂	t°	% FeK ₂ (SO ₄) ₂
0.5	22.94	80	40.46
17.2	26.79	90	43.82
40.1	32.41	95	44.11
60	35.68

(Küster and Thiel.)

+6H₂O. 100 pts. H₂O dissolve at t°:

0° 10° 14.5° 16° 25°
19.6 24.5 29.1 30.9 36.5 pts. anhydrous salt,

35° 40° 55° 65° 70°
41 45 56 59.3 64.2 pts. anhydrous salt.
(Tobler, A. 95. 193.)

Solubility in H₂O at t°.

t°	% FeK ₂ (SO ₄) ₂	t°	% FeK ₂ (SO ₄) ₂
0.5	18.36	60	42.93
17.2	25.16	80	45.29
40.1	36.72

(Küster and Thiel.)

Iron (ferric) potassium sulphate, basic,
4Fe₂O₃, K₂O, 7SO₃+9H₂O=4(Fe₂O₃,
2H₂O, SO₃), K₂SO₄+7H₂O.

Insol. in boiling H₂O. Sl. sol. in HCl+aq, more readily in aqua regia. (Rammelsberg.)
3Fe₂O₃, K₂O, 4SO₃+6H₂O=K(FeO)₃(SO₄)₂+3H₂O. Min. *Jarosite*.

Fe₂O₃, H₂O, 2SO₃, 2K₂SO₄+5H₂O. Sol. in 6 pts. cold H₂O. Solution soon decomposes. (Maus, Pogg. 11. 78.)

Sol. in 12.5 pts. H₂O at 10°. (Anthon, Rept. 76. 361.)

Formula is given as 3Fe₂O₃, 5K₂O, 12SO₃+18H₂O by Marignac.

3Fe₂O₃, 6SO₃, 2K₂SO₄+22H₂O. Sol. when moist in H₂O. Solution soon decomposes. Insol. in alcohol. (Soubeiran, A. ch. 44. 329.)

3Fe₂O₃, 7SO₃, 5K₂SO₄+12H₂O, and +17H₂O. (Scheerer, Pogg. 87. 81.)

2Fe₂O₃, 5SO₃, 3K₂SO₄+9H₂O. (S.)
3Fe₂O₃, 8SO₃, 4K₂SO₄+20H₂O and 24H₂O. (S.)

Iron (ferric) potassium sulphate, K₂SO₄, 2Fe₂(SO₄)₃.

Insol. in H₂O, but is gradually decomp. thereby. (Grimm and Ramdohr, A. 98. 127.)
+2H₂O. Nearly insol. in H₂O. (Weinland, Z. anorg. 1913, 84. 364.)

K₂Fe₂(SO₄)₄+24H₂O. *Iron alum*.
Sol. in 5 pts. H₂O at 12.5°. (Anthon.)

Aqueous solution is decomp. by heating. Insol. in alcohol.

Sp. gr. of aqueous solution. According to

G=Gerlach, at 15° (Z. anal. 28. 496); F=Franz, at 17.5° (J. pr. (2) 5. 288), containing:

5 10 15% K₂Fe₂(SO₄)₄+24H₂O,
F 1.0268 1.0466 1.0672
G 1.025 1.0507 1.0773

20 25 30% K₂Fe₂(SO₄)₄+24H₂O,
F 1.0894 1.1136 1.1422
G 1.1050 1.1340 1.1645

35 % K₂Fe₂(SO₄)₄+24H₂O.
G 1.1967.

Melts in crystal H₂O at 28°. (Locke, Am. Ch. J. 1901, 26. 183.)

Fe₂(SO₄)₃, 3K₂SO₄. Insol. in H₂O, but slowly decomp. thereby. (Étard, C. R. 84. 1089.)

Iron (ferric) potassium sulphate sulphite.

See Sulphite sulphate, ferric potassium.

Iron (ferrous) potassium zinc sulphate, FeSO₄, 2K₂SO₄, ZnSO₄+12H₂O.

Sol. in H₂O.

Iron (ferrous) rubidium sulphate, FeSO₄, Rb₂SO₄+6H₂O.

Sol. in H₂O. (Tutton, Chem. Soc. 63. 337.)
1 l. H₂O dissolves 242 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Iron (ferric) rubidium sulphate, Rb₂Fe₂(SO₄)₃+24H₂O.

Solubility in H₂O.

Temp.	G. per litre	Gram mols. of anhydrous salt per litre
25	97.4	0.294
30	202.4	0.617
35	Basic salt formed	...

(Locke, Am. Ch. J. 1901, 26. 180.)

Iron (ferrous) sodium sulphate, FeSO₄, Na₂SO₄+4H₂O.

Sol. in H₂O. (Marignac, Ann. Min. (5) 9. 25.)

Solubility of FeNa₂(SO₄)₂+4H₂O in 100 g. H₂O at t°.

t°	Grams FeSO ₄	Grams Na ₂ SO ₄
21.8	24.34	22.51
24.92	23.62	22.04
34.95	23.91	21.83
40	24.01	22.62

(Koppel, Z. phys. Ch. 1905, 52. 408.)

Solubility of $\text{FeNa}_2(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ in 100 g. H_2O at t° .

t°	grams FeSO_4	grams Na_2SO_4	t°	grams FeSO_4	grams Na_2SO_4
18.8	26.63	20.28	31°	33.99	14.41
23	28.82	18.40	35°	35.66	13.85
27	30.95	16.68	40°	39.98	11.92

(Koppel.)

Solubility of $\text{FeNa}_2(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$ in 100 g. H_2O at t° .

t°	grams FeSO_4	grams Na_2SO_4	t°	grams FeSO_4	grams Na_2SO_4
18.8	27.23	22.16	28°	11.28	35.94
23	20.31	26.48	31°	6.95	44.75

(Koppel.)

Solubility of $\text{FeNa}_2(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{Na}_2\text{SO}_4$ (anhydrous) in 100 g. H_2O at t° .

t°	grams FeSO_4	grams Na_2SO_4
35	6.16	46.58
40	6.27	46.99

(Koppel.)

See also under FeSO_4 .

Iron (ferric) sodium sulphate, basic, $2\text{Na}_2\text{O}$, Fe_2O_3 , $4\text{SO}_3 + 7\text{H}_2\text{O}$.

Only sl. sol. in H_2O with decomp. (Skrabal, Z. anorg. 1904, 38. 319.)

+ $8\text{H}_2\text{O}$. Min. *Urusite*. Insol. in H_2O ; easily sol. in $\text{HCl} + \text{Aq}$.

$3\text{Na}_2\text{O}$, Fe_2O_3 , $6\text{SO}_3 + 6\text{H}_2\text{O}$. Sol. in H_2O with decomp. (Skrabal.)

$4\text{Fe}_2\text{O}_3$, Na_2O , $5\text{SO}_3 + 9\text{H}_2\text{O}$.

Insol. in H_2O ; difficultly sol. in $\text{HCl} + \text{Aq}$. (Scheerer, Pogg. 45. 190.)

Iron (ferric) sodium sulphate, $[\text{Fe}(\text{SO}_4)_3]\text{Na}_3 + 3\text{H}_2\text{O}$.

Ppt. Nearly insol. in H_2O . (Weinland, Z. anorg. 1913, 84. 365.)

Iron (ferrous) thallium sulphate, FeSO_4 , $\text{Tl}_2\text{SO}_4 + 6\text{H}_2\text{O}$.

Easily decomp. by solution in H_2O . (Willm, A. ch. (4) 5. 56.)

Iron (ferric) thallium sulphate, $\text{Tl}_2\text{Fe}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

Not efflorescent. Very easily sol. in H_2O .

361.5 g. anhydrous, or 646 g. hydrated salt are sol. in 1 l. H_2O at 25° , or 0.799 mol. of the anhydrous salt is sol. in 1 l. H_2O at 25° . Melts in crystal H_2O at 37° . (Locke, Am. Ch. J. 1901, 26. 175.)

Iron (ferrous) zinc sulphate, FeSO_4 , $\text{ZnSO}_4 + 14\text{H}_2\text{O}$.

2FeSO_4 , 2ZnSO_4 , H_2SO_4 . (Étard, C. R. 87. 602.)

Iron (ferric) zinc sulphate, $\text{Fe}_2(\text{SO}_4)_3$, $\text{ZnSO}_4 + 24\text{H}_2\text{O}$.

(Bastick.)

Iron (ferrous) sulphate nitric oxide, FeSO_4 , NO .

The solubility of NO in $\text{FeSO}_4 + \text{Aq}$ is diminished by the presence of H_2SO_4 , HCl , phosphoric acid and by the presence of certain salts. (Manchot, A. 1910, 372. 157.)

$\text{Fe}(\text{NO})\text{SO}_4$, $\text{FeSO}_4 + 13\text{H}_2\text{O}$. Decomp. in the air. Sol. in water. (Manchot.)

Lanthanum sulphate, basic, $2\text{La}_2\text{O}_3$, $3\text{SO}_3 + 3\text{H}_2\text{O}$.

Precipitate. (Frerichs and Smith.)

Formula is $3\text{La}_2\text{O}_3$, $\text{SO}_3 + x\text{H}_2\text{O}$. (Cleve, B. 11. 910.)

Lanthanum sulphate, $\text{La}_2(\text{SO}_4)_3$.

Anhydrous. Much less sol. in warm than in cold H_2O . 1 pt. is sol. in less than 6 pts. H_2O , if added in small portions thereto at $2-3^\circ$, and the temperature not allowed to rise to 13° ; but if heated to 30° , $\text{La}_2(\text{SO}_4)_3 + 9\text{H}_2\text{O}$ separates out until the solution is solid (Mosander.)

100 pts. H_2O dissolve 2.208 pts. $\text{La}_2(\text{SO}_4)_3$ at 16.5° ; 2.130 pts. at 18° ; 1.641 pts. at 34° .

See also under $+9\text{H}_2\text{O}$.

The solubility of $\text{La}_2(\text{SO}_4)_3$ in H_2O is diminished by the presence of $(\text{NH}_4)_2\text{SO}_4$, K_2SO_4 , or Na_2SO_4 . (Barre, C. R. 1910, 151. 871.)

Solubility in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ at 18° .

Pts. per 100 pts. H_2O		Solid phase
$(\text{NH}_4)_2\text{SO}_4$	$\text{La}_2(\text{SO}_4)_3$	
0.00	2.130	$\text{La}_2(\text{SO}_4)_3 + 9\text{H}_2\text{O}$
4.011	0.393	$\text{La}_2(\text{SO}_4)_3$, $(\text{NH}_4)_2\text{SO}_4 + 2\text{H}_2\text{O}$
8.727	0.279	"
18.241	0.253	"
27.887	0.476	"
36.112	0.277	"
47.486	0.137	$2\text{La}_2(\text{SO}_4)_3$, $5(\text{NH}_4)_2\text{SO}_4$
53.823	0.067	$\text{La}_2(\text{SO}_4)_3$, $5(\text{NH}_4)_2\text{SO}_4$
65.286	0.0117	"
73.782	0.0033	"

(Barre.)

Solubility in $\text{K}_2\text{SO}_4 + \text{Aq}$ at 16.5° .

Pts. per 100 pts. H_2O		Solid phase
K_2SO_4	$\text{La}_2(\text{SO}_4)_3$	
0.00	2.198	$\text{La}_2(\text{SO}_4)_3 + 9\text{H}_2\text{O}$
0.247	0.727	$\text{La}_2(\text{SO}_4)_3$, $\text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}$
0.496	0.269	"
0.846	0.185	"
1.029	0.054	$\text{La}_2(\text{SO}_4)_3$, $5\text{K}_2\text{SO}_4$
1.516	0.022	"

(Barre.)

Solubility in $\text{Na}_2\text{SO}_4 + \text{Aq}$ at 18° .

Pts. per 100 pts. H_2O		Solid phase
Na_2SO_4	$\text{La}_2(\text{SO}_4)_3$	
0.00	2.130	$\text{La}_2(\text{SO}_4)_3 + 9\text{H}_2\text{O}$
0.395	0.997	$\text{La}_2(\text{SO}_4)_3, \text{Na}_2\text{SO}_4$
0.689	0.353	"
0.774	0.299	"
1.136	0.129	"
2.480	0.044	"
3.802	0.019	"
5.548	0.016	"

(Barre.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+9 H_2O . Sol. in 42.5 pts. H_2O , calculated as anhydrous salt, at 23° , and 115 pts. H_2O at 100° . (Mosander.)

Solubility in H_2O .

100 pts. H_2O dissolve pts. $\text{La}_2(\text{SO}_4)_3$ at t° .

t°	Pts. $\text{La}_2(\text{SO}_4)_3$
0	3.02
14	2.60
30	1.90
50	1.49
75	0.94
100	0.68

(Muthmann and Rölig, B. 1898, 31. 1723.)

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

Normality H_2SO_4	In 100 g. of the liquid are dissolved		Solid phase
	g. oxide	g. sulphate	
0	1.43	2.483	$\text{La}_2(\text{SO}_4)_3 + 9\text{H}_2\text{O}$
0.505	1.69	2.934	"
1.10	1.796	3.118	"
2.16	1.818	3.156	"
3.39	1.42	2.465	"
4.321	1.11	1.927	"
6.685	0.5309	0.9217	"
9.68	0.2659	0.4617	"
12.60	0.2136	0.3709	"
15.15	0.177	0.3073	"

(Wirth, Z. anorg. 1912, 76. 189.)

Lanthanum hydrogen sulphate, $\text{La}(\text{SO}_4\text{H})_3$.
(Brauner, Z. anorg. 1904, 38. 330.)

Lanthanum potassium sulphate, $\text{La}_2(\text{SO}_4)_3, \text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}$.

$\text{La}_2(\text{SO}_4)_3, 5\text{K}_2\text{SO}_4$. (Barre, C. R. 1910, 151. 872.)

$\text{La}_2(\text{SO}_4)_3, 3\text{K}_2\text{SO}_4$. Sl. sol. in H_2O . Insol. in sat. $\text{K}_2\text{SO}_4 + \text{Aq}$. (Cleve.)

$\text{La}_2(\text{SO}_4)_3, 4\text{K}_2\text{SO}_4$. As above. (Cleve.)

$2\text{La}_2(\text{SO}_4)_3, 9\text{K}_2\text{SO}_4$. As above. (Cleve.)

Lanthanum rubidium sulphate,

$\text{La}_2(\text{SO}_4)_3, \text{Rb}_2\text{SO}_4$.

(Baskerville, J. Am. Chem. Soc. 1904, 26. 67.)

+2 H_2O . (Baskerville.)

$3\text{La}_2(\text{SO}_4)_3, 2\text{Rb}_2\text{SO}_4$. (Baskerville.)

Lanthanum sodium sulphate, $\text{La}_2(\text{SO}_4)_3, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Cleve.) (Barre, C. R. 1910, 151. 872.)

Lead sulphate, basic, $2\text{PbO}, \text{SO}_3$.

Not completely insol. in H_2O . Decomp. by acids, even dil. $\text{HCl}, \text{H}_2\text{O}_2 + \text{Aq}$, with formation of PbSO_4 . (Barfoed, 1869.)

0.050 millimole calc. as Pb is sol. in 1 l. H_2O at 18° . (Pleissner, C. C. 1907, II. 1056.)

$5\text{PbO}, 3\text{SO}_3$. (Frankland, Proc. Roy. Soc. 46. 364.)

$\text{Pb}_3\text{O}_4, 2\text{SO}_3$. (Frankland.)

$3\text{PbO}, \text{PbSO}_4 + \text{H}_2\text{O}$. Ppt. (Strömholm, Z. anorg. 1904, 38, 442.)

$\text{Pb}_4(\text{SO}_4)(\text{OH})_3$. 0.106 millimole calc. as Pb is sol. in 1 liter H_2O at 18° . (Pleissner, C. C. 1907. II, 1056.)

Lead sulphate, PbSO_4 .

Sol. in 22,816 pts. H_2O at 11° . (Fresenius, A. 59. 125.)

Sol. in 31,569 pts. H_2O at 15° . (Rodwell, C. N. 11. 50.)

Sol. in 13,000 pts. H_2O . (Kremers, Pogg. 85. 247.)

Calculated from electrical conductivity of $\text{PbSO}_4 + \text{Aq}$, 1 l. H_2O dissolves 46 mg. PbSO_4 at 18° . (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

4.23×10^{-2} gr. are dissolved in 1 liter of sat. solution at 20° ; 4.41×10^{-2} at 25° . (Böttger, Z. phys. Ch. 1903, 46. 604.)

1 l. H_2O dissolves 41 mg. PbSO_4 at 18° . (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)

0.126 millimole Pb is sol. in 1 liter H_2O at 18° . (Pleissner, C. C. 1907, II. 1056.)

40 mg. are dissolved in 1 l. of sat. solution at 18° . (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

0.0824 g. PbSO_4 is sol. in 1000 cc. H_2O at 18° and also at 100° . The fact that PbSO_4 dissolves in H_2O is ascribed to hydrolysis, and in support of this it is shown that the solubility of hydrated oxide of lead, $\text{PbO}, \text{H}_2\text{O}$, in dil. H_2SO_4 is the same as the solubility of PbSO_4 in H_2O . (Sehnal, C. R. 1909, 148. 1395.)

1 l. H_2O dissolves 26 mg. at 18° ; 30 mg. at 25° ; 38 mg. at 37° . (Beck and Stegmüller, Arb. K. Gesund. Amt. 1910, 34. 447.)

Solubility in H ₂ O at t°. (Millimols per l.)	
t°	PbSO ₄
18	0.126
25	0.144
37	0.183

(Beck and Stegmüller, Arb. K. Gesund. Amt. 1910, **34**. 446.)

Sol. in hot conc. HCl+Aq. (Fresenius.)

Solubility of PbSO₄ in HCl+Aq.

Sp. gr. of HCl+Aq	% HCl in HCl+Aq	Pts. HCl+Aq for 1 pt. PbSO ₄
1.0519	10.602	681.89
1.0800	16.310	281.73
1.1070	22.010	105.65
1.1359	27.525	47.30
1.1570	31.602	35.03

(Rodwell, Chem. Soc. **15**. 59.)

Solubility of PbSO₄ in HCl+Aq at t°.
(Millimols. per l.)

t°	0.1N	0.2N	0.3N	0.4N
18	0.126	1.72	2.67	3.63
25	0.144	2.07	3.14	4.29
37	0.183	2.63	4.06	5.43

(Beck and Stegmüller, Arb. K. Gesund. Amt. 1910, **34**. 446.)

Above measurements in HCl+Aq show solubility directly proportional to the hydrogen ions. (Beck and Stegmüller.)

Sol. in HNO₃+Aq, and more sol. in hot or conc. than in cold or dil. HNO₃+Aq.

Sol. in 172 pts. HNO₃+Aq of 1.144 sp. gr. at 12.5°. (Bischof.)

Pptd. from HNO₃ solution by dil. H₂SO₄+Aq and not by H₂O. (Bischof, **1827**.)

Solubility of PbSO₄ in HNO₃+Aq.

Sp. gr. of HNO ₃ +Aq	% HNO ₃ in HNO ₃ +Aq	Pts. HNO ₃ +Aq for 1 pt. PbSO ₄
1.079	11.55	303.10
1.123	17.50	173.75
1.250	34.00	127.48
1.420	60.00	10282.78

(Rodwell, Chem. Soc. **15**. 59.)

Solubility in HNO₃ at 18°.
(Millimols per l.)

HNO ₃	PbSO ₄
0.1N	0.506
0.2N	0.844
0.3N	1.13
0.4N	1.44

(Beck and Stegmüller.)

Sol. in 36,504 pts. dil. H₂SO₄+Aq. (Fresenius.) See also under solubility in alcohol.

Sl. sol. in conc. H₂SO₄, from which it is partially pptd. by H₂O or completely by alcohol. (Fresenius.)

100 pts. conc. H₂SO₄ dissolve 6 pts. PbSO₄. (Schultz, Pogg. 133. 137.)

Conc. H₂SO₄ dissolves 0.005 pt. PbSO₄. (Ure.)

100 pts. H₂SO₄ dissolve 0.13 pt. PbSO₄, and 100 pts. fuming H₂SO₄ dissolve 4.19 pts. (Struve, Z. anal. 9. 31.)

More sol. in commercial H₂SO₄ than in the more conc. acid. (Hayes.)

100 pts. H₂SO₄+Aq of 1.841 sp. gr. dissolve 0.039 pts. PbSO₄; of 1.793 sp. gr. dissolve 0.011 pt. PbSO₄; of 1.540 sp. gr. dissolve 0.003 pt. PbSO₄.

Presence of SO₂ does not increase the solubility; HNO₃ increases the solubility somewhat, i. e., 100 pts. H₂SO₄+Aq of 1.841 sp. gr. with 5 pts. HNO₃ of 1.352 sp. gr. dissolve 0.044 pt. PbSO₄; 100 pts. H₂SO₄ of 1.749 sp. gr. with 5 pts. HNO₃ of 1.352 sp. gr. dissolve 0.014 pt. PbSO₄; 100 pts. H₂SO₄ of 1.512 sp. gr. with 5 pts. HNO₃ of 1.352 sp. gr. dissolve only a trace.

Nitrous oxides do not increase the action. (Kolb, Dingl. 209. 268.)

Solubility in dil. H₂SO₄+Aq at 18°.
(G. per l.)

H ₂ SO ₄	PbSO ₄	H ₂ SO ₄	PbSO ₄
0	0.0382	0.0245	0.0194
0.0049	0.0333	0.0490	0.0130
0.0098	0.0306	0.4904	0.0052

(Pleissner, Arb. K. Gesund. Amt. 1907, 28. 384.)

A trace of H₂SO₄ has a considerable effect in reducing the solubility of PbSO₄ in H₂O. (Sehnal, C. R. 1909, 148. 1395.)

Solubility in dil. H₂SO₄+Aq at 20°.
(G. per l.)

H ₂ SO ₄	PbSO ₄	H ₂ SO ₄	PbSO ₄
0	0.082	0.0980	0.013
0.0098	0.051	0.4900	0.006
0.0196	0.025	0.9800	0

(Sehnal.)

Pptd. from solution in H₂SO₄ by HCl. (Bolley, A. 91. 113.)

Not more insol. in dil. HC₂H₃O₂+Aq than in H₂O. (Bischof.)

Solubility in other acids is prevented by great excess of H₂SO₄. (Wackenroder.)

Sol. in warm NH₄OH+Aq, separating on cooling. Completely sol. in warm KOH or NaOH+Aq.

Decomp. by boiling with K_2CO_3 , Na_2CO_3 , and $(NH_4)_2CO_3 + Aq$.
Sol. in NH_4 salts + Aq, but repptd. by $H_2SO_4 + Aq$. (Fresenius, A. 59. 125.)
The best solvents of the NH_4 salts are the nitrate, citrate, and tartrate; the two latter should be strongly alkaline with $NH_4OH + Aq$. (Wackenroder.)
Sol. in $NH_4Cl + Aq$ at 12.5–25°.
Sl. decomp. by $NaCl + Aq$. (Bley.)
1 l. sat. $NaCl + Aq$ dissolves 0.66 g. $PbSO_4$. (Becquerel.)
Sol. in 100 pts. cold conc. $NaCl + Aq$, and $PbCl_2$ is deposited after a few hours. (Field.)
Solubility of $PbSO_4$ in $NaCl + Aq$ at 18°.
(Millimols per l.)

NaCl	PbSO ₄
0.1N	0.546
0.2N	0.904
0.3N	1.28
0.4N	1.68

(Beck and Stegmüller, Arb. K. Gesund. Amt. 1910, 34. 446.)

Sol. in $Fe_2Cl_6 + Aq$. (Fresenius, Z. anal. 19. 419.)
Sol. in $Na_2S_2O_3 + Aq$. (Löwe.)
Sol. in $(NH_4)_2SO_4 + Aq$. (Rose.)

Solubility of $PbSO_4 + PbSO_4$, K_2SO_4 in H_2O .

t°	K ₂ SO ₄		Solid phase
	g. in 100 cc. of solution	Mol. in 100 cc. of solution	
0	0.195	0.0112	K_2SO_4 , $PbSO_4 + PbSO_4$
22	0.396	0.0227	"

(Brönsted, Z. phys. Ch. 1911, 77. 316.)
Sol. in 47 pts. $NH_4C_2H_3O_2 + Aq$ (1.036 sp. gr.), and 969 pts. $NH_4NO_3 + Aq$ (1.269 sp. gr.); from the solution in $NH_4C_2H_3O_2$ it is pptd. by H_2SO_4 or K_2SO_4 ; from solution in NH_4NO_3 by K_2SO_4 , but not by H_2SO_4 . (Bischof.)
Sol. in acetates of NH_4 , Na , K , Ca , Al , and Mg . (Mercer.)

Solubility in $NH_4C_2H_3O_2 + Aq$. Excess of $PbSO_4$ was boiled with solution of $NH_4C_2H_3O_2 + Aq$ of varving conc.

G. $NH_4C_2H_3O_2$ in 100 cc.	g. $PbSO_4$ contained in 5 cc. solution		
	Hot	Cooled	Cooled 24 hrs.
28	0.356
30	0.418	...	0.224
32	0.494	0.451	0.242
35	0.513	0.452	...
37	0.529	...	0.238
40	0.539	...	0.263
45	0.555	0.488	...

(Dunnington and Long, Am. Ch. J. 1899, 22. 218.)

Solubility in ammonium acetate + Aq at 25°.

$NH_4C_2H_3O_2$ Millimol. per l.	Solubility of $PbSO_4$	
	Millimols. per l.	g. per l.
0.0	0.134	0.041
103.5	2.10	0.636
207.1	4.55	1.38
414.1	10.10	3.06

(Noyes and Whitcomb, J. Am. Chem. Soc. 1905, 27. 756.)

Solubility in $KC_2H_3O_2 + Aq$ at 25°.
Solid phase, $PbSO_4 + PbK_2(SO_4)_2$.

Composition of the solutions			
% $KC_2H_3O_2$	% $Pb(C_2H_3O_2)_2$	% $KC_2H_3O_2$	% $Pb(C_2H_3O_2)_2$
4.33	2.54	26.58	9.83
9.03	3.55	28.82	11.40
17.81	5.43	28.93	19.41
22.07	5.95		

(Fox, Chem. Soc. 1909, 95. 887.)

100 pts. H_2O containing a drop of $HC_2H_3O_2$ and 2.05 pts. $NaC_2H_3O_2$ dissolve 0.054 pt. $PbSO_4$; containing 8.2 pts. $NaC_2H_3O_2$ dissolve 0.900 pt. $PbSO_4$; containing 41.0 pts. $NaC_2H_3O_2$ dissolve 11.200 pts. $PbSO_4$. (Dibbits, Z. anal. 1874, 13. 139.)

Solubility in $NaC_2H_3O_2 + Aq$ at 25°.

Composition of the solutions			
% Na acetate	% Pb acetate	% Na_2SO_4	% H_2O
6.69	0.78	0.34	92.19
11.76	2.73	1.26	84.25
16.90	5.70	2.49	74.91
19.92	8.24	3.60	68.24
21.51	10.75	4.68	63.10
6.95	0.81	0.35	91.90

The proportion of sulphate in solution in each case corresponded with the amount of Pb present, but was calculated to sodium sulphate, since $Na_2SO_4 + 10H_2O$ cryst. from the solutions on cooling. The solid phase in these solutions was $PbSO_4$.

(Fox, Chem. Soc. 1909, 95. 887.)

Sol. in $Mn(C_2H_3O_2)_2$, $Zn(C_2H_3O_2)_2$, $Ni(C_2H_3O_2)_2$, and $Cu(C_2H_3O_2)_2$, but not in $Hg(C_2H_3O_2)_2$ or $AgC_2H_3O_2 + Aq$.

Solubility in $KC_2H_3O_2 + Aq$ is not less than that in $NaC_2H_3O_2 + Aq$. (Dibbits, Z. anal. 13. 137.)

Insol. in $Pb(C_2H_3O_2)_2 + Aq$. (Smith.)
Sol. in basic lead acetate + Aq, but not in neutral $Pb(C_2H_3O_2)_2 + Aq$. (Stammer, Z. anal. 23. 67.)

12.2 pts. $Ca(C_2H_3O_2)_2$ in very dil. solution dissolve 1 pt. $PbSO_4$. (Städel, Z. anal. 2. 180.)
Sol. in $Al(C_2H_3O_2)_3 + Aq$. (Lennsen.)

Very easily and abundantly sol. in NH_4 tartrate + Aq. (Wöhler, A. **34**. 235.)

Even when native, easily sol. in NH_4 citrate + Aq. (Smith.)

Insol. in alcohol (18%) and H_2SO_4 when NH_4 acetate, K tartrate, or NH_4 succinate are present. Insol. in alcohol (18%) and H_2SO_4 or $(\text{NH}_4)_2\text{SO}_4$ when Na acetate, Na or NH_4 oxalate are present. Sol. in NH_4 dicitrate and K tricitrate in presence of H_2SO_4 ; in NH_4 succinate and NH_4 acetate in presence of $(\text{NH}_4)_2\text{SO}_4$; and in NH_4 citrate in presence of H_2SO_4 or $(\text{NH}_4)_2\text{SO}_4$. (Storer, C. N. **21**. 17.)

Alcohol (59%) alone, or with ethylsulphuric acid or sugar, does not dissolve Pb by 3 months action. (Storer.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329); methyl acetate. (Naumann, B. 1909, **42**. 3790); ethyl acetate. (Naumann, B. 1910, **43**. 314.)

Min. *Anglesite*. Sol. in cold citric acid + Aq. (Bolton, C. N. **37**. 14.)

Lead hydrogen sulphate, PbSO_4 , $\text{H}_2\text{SO}_4 + \text{H}_2\text{O}$.
Decomp. by H_2O .

Lead pyrosulphate, $\text{Pb S}_2\text{O}_7$.
Decomp. by H_2O . (Schultz.)

Lead potassium sulphate, PbSO_4 , K_2SO_4 .
When PbSO_4 is added to potassium acetate + Aq at 25° a double salt, $\text{PbK}_2(\text{SO}_4)_2$ is formed. This salt is insol. in the solution which contains only potassium acetate and lead acetate. (Fox, Chem. Soc. 1909, **95**. 882.)

Decomp. by H_2O . Stable only in solutions of K_2SO_4 , containing at least 0.56% K_2SO_4 at 7° ; 0.62% at 17° ; 1.09% at 50° ; 1.37% at 75° ; 1.69% at 100° . (Barre, C. R. 1909, **149**. 294.)

Lead sulphate chloride, PbSO_4 , $2\text{PbCl}_2 + \text{H}_2\text{O}$.
Insol. in H_2O or $\text{NaCl} + \text{Aq}$. (Becquerel, C. R. **20**. 1523.)

Lead sulphate fluoride, PbSO_4 , 2PbF_2 .
Not decomp. by H_2SO_4 . (Lonyet, C. R. **24**. 434.)

Lithium sulphate, Li_2SO_4 .
More sol. in cold than in hot H_2O .
100 pts. H_2O dissolve 34.6 pts. Li_2SO_4 at 18° . (Wittstein.)

100 pts. H_2O dissolve pts. Li_2SO_4 at t°

t°	Pts. Li_2SO_4	t°	Pts. Li_2SO_4	t°	Pts. Li_2SO_4
0	35.34	45	32.38	100	29.24
20	34.36	65	30.3

(Kremers, Pogg. **95**. 468.)

Sat. $\text{Li}_2\text{SO}_4 + \text{Aq}$ contains at:

-20°	-16°	-15°	-12°
18.4	22.5	22.6	24.4 % Li_2S

-4°	$+15^\circ$	$+90^\circ$
25.7	25.3	23.9 % Li_2SO_4

(Étard, A. ch. 1894, (7) **2**. 547.)

Sat. solution boils at 105° . (Kremer)

Sp. gr. of $\text{Li}_2\text{SO}_4 + \text{Aq}$ at 19.5° contains

6.5	7.4	12.5	15.3 % Li_2S
1.05	1.06	1.098	1.118

22.6	24.4	29.4 % Li_2SO_4
1.167	1.178	1.208

(Kremers, Pogg. **114**. 47.)

Sp. gr. of $\text{Li}_2\text{SO}_4 + \text{Aq}$ at 15° contains $\text{Li}_2\text{SO}_4 = 1.0430$; 10% $\text{Li}_2\text{SO}_4 = 1.087$ (Kohlrausch, W. Ann. **1879**. 1.)

Sp. gr. of $\text{Li}_2\text{SO}_4 + \text{Aq}$ at 25° .

Concentration of $\text{Li}_2\text{SO}_4 + \text{Aq}$	Sp. gr.
1—normal	1.0453
$\frac{1}{2}$ —"	1.0234
$\frac{1}{4}$ —"	1.0115
$\frac{1}{8}$ —"	1.0057

(Wagner, Z. phys. Ch. 1890, **5**. 38.)

Sp. gr. of $\text{Li}_2\text{SO}_4 + \text{Aq}$.

$\frac{1}{2}$ Li_2SO_4 g. in 1000 g. of solution	Sp. gr. $16^\circ 16'$
0	1.000000
2.9198	1.002589
16.0461	1.014093

(Dijken, Z. phys. Ch. 1897, **24**. 100.)

Sp. gr. of $\text{Li}_2\text{SO}_4 + \text{Aq}$ at 20° .

Normality of $\text{Li}_2\text{SO}_4 + \text{Aq}$	% Li_2SO_4	Sp. gr.
2.60	23.48	1.233
1.96	18.53	1.165
1.708	16.41	1.144
1.320	13.01	1.113
0.747	7.71	1.067

(Forchheimer, Z. phys. Ch. 1900, **34**.

Insol. in SO_2 . (Weber, B. **17**. 2497.)
10 ccm. of sat. Li_2SO_4 in absolute H contain approx. 2.719 g. Li_2SO_4 . (Ber Z. phys. Ch. 1910, **72**. 355.)

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 30° .

Composition of the solution		Solid phase
% by wt. H_2SO_4	% by wt. Li_2SO_4	
5.05	22.74	$\text{Li}_2\text{SO}_4, \text{H}_2\text{O}$
12.23	20.45	"
15.37	19.11	"
16.60	19.10	"
32.70	13.37	"
36.90	11.90	"
42.98	10.57	"
48.00	10.20	"
52.72	11.44	"
54.54	12.92	"
55.08	13.69	Li_2SO_4
56.30	13.87	"
61.46	17.10	"
61.82	17.00	"
62.14	17.97	"
62.49	18.89	$\text{Li}_2\text{SO}_4, \text{H}_2\text{SO}_4$
65.70	16.55	"
69.40	13.75	"
77.30	11.31	"
78.23	11.64	"
81.20	13.28	"
81.70	13.85	"
82.30	15.50	"
83.43	15.65	"

(Van Dorp, Z. phys. Ch. 1910, **73**. 289.)

Solution in H_2SO_4 contains 17.2% Li_2SO_4 at 30° . (Van Dorp, Z. phys. Ch. 1913, **86**. 112.)

Solubility of $\text{Li}_2\text{SO}_4 + \text{Th}(\text{SO}_4)_2$ in H_2O at 25° .

Solid phase, $\text{Th}(\text{SO}_4)_2$.
G. in 100 g. H_2O .

Li_2SO_4	$\text{Th}(\text{SO}_4)_2$	Li_2SO_4	$\text{Th}(\text{SO}_4)_2$
0.0	1.722	11.13	11.05
2.57	4.13	13.18	12.54
4.93	6.20	16.12	14.52
6.98	7.95	20.49	16.92
9.23	9.68	16.92	18.87

(Barre, Bull. Soc. 1912, (4) **11**. 647.)

Easily sol. (Kastner), sl. sol. (Berzelius) in alcohol.

Solubility of Li_2SO_4 in alcohol + Aq at 30° .Solid phase $\text{Li}_2\text{SO}_4, \text{H}_2\text{O}$.

G. per 100 g. sat. solution			
$\text{C}_2\text{H}_5\text{OH}$	Li_2SO_4	$\text{C}_2\text{H}_5\text{OH}$	Li_2SO_4
0	25.1	47.28	3.04
11.75	16.16	58.59	1.22
21.19	11.52	69.39	0.4
29.40	8.17	80.74	0
33.31	6.66	94.11	0

(Schreinemakers and van Dorp, Chem. Weekbl. 1906, **3**. 557.)

Insol. in methyl acetate (Naumann, B. 1909, **42**. 3790); ethyl acetate (Naumann, B. 1904, **37**. 3601); acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, **37**. 4329.) + H_2O . Very sl. efflorescent. (Rammelsberg.)

Aq. solution contains 25.1% Li_2SO_4 at 30° . (Schreinemakers, C. C. 1910, I. 1801); 24.3 g. at 50° . (Schreinemakers and Cocheret, Chem. Weekbl. 1905, **2**. 771.)

Lithium hydrogen sulphate, LiHSO_4 .Decomp. by H_2O .Cryst. from H_2SO_4 . (Gmelin.)

$\text{LiH}_2(\text{SO}_4)_2$. Cryst. from H_2SO_4 . (Schultz, Pogg. **133**. 137.)

$\text{Li}_2\text{SO}_4, 7\text{H}_2\text{SO}_4$. (Bergius, Z. phys. Ch. 1910, **72**. 355.)

Lithium potassium sulphate, $\text{Li}_2\text{SO}_4, \text{K}_2\text{SO}_4$.

This is the only compd. of Li_2SO_4 and K_2SO_4 which exists below 100° . (Spielrein, C. R. 1913, **157**. 48.)

$\text{K}_4\text{Li}_2(\text{SO}_4)_2$. (Knobloch.) Has the formula $\text{K}_2\text{Li}_2(\text{SO}_4)_2 + 8\text{H}_2\text{O}$, according to Rammelsberg.

Lithium sodium sulphate, $\text{Na}_2\text{Li}(\text{SO}_4)_2 + 6\text{H}_2\text{O}$. $\text{Na}_4\text{Li}_2(\text{SO}_4)_2 + 9\text{H}_2\text{O}$. $\text{Na}_2\text{Li}_2(\text{SO}_4)_2 + 5\text{H}_2\text{O}$. (Rammelsberg.)

Do not exist. (Troost.)

$\text{Li}_2\text{SO}_4, \text{Na}_2\text{SO}_4 + 5.5\text{H}_2\text{O}$. Exists from 0° – 16° .

+ $3\text{H}_2\text{O}$. Exists from 32° – 100° .

$\text{Li}_2\text{SO}_4, 3\text{Na}_2\text{SO}_4 + 12\text{H}_2\text{O}$. Exists from 16° – 24° .

$4\text{Li}_2\text{SO}_4, \text{Na}_2\text{SO}_4 + 5\text{H}_2\text{O}$. Exists from 24° – 32° .

(Spielrein, C. R. 1913, **157**. 47.)Lithium thallic sulphate, $\text{LiTl}(\text{SO}_4)_2 + 3\text{H}_2\text{O}$.(Meyer and Goldschmidt, C. C. 1903, **V**. 495.)

Lithium titanium sulphate,
 $\text{Li}_2\text{TiO}_2(\text{SO}_4)_2 + 7\text{H}_2\text{O}$.
Less hygroscopic than K compound.
(Mazzuchelli and Pontanelli, C. C. 1909, II. 420.)

Lithium uranyl sulphate, $\text{Li}_2\text{SO}_4, \text{UO}_2\text{SO}_4 + 4\text{H}_2\text{O}$.
(de Coninck, Chem. Soc. 1905, 88. (2) 530.)

Magnesium sulphate basic,
 $6\text{Mg}(\text{OH})_2, \text{MgSO}_4 + 3\text{H}_2\text{O}$.
Sl. sol. in cold or hot H_2O .
Sol. in $\text{HCl} + \text{Aq}$. (Thugutt, Z. anorg. 1892, 2. 150.)

Magnesium sulphate, MgSO_4 .
Anhydrous. Very slowly sol. in H_2O ; sol. in hot conc. H_2SO_4 , less in HCl , and $\text{HNO}_3 + \text{Aq}$.
 $+ \text{H}_2\text{O}$. Min. *Kieserite*. Easily sol. in warm, but slowly dissolved by cold H_2O .
100 g. sat. solution at 83° contain 40.2 g. MgSO_4 . (Geiger, Dissert. 1904.)
 $+ 6\text{H}_2\text{O}$, and $+ 7\text{H}_2\text{O}$. The latter exists in two modifications; (a) hexagonal, and (b) the ordinary or rhombic salt.
 $\text{MgSO}_4 + \text{Aq}$, which on cooling or keeping in closed vessels has deposited $\text{MgSO}_4 + 6\text{H}_2\text{O}$, always contains for 100 pts. H_2O at:

0°	10°	20°
40.75	42.23	43.87 pts. MgSO_4 .

If only hexagonal $\text{MgSO}_4 + 7\text{H}_2\text{O}$ has been deposited, then the mother liquor contains for 100 pts. H_2O at:

0°	10°	20°
34.67	38.71	42.84 pts. MgSO_4 .

Solutions prepared from rhombic $\text{MgSO}_4 + 7\text{H}_2\text{O}$ contain for 100 pts. H_2O at:

0°	10°	20°
26.0	30.9	35.6 pts. MgSO_4 . (Löwel.)

These results may be given in tabular form as follows:

Temp.	A sat. aqueous solution of $\text{MgSO}_4 + 7\text{H}_2\text{O}$ (b) contains for 100 pts. H_2O	
	Anhydrous MgSO_4	$7\text{H}_2\text{O}$ (b) salt
0°	26.0	73.31
10°	30.9	93.75
20°	35.6	116.54

Temp.	A sat. aqueous solution of $\text{MgSO}_4 + 7\text{H}_2\text{O}$ (a) contains for 100 pts. H_2O	
	Anhydrous MgSO_4	$7\text{H}_2\text{O}$ (a) salt
0°	34.67	111.74
10°	38.71	133.67
20°	42.84	159.61

Temp.	A sat aqueous solution of $\text{MgSO}_4 + 6\text{H}_2\text{O}$ contains for 100 pts. H_2O		
	Anhydrous MgSO_4	$6\text{H}_2\text{O}$ salt	$7\text{H}_2\text{O}$ salt
0°	40.75	122.22	146.02
10°	42.32	129.44	155.53
20°	43.87	137.72	167.97

It is seen from table that at the same temp. the $6\text{H}_2\text{O}$ salt is more sol. than the $7\text{H}_2\text{O}$ (a) salt, and the latter is more sol. than $7\text{H}_2\text{O}$ (b) salt; that the solubility of the $7\text{H}_2\text{O}$ (b) salt increases rapidly from 0° to 20° ; that the $6\text{H}_2\text{O}$ salt is not much more sol. at 20° than at 0° , and at 20° the $7\text{H}_2\text{O}$ (b) salt is nearly as sol. as the $6\text{H}_2\text{O}$ salt. (Löwel, A. ch. 13. 42. 405.)

100 pts. H_2O at t° dissolve pts. MgSO_4 . G L = according to Gay-Lussac (A. ch. (2) 11. 311); T = according to Tobler (A. 98. 198).

t°	G L	T	t°	G L	T
0	25.8	24.7	50	49.7	52.5
10	30.5		55		
20	35.0		60	55.9	
25		37.1	70	60.4	
30	39.8		80	65.1	
40	45.2		90	70.3	

100 pts. H_2O at 105.5° dissolve 135.2 pts. MgSO_4 . (Griffiths.)

$\text{MgSO}_4 + \text{Aq}$ sat. at 17.5° has sp. gr. = 1.2932, and contains 55.57 % $\text{MgSO}_4 + 7\text{H}_2\text{O}$, or 100 pts. H_2O dissolve 125.06 pts. $\text{MgSO}_4 + 7\text{H}_2\text{O}$, or 60 pts. MgSO_4 at 17.5° . (Karsten.)

100 pts. H_2O at 0° dissolve 53.8 pts., and 123 pts. at ord. temp. (Otto-Graham.)

Sol. in 2 pts. cold, and 0.5 pt. boiling H_2O . (Fourcroy.)

The aqueous solution contains for 100 pts. H_2O 92.217 pts. $\text{MgSO}_4 + 7\text{H}_2\text{O}$ at 15° . (Michel and Kraft.)

1 pt. $\text{MgSO}_4 + 7\text{H}_2\text{O}$ is sol. in 0.933 pt. H_2O at 15° (Gerlach); in 0.92 pt. H_2O at 23° (Schiff).

100 pts. H_2O dissolve 28.067 pts. MgSO_4 at 17° . (Pfaff, A. 99. 224.)

100 pts. H_2O dissolve pts. MgSO_4 at t° .	
t°	Pts. MgSO_4
0	26.37
17.9	33.28
24.1	35.98

(Diacon, J. B. 1886. 62.)
100 pts. $\text{MgSO}_4 + \text{Aq}$ sat. at $18-20^\circ$ contain 25.67-26.38 pts. MgSO_4 . (v. Hauer, J. pr. 98. 137.)

Solubility in 100 pts. H₂O at t°, using
MgSO₄+7H₂O.

t°	Pts. MgSO ₄	t°	Pts. MgSO ₄	t°	Pts. MgSO ₄
0	26.9	37	44.2	74	61.4
1	27.4	38	44.7	75	61.9
2	27.9	39	45.2	76	62.3
3	28.3	40	45.6	77	62.8
4	28.8	41	46.1	78	63.2
5	29.3	42	46.5	79	63.7
6	29.7	43	47.0	80	64.2
7	30.2	44	47.5	81	64.6
8	30.6	45	48.0	82	65.1
9	31.1	46	48.4	83	65.6
10	31.5	47	48.9	84	66.0
11	32.0	48	49.3	85	66.5
12	32.4	49	49.8	86	67.0
13	32.9	50	50.3	87	67.5
14	33.4	51	50.7	88	68.0
15	33.8	52	51.2	89	68.4
16	34.3	53	51.7	90	68.9
17	34.7	54	52.2	91	69.4
18	35.2	55	52.7	92	69.9
19	35.7	56	53.2	93	70.4
20	36.2	57	53.6	94	70.9
21	36.7	58	54.1	95	71.4
22	37.1	59	54.5	96	71.9
23	37.6	60	55.0	97	72.4
24	38.0	61	55.5	98	72.8
25	38.5	62	55.9	99	73.3
26	39.0	63	56.4	100	73.8
27	39.5	64	56.8	101	74.3
28	39.9	65	57.3	102	74.8
29	40.4	66	57.7	103	75.2
30	40.9	67	58.2	104	75.7
31	41.4	68	58.6	105	76.2
32	41.8	69	59.1	106	76.7
33	42.3	70	59.6	107	77.2
34	42.8	71	60.0	108	77.7
35	43.3	72	60.5	108.4	77.9
36	43.7	73	61.0

(Mulder, calculated from his own and other observations, Scheik. Verhandel. 1864. 52).

100 pts. H₂O dissolve 72.4 pts. MgSO₄+7H₂O at 0°, 178 pts. at 40°; and 212.6 pts. at 49°. (Tilden, Chem. Soc. 45. 409.)

Supersat. MgSO₄+Aq is brought to crystallisation by addition of crystal of MgSO₄+7H₂O, or an isomorphous substance as ZnSO₄+7H₂O, NiSO₄+7H₂O, FeSO₄+7H₂O, or CoSO₄+7H₂O. (Thomson, Chem. Soc. 35. 199.)

Sat. MgSO₄+Aq contains at:

2° 7° 23° 67° 81°
20.9 22.5 26.0 35.6 38.6% MgSO₄,

94° 130° 145° 164° 188°
41.5 45.3 38.0 29.3 20.4% MgSO₄.

Readily forms supersat. solutions.

(Étard, A. ch. 1894, (7) 2. 551.)

M.-pt. of MgSO₄+7H₂O is 70°. (Tilden, Chem. Soc. 45. 409.)

MgSO₄+Aq with sp. gr. 1.50 contains 44.4 % MgSO₄; sp. gr. 1.42, 39%; sp. gr. 1.30, 30% MgSO₄. (Dalton.)

Sp. gr. of MgSO₄+Aq sat. at 15°=1.275 (Michel and Krafft); at 8°=1.267 (Anthon); at 18.75°=1.293 (Karsten.).

Sp. gr. of MgSO₄+Aq at 15°.

% MgSO ₄	Sp. gr.	% MgSO ₄	Sp. gr.
5	1.054	30	1.326
10	1.108	35	1.384
15	1.161	40	1.446
20	1.215	45	1.511
25	1.269	50	1.580

(Calculated from Anthon by Schiff, A. 107. 303.)

Sp. gr. of MgSO₄+Aq at 23°.

% MgSO ₄ +7H ₂ O	Sp. gr.	% MgSO ₄ +7H ₂ O	Sp. gr.
1	1.0048	28	1.1426
2	1.0096	29	1.1481
3	1.0144	30	1.1536
4	1.0193	31	1.1592
5	1.0242	32	1.1648
6	1.0290	33	1.1704
7	1.0339	34	1.1760
8	1.0387	35	1.1817
9	1.0437	36	1.1875
10	1.0487	37	1.1933
11	1.0537	38	1.1991
12	1.0587	39	1.2049
13	1.0637	40	1.2108
14	1.0688	41	1.2168
15	1.0739	42	1.2228
16	1.0790	43	1.2288
17	1.0842	44	1.2349
18	1.0894	45	1.2410
19	1.0945	46	1.2472
20	1.0997	47	1.2534
21	1.1050	48	1.2596
22	1.1103	49	1.2659
23	1.1156	50	1.2722
24	1.2109	51	1.2786
25	1.1261	52	1.2850
26	1.1316	53	1.2915
27	1.1371	54	1.2980

(Schiff, A. 113. 185.)

Sp. gr. of MgSO ₄ +Aq at 12°.			
% MgSO ₄ +7H ₂ O	Sp. gr.	% MgSO ₄ +7H ₂ O	Sp. gr.
1	1.0046	21	1.1071
2	1.0096	22	1.1125
3	1.0146	23	1.1179
4	1.0196	24	1.1234
5	1.0246	25	1.1289
6	1.0296	26	1.1344
7	1.0346	27	1.1399
8	1.0396	28	1.1454
9	1.0446	29	1.1510
10	1.0497	30	1.1566
11	1.0548	31	1.1622
12	1.0599	32	1.1679
13	1.0650	33	1.1736
14	1.0702	24	1.1793
15	1.0754	35	1.1850
16	1.0807	36	1.1908
17	1.0859	37	1.1965
18	1.0911	38	1.2023
19	1.0964	39	1.2082
20	1.1018	40	1.2140

(Oudemans, Z. anal. 7. 419.)

Sp. gr. of MgSO ₄ +Aq at 15°.			
% MgSO ₄	Sp. gr.	% MgSO ₄	Sp. gr.
1	1.01031	14	1.15083
2	1.02062	15	1.16222
3	1.03092	16	1.17420
4	1.04123	17	1.18618
5	1.05154	18	1.19816
6	1.06229	19	1.21014
7	1.07304	20	1.22212
8	1.08379	21	1.23465
9	1.09454	22	1.24718
10	1.10529	23	1.25972
11	1.11668	24	1.27225
12	1.12806	25	1.28478
13	1.13945	25.248	1.28802

(Gerlach, Z. anal. 8. 287.)

Sp. gr. of MgSO₄+Aq at 23.5°. a=no. of 1/2 mols. in grms. dissolved in 1000 g. H₂O; b=sp. gr. if a is MgSO₄+7H₂O, 1/2 mol. wt.=123; c=sp. gr. if a is MgSO₄, 1/2 mol. wt.=60.

a	b	c	a	b	c
1	1.056	1.059	5	1.203	1.260
2	1.103	1.114	6	1.229	...
3	1.141	1.166	7	1.252	...
4	1.174	1.214	8	1.273	...

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of MgSO ₄ +Aq at 15°.			
% MgSO ₄	Sp. gr.	% MgSO ₄	Sp. gr.
5	1.0510	20	1.2200
10	1.1052	25	1.2861
15	1.1602

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of MgSO₄+Aq at 0°. S=pts. MgSO₄ in 100 pts. solution.

S	Sp. gr.	S	Sp. gr.
13.800	1.1586	7.4046	1.0826
11.7458	1.1329	5.0447	1.0557
9.6218	1.1072	2.5907	1.0284

(Charpy, A. ch. (6) 29. 26.)

Sat. MgSO₄+Aq boils at 105° (Griffiths; 108.4° (Mulder).

Crust forms at 103.5° (solution containing 48.4 pts. MgSO₄ to 100 pts. H₂O); highest temp. observed, 105°. (Gerlach, Z. anal. 28. 426.)

B.-pt. of MgSO₄+Aq containing pts. MgSO₄ to 100 pts. H₂O.

B.-pt.	Pts. MgSO ₄	B.-pt.	Pts. MgSO ₄	B.-pt.	Pts. MgSO ₄
100.5°	8.8	102.5°	34.7	104.5°	51.3
101.0	16.7	103.0	39.5	105	54.6
101.5	23.5	103.5	43.8	106	75?
102.0	29.5	104.0	47.7

(Gerlach, Z. anal. 28. 432.)

Sp. gr. of MgSO₄+Aq at 9.5°.

Mass of salt per unit mass of solution	Density of solution (g. per cc.).
0.00191	1.00170
0.00380	1.00346
0.00569	1.00526
0.00758	1.00705
0.01132	1.01060

(McGregor, C. N. 1887, 55.6.)

Sp. gr. of MgSO₄+Aq at 25°.

Concentration of MgSO ₄ +Aq	Sp. gr.
1—normal	1.0584
1/2—"	1.0297
1/3—"	1.0152
1/4—"	1.0076

(Wagner, Z. phys. Ch. 1890, 5. 38.)

Sp. gr. at 16°/4° of MgSO₄+Aq containing 11.0222% MgSO₄=1.11471; containing 8.343% MgSO₄=1.08558. (Schönrock, Z. phys. Ch. 1893, 11. 782.)

Sp. gr. of MgSO ₄ +Aq.		
G.-equivalents MgSO ₄ per liter	t°	Sp. gr. t°/t°
0.002548	17.989	1.0001625
0.005093	18.020	1.000324
0.01015	17.995	1.000639
0.02023	17.980	1.001274
0.05023	18.047	1.003117
0.09950	18.033	1.006122
0.19773	18.014	1.012035
0.29459	17.997	1.017806
0.48671	17.994	1.029101
0.5022	17.90	1.03000
5.0220	17.95	1.26970
0.002616	14.096	1.0001672
0.005230	14.109	1.0003311
0.01042	14.098	1.000659
0.02077	14.092	1.001306
0.12462	14.199	1.007682
0.24567	14.092	1.014980

(Kohlrausch, W. Ann. 1894, 53. 27.)

Sp. gr. of MgSO ₄ +Aq.	
½ MgSO ₄ g. in 1000 g. of solution	Sp. gr. 16°/16°
0	1.000000
0.5368	1.000570
1.0917	1.001157
"	1.001141
2.1076	1.002234
4.1367	1.004372
9.0608	1.009523
18.0846	1.018954
37.1342	1.038983
52.1362	1.054867

(Dijken, Z. phys. Ch. 1897, 24. 108.)

Sp. gr. of MgSO₄+Aq at 18.2°, when p = per cent strength of the solution; d = observed density; and w = volume conc. in grams per cc. ($\frac{pd}{100} = w$.)

p	d	w
26.25	1.2903	1.3374
25.91	1.2860	1.3319
24.53	1.2693	1.3101
21.60	1.2330	1.2650
18.41	1.1950	1.2187
13.79	1.1423	1.1562
12.63	1.1291	1.1413
11.29	1.1147	1.1246
8.08	1.0803	1.0859
2.01	1.0204	1.0191

(Barnes, J. phys. Chem. 1898, 2. 542.)

Sp. gr. of MgSO ₄ +Aq at 20°.		
Normality of MgSO ₄ +Aq	% MgSO ₄	Sp. gr.
2.73	25.46	1.2879
1.86	18.61	1.2019
0.934	10.14	1.1049

(Forchheimer, Z. phys. Ch. 1900, 34. 24.)

Sp. gr. of dil. MgSO₄+Aq at 20.004°.
Conc. = g. equiv. per l. at 20.004°.
Sp. gr. compared with H₂O at 20.004° = 1.

Conc.	Sp. gr.
0.0000	1.000,000,0
0.0001	1.000,006,4
0.0002	1.000,012,9
0.0003	1.000,019,4
0.0004	1.000,025,9
0.0005	1.000,032,4
0.0010	1.000,064,8
0.0020	1.000,129,4
0.0050	1.000,322,4
0.0100	1.000,642,1

(Lamb and Lee, J. Am. Chem. Soc. 1913, 35. 1684.)

More sol. in HCl+Aq than in H₂O. (Rich-ter.)

In sat. HCl+Aq, anhydrous MgSO₄ is scarcely sol.; MgSO₄+7H₂O dissolves, but is precipitated by a current of HCl gas. (Hens-gen, B. 10. 259.)

Margueritte (C. R. 43. 50) denies the pre-cipitation.

For solubility in H₂SO₄, see MgH₂(SO₄)₂.

Completely pptd. from MgSO₄+Aq by conc. HC₂H₃O₂+Aq. (Persoz.)

Somewhat sol. in sat. NH₄Cl+Aq with separation of a double sulphate.

Rapidly sol. in KCl+Aq with separation of K₂SO₄.

Sol. in sat. NaCl+Aq without pptn. of the latter.

Easily sol. in sat. KNO₃+Aq without caus-ing any pptn.

Sol. in sat. NaNO₃+Aq. (Karsten.)

Rapidly sol. in sat. CuSO₄+Aq; when saturation is reached, a double salt separates out. (Karsten.)

100 pts. sat. MgSO₄+NiSO₄+Aq at 18-20° contain 30.93 pts. of the two salts; 100 pts. sat. MgSO₄+ZnSO₄+Aq at 18-20° contain 35.45 pts.; 100 pts. sat. MgSO₄+NiSO₄+ZnSO₄+Aq at 18-20° contain 35.62 pts. (v. Hauer, J. pr. 98. 137.)

100 pts. H₂O dissolve 14.1 pts. MgSO₄ and 9.8 pts. K₂SO₄, if sat. MgSO₄+Aq is sat. with K₂SO₄; 32.4 pts. MgSO₄ and 8.2 pts. K₂SO₄, if sat. K₂SO₄+Aq is sat. with MgSO₄, all at 15°. (Mulder, J. B. 1866.)

100 pts. H₂O dissolve 25.95 pts. MgSO₄ and 5.21 pts. Na₂SO₄ at 0°. (Diacon, J. B. 1888. 62.)

100 pts. H₂O dissolve 15.306 pts. MgSO₄ and 13.086 pts. Na₂SO₄ at 0°. (Pfaff, A. 99. 224.)
See also under MgNa₂(SO₄)₂+4H₂O.

Solubility of mixtures of MgSO₄ and MgNa₂(SO₄)₂+4H₂O at t°.

t°	g. per 100 g. H ₂ O	
	Na ₂ SO ₄	MgSO ₄
22	23.3	31.4
24.5	27.2	24.2
30	36.1	19.1
35	33.9	18.44

(Roozeboom, 1888, Z. phys. Ch. 2. 518.)
See also under MgNa₂(SO₄)₂+4H₂O.

Slowly sol. in sat. ZnSO₄+Aq without pptn. until saturation, when a double salt separates out.

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 828.)

100 pts. dil. alcohol containing at 15°:
10 20 40 % alcohol
contain 39.3 21.3 1.62% MgSO₄+7H₂O.
(Schiff, A. 118. 365.)

At higher temp. the solubility increases proportional to the temp. (Gerardin, A. ch. (4) 5. 145.)

100 pts. absolute methyl alcohol dissolve 1.18 pts. MgSO₄ at 18°. (de Bruyn, Z. phys. Ch. 10. 783.)

100 pts. absolute methyl alcohol dissolve 41 pts. MgSO₄+7H₂O at 17°; 100 pts. absolute methyl alcohol dissolve 29 pts. MgSO₄+7H₂O at 3-4°; 100 pts. 93% methyl alcohol dissolve 9.7 pts. MgSO₄+7H₂O at 17°; 100 pts. 50% methyl alcohol dissolve 4.1 pts. MgSO₄+7H₂O at 3-4°. (de Bruyn, R. t. c. 11. 112.)

100 pts. absolute ethyl alcohol dissolve 1.3 pts. MgSO₄+7H₂O at 3°. (de Bruyn.)

Insol. in CS₂. (Arctowski, Z. anorg. 1894, 6. 257.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790.); ethyl acetate (Naumann, B. 1910, 43. 314.); acetone. (Naumann, B. 1904, 37. 4329.)

100 g. 95% formic acid dissolve 0.34 g. MgSO₄ at 19°. (Aschan, Ch. Ztg. 1913, 37. 1117.)

100 g. sat. solution of MgSO₄ and sugar in H₂O contains 46.52 g. sugar +14.0 g. MgSO₄, or 100 g. H₂O dissolve 119.6 g. sugar +36.0 g. MgSO₄. (Köhler, Z. Ver. Zucker-ind, 1897, 47. 447.)

Magnesium hydrogen sulphate, MgH₂(SO₄)₂.
Decomp. by H₂O. Sol. in H₂SO₄. Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

MgH₂(SO₄)₂. Boiling H₂SO₄ dissolves about 2% MgSO₄, from which this compound crystallises. (Schultz, Pogg. 133. 137.)

Magnesium pyrosulphate, MgS₂O₇.
Decomp. by H₂O.

Magnesium manganous sulphate, MgSO₄·2MnSO₄+15H₂O.
Min. *Fauserite*.

Magnesium manganous zinc sulphate, MgSO₄·MnSO₄·ZnSO₄+21H₂O.
Sol. in H₂O. (Vohl, A. 99. 124.)

Magnesium nickel sulphate, MgSO₄·3NiSO₄+28H₂O.
Sol. in H₂O. (Schiff.)

Magnesium nickel potassium sulphate, MgSO₄·NiSO₄·2K₂SO₄+12H₂O.
Sol. in H₂O. (Vohl, A. 94. 57.)

Magnesium potassium sulphate, MgK₂(SO₄)₂+6H₂O.
100 pts. H₂O dissolve 22.7 pts. anhydrous salt at 16.5°. (Mulder.)
100 pts. H₂O dissolve at:
0° 10° 20° 30° 35°
14.1 19.6 25.0 30.4 33.3 pts. anhy-
drous salt,
45° 55° 60° 65° 75°
40.5 47.0 50.2 53.0 59.8 pts. anhy-
drous salt.
(Tobler, A. 96. 193.)

100 g. H₂O dissolve 30.52 g. MgK₂(SO₄)₂+6H₂O at 15°. (Lothian, Pharm. J. 1909, 82. 292.)

Solubility in H₂O at t°.

t°	Sat. solution contains		Mols. K ₂ SO ₄ : mols. MgSO ₄ in the solution	100 pts. H ₂ O dissolve	
	% K ₂ SO ₄	% MgSO ₄		K ₂ SO ₄	MgSO ₄
10	9.4	9.8	1:1.52	11.63	12.13
20	10.9	10.8	1:1.43	13.92	13.79
30	12.4	11.8	1:1.38	16.36	15.56
40	13.8	13.1	1:1.37	18.88	17.92
50	14.7	14.8	1:1.46	20.85	20.99
60	15.2	16.3	1:1.55	22.19	23.79
70	15.6	16.8	1:1.52	23.07	24.85
80	16.0	17.1	1:1.56	23.91	25.56
80	16.6	18.1	1:1.58	25.42	27.72
90	17.2	18.2	1:1.54	26.62	28.17

(Precht, B. 1882, 15. 1668.)

Sp. gr. of aqueous solution at 15° containing:

2 4 6 8% hydrous salt,
1.0129 1.0261 1.0394 1.053

10 12 14 16% hydrous salt,
1.0668 1.0808 1.095 1.1094

18 20 22% hydrous salt.
1.124 1.1388 1.1539
(Schiff, A. 113. 183, calculated by Gerlach,
Z. anal. 8. 287.)

Sp. gr. of $\text{MgK}_2(\text{SO}_4)_2 + \text{Aq}$ at 18°.

G-equiv. of salt per l.	Sp. gr.
1.0010	1.0633
0.8345	1.0531
0.6688	1.0427
0.3744	1.0243
0.0998	1.0040
0.02004	1.0015
0.01004	1.0004

These results lead the author to conclude that in dil. solutions the double salt is decomp. into its constituents. (McKay, Elektrochem. Zeit. 1899, 6. 115.)

Min. *Picromerite*.

+4H₂O. (van der Heide, B. 26. 414.)

2MgSO₄, K₂SO₄. Min. *Langbeinite*.

Deliquescent. Absorbs 56.26% H₂O from air to form K₂SO₄, MgSO₄+6H₂O. (Mallet, Chem. Soc. 1900, 77. 220.)

4MgSO₄, K₂SO₄+5H₂O. (van't Hoff and Kassatkin, B. A. B. 1889. 951.)

Magnesium potassium zinc sulphate, MgSO₄, 2K₂SO₄, ZnSO₄+12H₂O.

Sol. in H₂O. (Vohl, A. 94. 57.)

Magnesium potassium sulphate chloride, MgSO₄, K₂SO₄, MgCl₂+6H₂O.

Min. *Kainite*.

Magnesium rubidium sulphate, MgSO₄, Rb₂SO₄+6H₂O.

Sol. in H₂O. (Tutton, Chem. Soc. 63. 337.)

1 l. H₂O dissolves 202 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

2MgSO₄, Rb₂SO₄. Deliquescent. (Mallet, Chem. Soc. 1900, 77. 223.)

Magnesium sodium sulphate, MgSO₄, Na₂SO₄+4H₂O.

Min. *Blödite*, *Simonyite*.

Blödite is efflorescent; Simonyite, deliquescent.

+5H₂O. Min. *Löwite*.

+6H₂O. Decomp. on air. Sol. in 3 pts. cold H₂O.

Na₄Mg(SO₄)₄. Min. *Vanthoffite*. (van't Hoff, B. A. B. 1902. 414.)

MgNa₂(SO₄)₂+4H₂O. Min. *Astrakanite*.

100 mols. H₂O hold mols. salt in solution at t°.

t°	MgSO ₄	Na ₂ SO ₄
22	4.70	2.95
24.5	3.68	3.45
30	3.60	3.60
35	3.69	3.69
47	3.60	3.60

(Roozeboom, R. t. c. 1887, 6. 333.)

Solubility of mixtures of MgNa₂(SO₄)₂+4H₂O and Na₂SO₄ at t°.

t°	g. per 100 g. H ₂ O	
	Na ₂ SO ₄	MgSO ₄
18.5	43.0	45.5
22	35.2	48.9
24.5	32.5	50.3
30	25.9	55.0
35	23.5	59.4

(Roozeboom, Z. phys. Ch. 1888, 2. 518.)

See also under MgSO₄.

Magnesium thallous sulphate, MgSO₄, Tl₂SO₄+6H₂O.

Sol. in H₂O, but decomp. by repeated recrystallisations. (Werther.)

Magnesium uranyl sulphate, MgSO₄, (UO₂)SO₄+5H₂O.

(de Coninck, Chem. Soc. 1905, 88. (2) 530.)

Magnesium zinc sulphate, MgSO₄, ZnSO₄+14H₂O.

Sol. in H₂O. (Pierre, A. ch. (3) 16. 244.)

+10H₂O. (Pierre.)

3ZnSO₄, 5MgSO₄+56H₂O. (Schiff.)

There are only two compounds, 2(MgSO₄, 7H₂O), ZnSO₄, 7H₂O and MgSO₄, 7H₂O, ZnSO₄, 7H₂O. (Hollmann, Z. phys. Ch. 1901, 37. 212, and 1902, 40. 577.)

Magnesium sulphate potassium chloride, MgSO₄, KCl+3H₂O or MgSO₄, K₂SO₄, MgCl₂+6H₂O.

Min. *Kainite*.

100 pts. H₂O dissolve 79.56 pts. at 18°. (Krause, Arch. Pharm. (3) 6. 326.)

Not sol. in a mixture of abs. alcohol and ether, which dissolves out MgCl₂. (Lehmann, J. B. 1867. 416.)

Alcohol dissolves out MgCl₂, also little H₂O. Much H₂O dissolves completely. (Zincken, Miner. Jahrb. 1865. 310.)

Magnesium sulphate potassium chromate, 2MgSO₄, K₂CrO₄+9H₂O.

Sol. in H₂O. (Étard, C. R. 85. 443.)

Manganous sulphate, basic, 3MnO , 2SO_3 , + $3\text{H}_2\text{O}$.

Insol. in H_2O , but slowly decomp. thereby. (Gorgeu, C. R. 94. 1425.)

Manganous sulphate, MnSO_4 .

Anhydrous.

Absorbs H_2O from the air to form $\text{MnSO}_4 \cdot 4\text{H}_2\text{O}$.
1 pt. MnSO_4 is sol. in pts. H_2O at t° .

t°	Pts. H_2O	t°	Pts. H_2O	t°	Pts. H_2O
6.25	1.77	18.75	1.667	75	1.494
10	1.631	37.5	1.457	101.25	2.031

Or—

100 pts. H_2O dissolve pts. MnSO_4 at t° .

t°	Pts. MnSO_4	t°	Pts. MnSO_4	t°	Pts. MnSO_4
6.25	56.49	18.75	60.00	75	66.95
10	61.29	37.5	68.63	101.25	49.33

(Brandes, Pogg. 20. 575.)

Sol. in 2.5 pts. H_2O at 18.75° ; at 62.5° it is difficult to dissolve 1 pt. MnSO_4 in 3 pts. H_2O , but the sat. solution at 62.5° does not become cloudy on heating to 100° . (Jahn.)

100 pts. MnSO_4 + Aq sat. at 11 – 14° contain 37.5 pts. MnSO_4 . (v. Hauer, J. pr. 103. 114.)

Sat. MnSO_4 + Aq contains at:

-8°	-5°	$+5^\circ$	18°	22°	
30.0	31.0	34.1	38.3	38.2% MnSO_4	
23°	32°	45°	52°	70°	
39.1	41.7	44.2	36.4	41.1% MnSO_4	
83°	110°	115°	123°	130°	140°
36.3	18.4	21.5	16.7	13.6	9.4% MnSO_4

(Étard, A. ch. 1894, (7) 2. 553.)

Solubility in H_2O increases from 0 – 55° , and decreases from 55 – 145° . The increasing solubility is that of $\text{MnSO}_4 \cdot 5\text{H}_2\text{O}$, and MnSO_4 + $2\text{H}_2\text{O}$ separates out at 35° , and is completely insol. at 145° . (Étard.)

If solubility S = pts. anhydrous MnSO_4 in 100 pts. solution, $S = 30.0 + 0.2828t$ from -8° to 57° ; $S = 48.0 - 0.4585t$ from 57° to 150° .

Practically insol. in H_2O at 180° . (Étard, C. R. 106. 208.)

Solubility varies according to the hydrate used. Above results of Étard show the solubility of $\text{MnSO}_4 \cdot 7\text{H}_2\text{O}$ at 0° , and MnSO_4 + $3\text{H}_2\text{O}$ at 57° . Anhydrous MnSO_4 is stable only above 117° . (Linebarger.)

100 pts. H_2O dissolve pts. anhydrous MnSO_4 at t° .

t°	Pts. MnSO_4	t°	Pts. MnSO_4	t°	Pts. MnSO_4
120	67.18	141	41.18	155	26.49
132	63.16	146	38.83	170	16.15

(Linebarger, Am. Ch. J. 15. 225.)

+ H_2O . Stable only between 57° and 117° .

100 pts. H_2O dissolve pts. MnSO_4 from MnSO_4 + H_2O at t° .

t°	Pts. MnSO_4	t°	Pts. MnSO_4	t°	Pts. MnSO_4
48	87.98	78	79.13	115	69.78
53	86.10	90	75.63	117	68.81
65	84.33	100	71.27
72	82.73	106	70.14

(Linebarger.)

Min. Szmikite.

Solubility of MnSO_4 + H_2O in H_2O at t° .

t°	Pts. MnSO_4 per 100 pts. H_2O	t°	Pts. MnSO_4 per 100 pts. H_2O
41.5	61.06	75	49.45
50.1	58.01	84.8	44.87
67.1	51.37	95	38.71
...	...	99.6	34.27

Av. of varying results.

(Cottrell, J. phys. Ch. 1900, 4. 652.)

Linebarger's determinations are inaccurate (Cottrell.)

+ $2\text{H}_2\text{O}$. Stable between 40° and 57° .

100 pts. H_2O dissolve pts. MnSO_4 from MnSO_4 + $2\text{H}_2\text{O}$ at t° .

t°	Pts. MnSO_4	t°	Pts. MnSO_4	t°	Pts. MnSO_4
35	68.88	42	77.63	50	83.16
40	75.31	45	80.07	55	86.27

(Linebarger.)

+ $3\text{H}_2\text{O}$. Stable between 30° and 40° .

100 pts. H_2O dissolve pts. MnSO_4 from MnSO_4 + $3\text{H}_2\text{O}$ at t° .

t°	Pts. MnSO_4	t°	Pts. MnSO_4	t°	Pts. MnSO_4
5	54.68	25	66.85	68	71.89
12	60.56	30	67.38	53	72.81
16	63.41	35	68.31	57	73.17
19	65.12	40	70.63

(Linebarger.)

+ $4\text{H}_2\text{O}$. Sl. efflorescent. Less sol. in boiling than in cold H_2O .

100 pts. H_2O at 4.4° dissolve 31 pts. MnSO_4 + $4\text{H}_2\text{O}$. (Jahn.)

100 pts. H₂O at t° dissolve pts. MnSO₄ + 4H₂O.

t°	Pts. MnSO ₄ + 4H ₂ O	t°	Pts. MnSO ₄ + 4H ₂ O
6 25	113 22	37 5	149
10	123	75	144
18 75	122	101 25	93

(Brandes, Pogg. 20. 575.)

Solubility of MnSO₄ in 100 pts. H₂O at t°, using MnSO₄ + 4H₂O.

t°	Pts. MnSO ₄	t°	Pts. MnSO ₄	t°	Pts. MnSO ₄
0	55.4	35	71.9	70	61.5
1	55.9	36	72.2	71	61.5
2	56.5	37	72.4	72	61.5
3	57.1	38	72.7	73	61.5
4	57.7	39	72.9	74	61.5
5	58.2	40	73.1	75	61.5
6	58.8	41	73.3	76	61.5
7	59.4	42	73.5	77	61.5
8	60.0	43	73.7	78	61.5
9	60.5	44	73.9	79	61.5
10	61.1	45	74.0	80	61.5
11	61.7	46	74.2	81	61.5
12	62.2	47	74.4	82	61.5
13	62.7	48	74.6	83	61.5
14	63.3	49	74.7	84	61.4
15	63.8	50	74.8	85	61.3
16	64.3	51	74.9	86	61.2
17	64.8	52	75.1	87	61.0
18	65.3	53	75.2	88	60.8
19	65.8	54	75.3	89	60.6
20	66.3	55	74.7	90	60.3
21	66.7	56	74.0	91	60.0
22	67.2	57	72.9	92	59.6
23	67.6	58	71.5	93	59.2
24	68.1	59	69.5	94	58.6
25	68.5	60	65.9	95	57.9
26	68.9			96	57.2
27	69.3	63 5	61.3	97	56.3
28	69.7	64	61.5		55.4
29	70.0	65	61.5	99	54.3
30	70.4	66	61.5	100	52.9
31	70.7	67	61.5	101	51.2
32	71.0	68	61.5	102	49.3
33	71.3	69	61.5	102 5	47.4
34	71.6				

(Mulder, Scheik. Verhandel. 1864. 137.)

100 pts. H₂O dissolve pts. MnSO₄ from MnSO₄ + 4H₂O at t°

t°	Pts. MnSO ₄	t°	Pts. MnSO ₄	t°	Pts. MnSO ₄
2 2	57 88	25	72 23	48	84 33
7 3	61 78	30	74 67	52	86 16
11	64 01	35 5	78 41	56	88 19
15	67 12		79 63		
20	69 93	45	83 00		

(Lanebarger)

Stable in aqueous solution between 25° and 31°. (Schieber, M. 1898, 19. 281.)

Solubility of MnSO₄ + 4H₂O in H₂O at t°.

t°	Pts. MnSO ₄ per 100 pts. H ₂ O	t°	Pts. MnSO ₄ per 100 pts. H ₂ O
16.0	63.97	35 0	67.87
17 7	64.16	35 5	68.09
18 5	64.19	39 9	68.81
25.0	65.32	49 9	72.48
30 0	66.43	50 0	72.62
32.2	66.83		

(Cottrell, J. phys. Ch. 1900, 4. 651.)

Lanebarger's determinations are inaccurate. (Cottrell.)

Solubility in H₂O at t°.

t°	g. MnSO ₄ for 100 g. H ₂ O
30.15	66.38
35	68.22

(Richards and Fraprie, Am. Ch. J. 1901, 26. 77.)

+5H₂O. Sol. in 1 pt H₂O at 18 75° (Jahn, A 28. 110.)
Stable from 8° to 18°.100 pts. H₂O dissolve pts. MnSO₄ from MnSO₄ + 5H₂O at t°.

t°	Pts. MnSO ₄	t°	Pts. MnSO ₄	t°	Pts. MnSO ₄
0	58.05	20	75.16	40	84.63
2 5	62.41	25	78.63	42	85.27
4	64.22	30	79.16	45	86.16
7	66.83	32	80.38	47.7	86.95
10	68.05	34	82.04	53	88.89
15	72.33	37	83.91	54	89.08

(Lanebarger.)

Stable in aqueous solution between 15° and 20°. (Schieber, M. 1898, 19. 281.)

Solubility of MnSO₄ + 5H₂O at t°.

t°	Pts. MnSO ₄ per 100 pts. H ₂ O	t°	Pts. MnSO ₄ per 100 pts. H ₂ O
5	58.06	16	61.59
9	59.23	25	64.78
12	60.19	30	67.76
12 3	60.16	31.1	67.92
15	61.08	35.5	71.61

(Cottrell, J. phys. Ch. 1900, 4. 651.)

Lanebarger's determinations are inaccurate. (Cottrell.)

Solubility in H₂O at 25° = 65.09 g. MnSO₄ for 100 g. H₂O. (Richards and Fraprie, Am. Ch. J. 1901, 28. 77.)

+6H₂O. Stable from - 5° to +8°. 100 pts. H₂O dissolve pts. MnSO₄ from MnSO₄+6H₂O at t°.

t°	Pts. MnSO ₄	t°	Pts. MnSO ₄	t°	Pts. MnSO ₄
- 4	55.87	9	70.88	30	76.24
0	64.21	15	72.45	34	77.02
3	66.87	20	74.35	35	77.23
5	67.49	25	75.38	38	77.481

(Linebarger.)

+7H₂O. Efflorescent. Sol. in less than 0.5 pt. H₂O at 18.75°. (Jahn.) Stable between - 10° and - 5°.

100 pts. H₂O dissolve pts. MnSO₄ from MnSO₄+7H₂O at t°.

t°	Pts. MnSO ₄	t°	Pts. MnSO ₄	t°	Pts. MnSO ₄
- 10	50.11	0	53.61	10	59.91
- 8	50.93	5	54.83	15	64.34
- 5	51.53	7	56.62

(Linebarger.)

Stable in aqueous solution below 0°. (Schieber, M. 1898, 19. 281.)

Solubility of MnSO₄+7H₂O in H₂O at t°.

t°	Pts. MnSO ₄ per 100 pts. H ₂ O	t°	Pts. MnSO ₄ per 100 pts. H ₂ O
- 10	47.96	9	59.33
0	56.23	12	61.78
5	56.38	14.3	63.93

(Cottrell, l. c.)

M.-pt. of MnSO₄+7H₂O is 54°. (Tilden, Chem. Soc. 45. 409.)

Sp. gr. of MnSO₄+Aq at 15°.

% MnSO ₄ +4H ₂ O	Sp. gr.	% MnSO ₄ +4H ₂ O	Sp. gr.
1	1.006	29	1.206
2	1.013	30	1.2150
3	1.020	31	1.224
4	1.025	32	1.231
5	1.0320	33	1.244
6	1.038	34	1.250
7	1.044	35	1.2579
8	1.050	36	1.268
9	1.056	37	1.276
10	1.0650	38	1.285
11	1.072	39	1.296
12	1.079	40	1.3038
13	1.085	41	1.313
14	1.093	42	1.322
15	1.1001	43	1.331
16	1.106	44	1.340
17	1.114	45	1.3495
18	1.121	46	1.360
19	1.129	47	1.370
20	1.1363	48	1.380
21	1.144	49	1.389
22	1.150	50	1.3986
23	1.160	51	1.410
24	1.166	52	1.420
25	1.1751	53	1.430
26	1.183	54	1.440
27	1.190	55	1.4514
28	1.200

(Gerlach, Z. anal. 8. 288.)

Sp. gr. of MnSO₄+Aq at 23°. a=no. of ½ mols. in grms. dissolved in 1000 g. H₂O; b=sp. gr. if a is MnSO₄+5H₂O, ½ mol. wt.=120.5; c=sp. gr. if a is MnSO₄ ½ mol. wt.=75.5.

a	b	c	a	b	c
1	1.068	1.071	6	1.306	3.576
2	1.128	1.139	7	1.341	1.429
3	1.181	1.202	8	1.371	...
4	1.227	1.262	9	1.399	..
5	1.269	1.320	10	1.428	..

(Favre and Valson, C. R. 79. 968.)

Above table recalculated by Gerlach (Z. anal 28. 475.)

% MnSO ₄ +5H ₂ O	Sp. gr.	% MnSO ₄ +5H ₂ O	Sp. gr.
10	1.0630	40	1.2900
20	1.1325	50	1.3800
30	1.2070

Sp. gr. of $\text{MnSO}_4 + \text{Aq}$ at 15° . $a = \%$; $b =$ sp. gr. if a is MnSO_4 ; $c =$ sp. gr. if a is $\text{MnSO}_4 + 4\text{H}_2\text{O}$; $d =$ sp. gr. if a is $\text{MnSO}_4 + 5\text{H}_2\text{O}$; $e =$ sp. gr. if a is $\text{MnSO}_4 + 7\text{H}_2\text{O}$.

a	b	c	d	e
5	1.0500	1.0340	1.0310	1.0270
10	1.1035	1.0690	1.0630	1.0545
15	1.1605	1.1055	1.0965	1.0830
20	1.2215	1.1435	1.1315	1.1130
25	1.2870	1.1835	1.1685	1.1440
30	1.3575	1.2255	1.2070	1.1765
35	...	1.2695	1.2470	1.2105
40	...	1.3155	1.2885	1.2455
45	...	1.3640	1.3315	1.2815
50	1.3760	1.3185
55	1.3565

(Gerlach, Z. anal. 28. 475.)

Sp. gr. of $\text{MnSO}_4 + \text{Aq}$ at 0° . $S =$ pts. MnSO_4 in 100 pts. solution.

S	Sp. gr.	S	Sp. gr.
16.7450	1.1834	8.8295	1.0928
14.0462	1.1519	6.0172	1.0622
11.5804	1.1239	3.0865	1.0315

(Charpy, A. ch. (6) 29. 26.)

Sp. gr. of $\text{MnSO}_4 + \text{Aq}$ at room temp. containing:

11.45 18.8 22.08 % MnSO_4 .
1.1469 1.2513 1.3082

(Wagner, W. Ann. 1883, 18. 271.)

Sp. gr. of $\text{MnSO}_4 + \text{Aq}$ at 25° .

Concentration of $\text{MnSO}_4 + \text{Aq}$	Sp. gr.
1—normal	1.0728
$\frac{1}{2}$ —"	1.0365
$\frac{1}{4}$ —"	1.0179
$\frac{1}{8}$ —"	1.0087
$\frac{1}{16}$ —"	1.0041

(Wagner, Z. phys. Ch. 1890, 5. 39.)

Sp. gr. at $16^\circ/4^\circ$ of $\text{MnSO}_4 + \text{Aq}$ containing 30.819% $\text{MnSO}_4 = 1.36267$. (Schönrock, Z. phys. Ch. 1893, 11, 781.)

Sat. $\text{MnSO}_4 + \text{Aq}$ boils at 102.4° ; crust forms at 101.6° , and solution contains 48.7 pts. MnSO_4 to 100 pts. H_2O .

B.-pt. of $\text{MnSO}_4 + \text{Aq}$ containing pts. MnSO_4 to 100 pts. H_2O .

B.-pt.	Pts. MnSO_4	B.-pt.	Pts. MnSO_4
100.5°	17.1	102.0°	58.9
101.0	32.1	102.4	68.4
101.5	46.2

(Gerlach, Z. anal. 26. 434.)

Sol. in about 20 pts. boiling H_2SO_4 , and more sol. in boiling $\text{H}_2\text{SO}_4 + \text{Aq}$ of 1.6 sp. gr. (Schultz, Pogg. 183. 137.)

Completely pptd. from solution by $\text{HC}_2\text{H}_3\text{O}_2$. (Persoz.)

For solubility in $(\text{NH}_4)_2\text{SO}_4$, see under $(\text{NH}_4)_2\text{SO}_4$.

$\text{MnSO}_4 + \text{Aq}$ sat. at 10° , then sat. with K_2SO_4 at same temp. contains for 100 pts. H_2O 16.7 pts. MnSO_4 and 44.3 pts. K_2SO_4 (Mulder.)

Solubility of $\text{MnSO}_4 + \text{Na}_2\text{SO}_4$ in H_2O at 35° .

g. per 100 g. sat. solution		
MnSO_4	Na_2SO_4	Solid phase
39.45	0	$\text{MnSO}_4, \text{H}_2\text{O}$
33.92	5.23	"
33.06	7.97	$\text{MnSO}_4, \text{H}_2\text{O} + 9\text{MnSO}_4, 10\text{Na}_2\text{SO}_4$
32.92	7.42	"
31.05	9.20	$9\text{MnSO}_4, 10\text{Na}_2\text{SO}_4$
27.67	10.76	"
22.14	14.28	"
14.58	20.01	"
13.96	21.91	"
12.19	22.49	$9\text{MnSO}_4, 10\text{Na}_2\text{SO}_4 + \text{MnSO}_4, 3\text{Na}_2\text{SO}_4$
10.45	23.41	$\text{MnSO}_4, 3\text{Na}_2\text{SO}_4$
7.43	26.58	"
5.69	29.31	"
5.11	30.52	$\text{MnSO}_4, 3\text{Na}_2\text{SO}_4 + \text{Na}_2\text{SO}_4$
2.96	31.33	Na_2SO_4
0	33	"

(Schreinemakers and Provije, Proc. Ak. Wet. Amsterdam, 1913, 15. 326.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Anhydrous MnSO_4 in insol. in absolute alcohol.

1000 pts. alcohol of 0.872 sp. gr. dissolve 6.3 pts. MnSO_4 .

Sol. in 50 pts. of 50% alcohol. Insol. in absolute alcohol. (Brandes, Pogg. 20. 556.)

100 pts. solution saturated at 15° in dil. alcohol containing:

0 10 50 60 % alcohol, contain
56.25 51.4 2.0 0.66 pts. $\text{MnSO}_4 + 5\text{H}_2\text{O}$.

(Schiff, A. 118. 365.)

When $\text{MnSO}_4 + 7\text{H}_2\text{O}$ is boiled with absolute alcohol none is dissolved, but $\text{MnSO}_4 + 3\text{H}_2\text{O}$ is formed.

When $\text{MnSO}_4 + 7\text{H}_2\text{O}$ is dissolved in 15–50% alcohol, the liquid separates into two layers, the lower containing less (12–14%) alcohol and more (47–49%) salt; the upper containing

more (50-55%) alcohol and less (1.3-2.2%) salt. If the alcohol has the above strength (15-50%) the separation takes place at ordinary temp., but with 13-14% or 60% or more alcohol, warming is necessary to effect the separation. (Schiff, A. 118, 363.)

Solubility of $\text{MnSO}_4 + \text{H}_2\text{O}$ in alcohol + Aq at t° .

Composition of two layers sat. with the solid salt at t° .

t°	Alcohol layer		Water layer	
	% alcohol	% MnSO_4	% alcohol	% MnSO_4
30	45.20	2.49	8.69	30.15
31	43.90	2.74	8.47	30.10
■	41.71	3.44	9.24	28.61
37	38.26	4.84	11.03	26.47
41	34.01	5.86	11.93	24.97
■	32.37	6.89	13.57	23.09
43	31.42	8.51	14.33	22.01

(Schreinemakers and Deuss, Z. phys. Ch. 1912, 79, 559.)

Composition of alcohol solutions sat. with $\text{MnSO}_4 + \text{H}_2\text{O}$ at t° .

t°	% H_2O	% alcohol	% MnSO_4
50	63.74	0	36.26
	65.21	6.67	28.12
	65.23	16.02	18.75
	64.83	22.63	12.54
	59.41	36.47	4.12
35	61.4	0	38.6
	62.13	5.50	32.37
	62.06	6.46	31.48
	62.01	7.48	30.51
	*62.15	9.24	28.61
	*54.85	41.71	3.44
	50.69	47.73	1.58
	50.16	48.27	1.57
30	61.4	0	38.6
	61.43	2.26	36.31
	61.25	5.09	33.66
	60.78	5.96	*33.26
	*61.16	8.69	30.15
	*52.31	45.20	2.49
	44.83	54.19	0.98
	30.95	68.97	0.08
	0.19	90.80	0.01

*The solutions also sat. with respect to one another.

(Schreinemakers and Deuss.)

Composition of the solutions sat. with respect to one another.

t°	water layer		alcohol layer	
	% alcohol	% MnSO_4	% alcohol	% MnSO_4
50	†5.68	34.95	†53.64	0.97
	†7.69	30.99	†45.83	2.19
	†8.70	29.20	†41.93	3.11
	†11.85	24.84	†35.15	5.95
35	†8.38	29.52	†42.38	3.07
	*9.24	28.61	*41.71	3.44
	10.75	26.33	36.89	5.19
	15.09	21.85	■	9.03
30	†7.60	32.40	†50.97	1.74
	*8.69	30.15	*45.20	2.49
	10.46	27.58	40.71	3.93
	11.86	25.75	37.54	5.20
	16.18	20.86	29.89	■

(Schreinemakers and Deuss.)

†Metastable solutions.

*Solutions also sat. with respect to $\text{MnSO}_4 + \text{H}_2\text{O}$.

Solubility of $\text{MnSO}_4 + 4\text{H}_2\text{O}$ in alcohol + Aq.

g. H_2O	g. alcohol	$\text{MnSO}_4 \cdot 4\text{H}_2\text{O}$
55.86	30.03	14.11
52.25	43.59	4.16
49.41	47.66	2.94
45.34	53.00	■
42.56	56.24	1.20

(Linebarger, Am. Ch. J. 1892, 14, 380.)

Solubility of $\text{MnSO}_4 + 5\text{H}_2\text{O}$ in alcohol + Aq.

Composition of the solution sat. with $\text{MnSO}_4 + 5\text{H}_2\text{O}$.

t°	alcohol layer		water layer	
	% alcohol	% MnSO_4	% alcohol	% MnSO_4
10	37.06	5.44	13.78	25.25
15	44.56	2.79	9.25	29.79
17.6	47.11	2.22	8.53	30.86
21	53.55	1.10	6.10	35.06
25	53.09	1.23	6.81	33.77

Composition of solution sat. with solid substance at 25° .

% H_2O	% alcohol	% MnSO_4	Solid phase
60.7	■	39.3	$\text{MnSO}_4 + 5\text{H}_2\text{O}$
*59.47	6.81	33.72	"
*45.68	53.09	1.23	"
42.05	57.39	0.56	"
23.30	76.70	0.0	$\text{MnSO}_4 + \text{H}_2\text{O}$

*The two liquids are sat. with respect to one another.

Position of the two solutions sat. with to one another at 25°.

Water layer		Alcohol layer	
sol.	% MnSO_4	% alcohol	% MnSO_4
1	*33.72	*53.09	1.23
2	31.51	49.76	1.83
3	22.61	32.75	8.01

sat. with $\text{MnSO}_4 \cdot 5\text{H}_2\text{O}$.

(Schreinemaker and Deuss.)

in absolute ether between 5° and 7°, crystal H_2O is removed thereby, boiling oil of turpentine, but 1 mol. H_2O is removed from $\text{MnSO}_4 + 4\text{H}_2\text{O}$ (Pogg. 20. 568)

in benzonitrile. (Naumann, B. 1914, 0.); ethyl acetate. Naumann, B. 314; acetone. (Naumann, B. 1904, 1; Eidmann, C.C. 1899, II. 1914)

sat. solution in glycol contain 0.5% (de Coninck, Bull. Ac. roy. Belg. 19)

$\text{Mn}_2\text{O}_3 + 7\text{H}_2\text{O}$ occurs as the min. Mallar-

omanganic sulphate, MnO , MnO_2 , H_2O .

rescent. Decomp. by H_2O . Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Fremy, C. R. 82.)

ic sulphate, $\text{Mn}_2(\text{SO}_4)_3$.

nely deliquescent. Sol. in H_2O with 1 of heat, and decomposition into a sulphate. Behaves similarly with H_2SO_4 . Sol. in traces in cold conc. H_2SO_4 . cold conc. $\text{HNO}_3 + \text{Aq}$. Sol. in conc. H_2SO_4 . Decomp. by absolute alcohol. A. 98. 53.)

yl sulphate, MnO_2 , SO_2 .

n H_2SO_4 , but solution decomp. if 10-60° Baumé. Solubility in 40° acid = 15%; 55°, 4-5%. Solution can be heated to 60-80° without decomp. (Bad. Sodafabrik, C. C. 1905, II. 1398.)

ous hydrogen sulphate.

Mn_2O_3 is sol. in 20 pts. boiling conc. H_2SO_4 ; in boiling $\text{H}_2\text{SO}_4 + \text{Aq}$ of 1.6 sp. gr.

(Schultz)

(SO_4)₂ and $+ \text{H}_2\text{O}$. Sol. in H_2O with decomp.

ic hydrogen sulphate, $\text{Mn}_2\text{H}_2(\text{SO}_4)_4 + \text{O}$.

rescent. Decomp. by H_2O . Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Francke, J. pr. (2) 36. 251.)

Manganous hydrazine sulphate, $\text{MnH}_2(\text{SO}_4)_2 \cdot 2\text{N}_2\text{H}_4$.

1 pt. is sol. in 60 pts. H_2O at 18°. Stable in the air at 100°. (Curtius, J. pr. 1894, (2) 60. 331.)

Manganous nickel potassium sulphate,

MnSO_4 , NiSO_4 , $2\text{K}_2\text{SO}_4 + 12\text{H}_2\text{O}$. Sol. in H_2O . (Vohl, A. 94. 57.)

Manganous potassium sulphate, K_2SO_4 , $\text{MnSO}_4 + 2\text{H}_2\text{O}$.

$+ 4\text{H}_2\text{O}$. Efflorescent. (Pierre, A. ch. (3) 16. 239.)

2MnSO_4 , K_2SO_4 . (Mallet, C. N. 1899, 80. 301.)

Manganic potassium sulphate, $\text{K}_2\text{Mn}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

Decomp. by dissolving in H_2O . (Mitscherlich.)

Manganomanganic potassium sulphate, $\text{Mn}_2(\text{SO}_4)_3$, $5\text{K}_2\text{SO}_4 = 3\text{Mn}(\text{SO}_4)_2$, 2MnSO_4 , $5\text{K}_2\text{SO}_4$.

Decomp. by much H_2O . Sol. in dil. or conc. H_2SO_4 . Insol. in alcohol or ether. (Francke, J. pr. (2) 36. 166.)

Manganous potassium zinc sulphate, MnSO_4 , $2\text{K}_2\text{SO}_4$, $\text{ZnSO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Vohl.)

Manganous rubidium sulphate, MnSO_4 , $\text{Rb}_2\text{SO}_4 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Tutton, Chem. Soc. 63. 337.)

1 l. H_2O dissolves 357 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

$+ 2\text{H}_2\text{O}$. (Wyruboff, Bull. Soc. Min. 1891, 14. 242.)

2MnSO_4 , Rb_2SO_4 . (Wyruboff.)

Manganic rubidium sulphate, $\text{Mn}_2(\text{SO}_4)_3$, $\text{Rb}_2\text{SO}_4 + 24\text{H}_2\text{O}$.

Deliquescent. (Christensen, Z. anorg. 1901, 27. 333.)

Manganous sodium sulphate, MnSO_4 , Na_2SO_4 , $+ 2\text{H}_2\text{O}$. Deliquescent in moist air.

(Geiger.)

$+ 4\text{H}_2\text{O}$. Sol. in 1.2 pts. boiling H_2O . (Geiger.)

Manganous sulphate ammonia, $\text{MnSO}_4 \cdot 4\text{NH}_3$.

Decomp. by H_2O . (Rose, Pogg. 20. 148.)

Manganous sulphate cupric oxide, MnSO_4 , $2\text{CuO} + 3\text{H}_2\text{O}$.

(Mailhe, A. ch. 1902, (7) 27. 392.)

MnSO_4 , $3\text{CuO} + z\text{H}_2\text{O}$. (Recours, C. R. 1901, 133. 1415.)

MnSO_4 , $24\text{CuO} + z\text{H}_2\text{O}$. (Recours.)

Manganous sulphate hydrazine, $\text{MnSO}_4 \cdot 2\text{N}_2\text{H}_4$.

Very unstable.

Somewhat sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Franzen, Z. anorg. 1908, 60. 285.)

Manganous sulphate hydroxylamine, $\text{MnSO}_4 \cdot \text{NH}_2\text{OH} + 2\text{H}_2\text{O}$.

Insol. in H_2O . (Feldt, B. 1894, 27: 405.)

Mercurous sulphate, basic, $2\text{Hg}_2\text{O} \cdot \text{SO}_3 + \text{H}_2\text{O}$.

Sol. in 25,000 pts. H_2O at 20° . (Gouy, C. R. 1900, 130. 1401.)

Mercurous sulphate, Hg_2SO_4 .

Sol. in 500 pts. cold, and 300 pts. hot H_2O . (Wackenroder, A. 41. 319.)

Solubility in H_2O at $18^\circ = 7.8 \times 10^{-4}$ g. mol. per liter. (Wilsmore, Z. phys. Ch. 1900, 35. 305.)

1 l. H_2O dissolves 11.71×10^{-4} g.-mol. Hg_2SO_4 at 25° . (Drucker, Z. anorg. 1901, 28. 362.)

Solubility in H_2O at t° .

t°	In 100 pts. of the solution	
	Hg_2SO_4	H_2SO_4
16.5	0.055	0.008
33	0.060	0.018
50	0.065	0.037
75	0.074	0.063
91	0.084	0.071
100	0.092	0.071

(Barre, A. ch. 1911, (8) 24. 203.)

Solubility in H_2O at 25° is 20% greater than at 18° and $= 11.71 \times 10^{-4}$ g. mol. per l. By addition of increasing amts. of H_2SO_4 the solubility is somewhat, but not regularly, decreased, K_2SO_4 lowers solubility less than H_2SO_4 . (Drucker, Z. anorg. 1901, 28. 362.)

Easily sol. in dil. $\text{HNO}_3 + \text{Aq.}$ from which solution it is separated by dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Wackenroder, A. 41. 319.)

Abundantly sol. in hot, less sol. in cold dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Berzelius.)

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

H_2SO_4 normality	g.-mol. Hg_2SO_4 per litre
0.0400	8.31×10^{-4}
0.1000	8.78×10^{-4}
0.2000	8.04×10^{-4}

(Drucker, Z. anorg. 1901, 28. 362.)

Partially decomp. by hot NH_4 salts + Aq. (Miahle, A. ch. (3) 5. 179.)

5 times more sol. in sat. $\text{CdSO}_4 + \text{Aq}$ than in H_2O . (Hulett, Phys. Rev. 1907, 25. 16.)

Sat. ZnSO_4 or $\text{CdSO}_4 + \text{Aq}$ attack much less

than pure H_2O , yet the solubility of Hg_2SO_4 in these solutions is greater, i. e., 0.8 g. in 1 l. $\text{ZnSO}_4 + \text{Aq}$ and 1.1 g. in 1 l. $\text{CdSO}_4 + \text{Aq}$ at 20° . (Gouy, C. R. 1900, 130. 1402.)

Solubility in $0.2\text{N } \text{K}_2\text{SO}_4 + \text{Aq} = 9.05 \times 10^{-4}$ g. mol. per l. at 25° . (Drucker, Z. anorg. 1901, 28. 362.)

Solubility of Hg_2SO_4 in $\text{K}_2\text{SO}_4 + \text{Aq.}$

t°	g. per 100 g. sat. solution		
	K_2SO_4	Hg_2SO_4	free H_2SO_4
15	2.90	0.0475	0.0080
	5.70	0.0703	0.0093
	8.22	0.0912	0.0098
	8.77	0.0994	
	9.44	0.1080	0.0110
33	2.94	0.0677	0.0250
	5.68	0.1015	0.0350
	8.30	0.1364	0.0441
	10.70	0.1724	0.0438
	11.90	0.1902	0.0420
75	3.10	0.1344	0.1681
	5.75	0.2120	0.2135
	8.50	0.2951	0.2514
	13.20	0.4610	0.2503
	17.30	0.6440	0.2225

(Barre, A. ch. 1911, (8) 24. 202.)

About 3 times as sol. in sat. $\text{ZnSO}_4 + \text{Aq}$ as in distilled H_2O . (Wright, Phil. Mag. 5 1885, 19. 29.)

Mercuric sulphate, basic, $2\text{HgO} \cdot \text{SO}_3$.

(Mailhe, A. ch. 1902, (7) 27. 394.)

$3\text{HgO} \cdot \text{SO}_3$. (Mineral turpeth.)

Sol. in 2000 pts. cold and 600 pts. boiling H_2O . (Fourcroy, A. ch. 19. 307.)

Sol. in 43,478 pts. H_2O at 16° when pptd. cold, and in 32,258 pts. at 16° when pptd. at 100° . (Cameron, Z. anal. 19. 144.)

Sl. sol. in warm dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Rose.)

Solubility in H_2O is increased by addition of H_2SO_4 up to an acid content of 4.3 mol SO_3 to 93.7 mols. H_2O . (Hoitsema, Z. phys. Ch. 1895, 17. 665.)

Sol. in warm conc. HCl or $\text{HBr} + \text{Aq.}$ (Ditte.)

Sol. in alkali chlorides + Aq. (Miahle.)

Sol. in dil. HNO_3 or in $\text{HCl} + \text{Aq.}$ More easily sol. in the warm acids. (Ray, Chem. Soc. 1897, 71. 1099.)

$3\text{HgO} \cdot 2\text{SO}_3 + 2\text{H}_2\text{O}$. (Hoitsema, Z. phys. Ch. 1895, 17. 659.)

$4\text{HgO} \cdot 3\text{SO}_3$. (Hopkins, Sil. Am. J. 18 364.)

Mercuric sulphate, HgSO_4 .

Decomp. by H_2O into $3\text{HgO} \cdot \text{SO}_3$ and a sol. acid salt. Sol. in dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ Decomp. by all acids. (Berzelius.)

Sol. in warm conc. HCl or HBr + Aq; very sl. sol. in boiling conc. HI + Aq. (Ditte, A. ch. (5) 17. 124.)

Very sl. sol. in hot conc. HF. (Ditte, A. ch. 1879, (5) 17. 125.)

Sol. in HCN + Aq. (Mohr.)

Sol. with decomp. in NaCl + Aq. (Miahle.)

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in conc. alcohol.

Insol. in acetone. (Naumann, B. 1904, 37. 4329.); methyl acetate. (Naumann, B. 1909, 42. 3790.); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

Insol. in pyridine. (Schroeder, Dissert. 1901.)

+H₂O. Decomp. by H₂O. (Eisfeldt, Pharm. Centr. 1853. 812.)

Mercuromercuric sulphate, Hg₂O, 2HgO, SO₃.

Insol. in cold H₂O; not decomp. by boiling H₂O. Decomp. by HCl + Aq. (Brooke, Pogg. 66. 63.)

Hg₂SO₄, HgSO₄. (Baskerville, J. Am. Chem. Soc. 1897, 19. 875.)

Mercuric hydrogen sulphate, HgH₂(SO₄)₂.

(Braham, C. N. 42. 163.)

Mercuric potassium sulphate, 3HgSO₄, K₂SO₄ + 2H₂O.

Sol. in H₂O. (Hirzel, J. B. 1850. 332.)

Mercuric sulphate chloride ammonium chloride, 2HgSO₄, HgCl₂, 2NH₄Cl.

Decomp. with H₂O. Ether dissolves out HgCl₂. (Kosmann, A. ch. (3) 27. 238.)

Mercuric sulphate cyanide, HgSO₄, Hg(CN)₂, + 5H₂O.

Decomp. by conc. or warm acids. (Rupp, Arch. Pharm. 1912, 250. 280.)

Mercuric sulphate hydrazine, HgSO₄, N₂H₄.

Ppt. (Hofmann and Marburg, A. 1899, 305. 216.)

Mercuric sulphate hydrobromide, HgSO₄, 2HBr.

Sol. in H₂O without separation of basic sulphate. (Ditte, A. ch. (5) 17. 122.)

3HgO, SO₃, 6HBr. Sol. in H₂O. (Ditte.)

Mercuric sulphate hydrochloride, HgSO₄, HCl

Not attacked by HCl. Sl. sol. in HNO₃. (Baskerville, J. Am. Chem. Soc. 1901, 23. 895.)

HgSO₄, 2HCl. Sol. in H₂O without separation of a basic salt. Very sol. in warm H₂SO₄, solidifying on cooling if very conc., or crystallising if dil. (Ditte.)

Very deliquescent.

Very sol. in H₂O. (Baskerville, J. Am. Chem. Soc. 1901, 23. 895.)

+H₂O. (Baskerville, J. Am. Chem. Soc. 1901, 23. 895.)

3HgO, SO₃, 6HCl. Sol. in H₂O. (Ditte.)

Mercuric sulphate hydroxylamine, HgSO₄, 2NH₂OH + H₂O.

Decomp. by cold H₂O. (Adams, Am. Ch. J. 1902, 28. 209.)

Mercuric sulphate iodate iodide, 6(3HgO, 2SO₃), 6HgI₂, Hg(IO₃)₂.

Decomp. by H₂O and acids. (Brückner, M. 1907, 28. 961.)

Mercuric sulphate iodide, basic, 3HgO, 2SO₃, HgI₂.

3(3HgO, 2SO₃), 2HgI₂ + 10H₂O.

2(3HgO, 2SO₃), HgI₂ + 10H₂O. Very sol. in hot conc. HNO₃.

3HgO, 2SO₃, HgSO₄, HgI₂ + 10H₂O. (Ditte, C. R. 1905, 140. 1167.)

Mercuric sulphate iodide, HgSO₄, HgI₂.

Decomp. by H₂O, not by alcohol or ether. (Riegel, J. B. pr. Pharm. 11. 396.)

3HgSO₄, HgI₂. Decomp. by cold or hot H₂O. Sol. in H₂SO₄ + Aq. (Ditte, C. R. 1905, 140. 1165.)

4HgSO₄, HgI₂ + 15H₂O, and + 18H₂O. Decomp. by cold or hot H₂O.

Sol. in H₂SO₄. (Ditte.)

Mercuric sulphate phosphide.

See *Dimercuriphosphonium mercuric sulphate*.

Mercuric sulphate sulphide, basic, 2HgO, SO₃, HgS.

Somewhat sol. in HCl, H₂SO₄ and HNO₃. (Jacobson, Pogg. 1846, 68. 412.)

4HgO, 3SO₃, 2HgS + 4H₂O. Sl. sol. in H₂SO₄. (Estrup, Z. anorg. 1909, 62. 169.)

Mercuric sulphate sulphide, 2HgSO₄, HgS.

Sl. sol. in hot HCl, H₂SO₄, or HNO₃ + Aq. Easily sol. in hot aqua regia. (Jacobson, Pogg. 68. 410.)

2HgSO₄, HgS. (Palm, C. C. 1863. 122.)

HgSO₄, 2HgS. (Barfoed, J. B. 1864. 282.)

Sol. in aqua regia. (Denigès, Bull. Soc. 1915, (4) 17. 355.)

HgSO₄, 3HgS. Insol. in H₂O. Easily sol. in aqua regia; decomp. by HNO₃ into—

3HgSO₄, HgS. Insol. in all acids except aqua regia. (Spring, A. 199. 116.)

Molybdenum sesquisulphate (?).

Basic. Insol. in H₂O.

Neutral. Decomp. by H₂O into acid and basic salts.

Acid. Sol. in H₂O. (Berzelius.)

Molybdenum disulphate (?).

Sol. in H₂O.

Molybdenum sulphate, Mo₂O₃, 2SO₃.

Very slowly sol. in cold, more quickly sol. in hot H₂O. (Bailhache, C. R. 1901, 132. 476.)

7MoO₃, 2MoO₃, 7SO₃+Aq. (Péchar, C. R. 1901, 132. 630.)

Molybdic sulphate, MoO₃, SO₃.

Deliquescent. Sol. in H₂O. (Schultz-Sellack, B. 4. 14.)

MoO₃, 3SO₃+2H₂O. Deliquescent. Partially sol. in H₂O. (Anderson, Berz. J. B. 22. 161.)

Does not exist. (Schultz-Sellack.)

Molybdenum sulphate ammonia,

5NH₃, MoO₃SO₃, 7MoO₃+8H₂O.

3NH₃, MoO₃SO₃, 7MoO₃+10H₂O.

Both very sol. in water but less sol. in H₂O containing ammonium salts. (Péchar, C. R. 1901, 132. 630.)

Neodymium sulphate, basic, Nd₂O₃, SO₃.

Insol. in H₂O.

Nearly insol. in dil. acids. (Wöhler, B. 1913, 46. 1730.)

Insol. in H₂O. (Matignon, C. R. 1902, 134. 658.)

Neodymium sulphate, Nd₂(SO₄)₃+8H₂O.

Solubility in 100 pts. H₂O at t°.

t°	pts. Nd ₂ (SO ₄) ₃
0	9.50
16	7.05
30	5.04
50	3.72
80	2.70
100	2.21

(Muthmann and Rolig, B. 1898, 31. 1728.)

Neodymium hydrogen sulphate, Nd(SO₄H)₃.

(Brauner, Z. anorg. 1904, 38. 331.)

Neodymium potassium sulphate.

Cryst. modification more sol. in cold than in hot H₂O. (Boudouard, C. R. 1898, 126. 901.)

Nickel sulphate, basic.

Very sl. sol. in H₂O. (Berzelius.)

6NiO, 5SO₃+4H₂O. (Athanasesco, C. R. 108. 271.)

7NiO, 7H₂O, SO₃+3H₂O. Nearly insol. in H₂O. (Habermann, M. 5. 432.)

5NiO, SO₃; 5NiO, 2SO₃; and 5NiO, 3SO₃. (Pickering, Chem. Soc. 1907, 91. 1985.)

6NiO, SO₃. (Strömholm. C.C. 1903, 1. 1222.)

Nickel sulphate, NiSO₄.

100 pts. H₂O dissolve pts. NiSO₄ at t°:

2° 16° 20° 23° 31°
30.4 37.4 39.7 41 45.3 pts. Ni

41° 50° 53° 60° 70°
49.1 52 54.4 57.2 61.9 pts. Ni

(Tobler, A. 98. 193.)

100 pts. of sat. solution contain: at 11-14° at 18 20°, 30.77 pts. anhydrous salt. (v. H. A. B. 53, 2. 221.)

100 pts. H₂O at 112.5° dissolve 185.71 pts. (Griffiths.)

NiSO₄+7H₂O is sol. in 3 pts. H₂O at 12.5° (puti.)

100 pts. H₂O at 15.5° dissolve 75.6 pts. 7H₂O.

Sat. NiSO₄+Aq contains at:

-3° +2° 5° 11° 17° 54°
21.7 22.7 23.1 25.2 26.6 33.6%

68° 74° 92° 97° 110° 117° 119°
38.2 38.7 42.4 44.2 46.5 48.8 49.4%

(Étard, A. ch. 1894, (7) 2. 552.)

See also below under hydrated salts.

Sp. gr. of NiSO₄+Aq containing g. 7H₂O in 1000 g. H₂O at 23.5°.

140.5 g. (= 1/2 mol.) 281 421.5

1.073 1.136 1.190

602.5 843 983.5 11

1.280 1.317 1.349 1.3

Containing NiSO₄ (anhydrous):

77.5 g. (= 1/2 mol.) 155 232.5 310 387

1.079 1.153 1.224 1.292 1.3

(Gerlach, Z. anal. 28. 468.)

Sp. gr. of NiSO₄+Aq at 0°. S = pts

in 100 pts. solution; S₁ = mols. N

100 mols. solution.

S	S ₁	Sp.
4.2930	0.581	1.0
3.9591	0.476	1.0
3.2845	0.392	1.0
2.5043	0.297	1.0
1.6131	0.189	1.0
0.8327	0.097	1.0

(Charpy, A. ch. (6) 29. 26.)

Sp. gr. of NiSO₄+Aq at room tem

taining:

10.62 18.19 25.35% NiSO₄

1.0925 1.1977 1.3187

(Wagner, W. Ann. 1883, 18. 272)

Sp. gr. of NiSO₄+Aq at 25°.

Concentration of NiSO ₄ +Aq	Sp. gr.
1-normal	1.077
1/2 " "	1.089
1/3 " "	1.019
1/4 " "	1.001

(Wagner, Z. phys. Ch. 1890, 5. 3

For solubility of $\text{NiSO}_4 + \text{Na}_2\text{SO}_4$ in H_2O , see under $\text{NiSO}_4 + 7\text{H}_2\text{O}$ and $\text{NiNa}_2(\text{SO}_4)_2$.

100 pts. sat. $\text{NiSO}_4 + \text{ZnSO}_4 + \text{Aq}$ at $18-20^\circ$ contain 35.45 pts. of the two salts. (v. Hauer.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

$\text{HC}_2\text{H}_3\text{O}_2$ precipitates it completely from aqueous solution. (Persoz.)

100 pts. absolute methyl alcohol dissolve 0.5 pt. NiSO_4 at 18° . (de Bruyn, Z. phys. Ch. 10. 783.)

Solubility of $\text{NiSO}_4, 3\text{CH}_3\text{O} + 3\text{H}_2\text{O}$ in methyl alcohol at 14° .

P = % anhydrous NiSO_4 in the sat. solutions.

Alcohol % by wt.	P	In 1000 mol. of the solution		
		Mol. NiSO_4	Mol. CH_3O	Mol. H_2O
100	3.72	7.75	969	23.2
97.5	0.77	1.65	950	48.5
95	0.455	0.96	908	91
92.5	0.50	1.0	871	128
90	0.70	1.6	830	168
89	1.01	2.0	814	184
88	1.25	2.4	800	198
87	1.48	2.9	781	216
86	1.73	3.2	767	230
85	1.93	3.6	755	241

(de Bruyn, R. t. c. 1903, 22. 418.)

This salt is more sol. in ethyl alcohol than in methyl alcohol.

See also under hydrated salts.

For solubility of NiSO_4 in ethyl alcohol, see under hydrated salts.

100 g. sat. solution in glycol contain 9.7 g. NiSO_4 at ord temp. (de Coninck, Bull. Ac. Roy. Belg. 1905. 359.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); ethyl acetate, (Naumann, B. 1904, 37. 3602.)

Very sl. sol. in acetone. (Krug and M'Elroy.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

+ H_2O . Very sl. sol. in abs. methyl alcohol. After standing in contact with it for 8-9 months, the solution contains 1.34% NiSO_4 . (de Bruyn, R. t. c. 1903, 22. 414.)

+ $2\text{H}_2\text{O}$. Cryst. from sat. $\text{NiSO}_4 + \text{Aq}$ at 131° . (Steele and Johnson.)

+ $4\text{H}_2\text{O}$. Solubility in methyl alcohol + Aq at 10° . Time = 24 hrs. P = % anhydrous salt in the sat. solutions.

Alcohol of 100% 80% 50% 20% water
P 7.38 0.66 1.43 14.8 25.1

(de Bruyn, R. t. c. 1903, 22. 414.)

+ $6\text{H}_2\text{O}$. Two modifications. α -blue, tetragonal; β -green, monoclinic.

Solubility of α - $\text{NiSO}_4, 6\text{H}_2\text{O}$ in H_2O at t° .

Salt used	t°	g. NiSO_4 in 100 g. H_2O	Salt remaining
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	32.3	43.57	$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)
"	33.0	43.35	"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$	34.0	43.84	"
"	35.6	43.79	"
$\text{NiSO}_4 + 6\text{H}_2\text{O}$	44.7	48.05	"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$	44.7	47.97	"
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	50.0	50.15	"
"	51.0	50.66	"
"	52.0	52.34	"
"	53.0	52.34	"

(Steele and Johnson.)

Solubility of β - $\text{NiSO}_4, 6\text{H}_2\text{O}$ in H_2O at t° .

Salt used	t°	g. NiSO_4 per 100 g. H_2O	Salt remaining
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	54.5	52.50	$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (green)
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	57.0	53.40	"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$			"
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	60.0	54.84	"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$			"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$	69.0	58.38	"
"	70.0	59.44	"
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	73.0	60.72	"
$\text{NiSO}_4 + 6\text{H}_2\text{O}$ (blue)	80.0	63.17	"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$			"
$\text{NiSO}_4 + 7\text{H}_2\text{O}$	89.0	67.90	"
"	99.0	76.71	"

(Steele and Johnson, Chem. Soc. 1904, 85. 118.)

Tr. point from α - to β - salt = 53.3° . (Steele and Johnson.)

100 pts. absolute methyl alcohol dissolve 31.6 pts. $\text{NiSO}_4 + 6\text{H}_2\text{O}$ at 17° ; 100 pts. 93.5% methyl alcohol dissolve 7.8 pts. $\text{NiSO}_4 + 6\text{H}_2\text{O}$ at 18° ; 100 pts. 50% methyl alcohol dissolve 1.9 pts. $\text{NiSO}_4 + 6\text{H}_2\text{O}$ at 18° . (de Bruyn, Z. phys. Ch. 10. 786.)

α -Salt. Solubility in methyl alcohol + Aq at 14° . Time = 5 to 6 hrs.

Alcohol % by wt.	% NiSO_4	In 1000 mol. of the solution		
		Mol. NiSO_4	Mol. CH_3O	Mol. H_2O
100	12.4	26	794	180
97.5	10.6	22.3	808	170
95	6.5	14	817	169
92.5	3.06	5.9	838	156
90	1.18	2.3	821	177
85	0.315	0.57	757	242
80	0.25	0.4	688	312
60	0.46	0.8	453	546
40	2.43	3.5	265	732
20	14.7	21	105	874
0 (water)	26.0	39	0	961

(de Bruyn, R. t. c. 1903, 22. 412.)

β -salt. Solubility in methyl alcohol + Aq at 14°. Time = 24 hrs.

Alcohol % by wt.	g. NiSO ₄	In 1000 mol. of the solution		
		Mol. NiSO ₄	Mol. CH ₃ O	Mol. H ₂ O
100	15.7	33.8	763	203
97.5	12.4	26	781	193
95	10.0	20.3	784	196
92.5	5.61	11.1	800	189
III	2.35	4.5	810	185
89	1.79
88	1.29
87	0.97
86	0.735
85	0.61	1.3	755	244
80	0.415	0.7	682	317
60	0.75	1.3	453	546
40	3.11	4.0	264	732
20	14.1	21.0	105	874
0 (water)	27.2	40.0	0	960

(de Bruyn.)

+7H₂O.
Solubility in 100 pts. H₂O at t°, using
NiSO₄ + 7H₂O.

t°	Pts. NiSO ₄	t°	Pts. NiSO ₄	t°	Pts. NiSO ₄
0	29.3	33	45.5	66	63.6
1	29.7	34	46.0	67	64.1
2	30.1	35	46.5	68	64.7
3	30.5	36	47.0	69	65.3
4	31.0	37	47.5	70	65.9
5	31.5	38	48.0	71	66.5
6	32.0	39	48.5	72	67.0
7	32.5	40	49.0	73	67.6
8	33.0	41	49.6	74	68.2
9	33.5	42	50.1	75	68.8
10	34.0	43	50.6	76	69.3
11	34.5	44	51.2	77	69.9
12	35.0	45	51.7	78	70.5
13	35.5	46	52.3	79	71.1
14	36.0	47	52.8	80	71.7
15	36.5	48	53.4	81	72.3
16	37.0	49	53.9	82	72.9
17	37.5	50	54.5	83	73.5
18	38.0	51	55.0	84	74.1
19	38.5	52	55.6	85	74.6
20	39.0	53	56.1	86	75.2
21	39.5	54	56.7	87	75.8
22	40.0	55	57.3	88	76.4
23	40.5	56	57.9	89	77.0
24	41.0	57	58.4	90	77.6
25	41.5	58	59.0	91	78.2
26	42.0	59	59.6	92	78.8
27	42.5	60	60.2	93	79.4
28	43.0	61	60.7	94	80.1
29	43.5	62	61.3	95	80.7
30	44.0	63	61.9	96	81.3
31	44.5	64	62.4	97	81.9
32	45.0	65	63.0	98	82.5

Solubility in 100 pts. H₂O at t°.—Continued

t°	Pts. NiSO ₄	t°	Pts. NiSO ₄	t°	Pts. NiSO ₄
99	83.1	103	85.6	107	86.1
100	83.7	104	86.2	IV	86.7
101	84.3	105	86.8	108	88.7
102	84.9	106	87.5		

(Mulder, calculated from his own and Tobler's determinations, Scheik. Verhandel 1864 70.)

Solubility in H₂O at t°

Salt used	t°	g. NiSO ₄ in 100 g H ₂ O	Salt remaining
NiSO ₄ + 7H ₂ O	-5	25.74	NiSO ₄ + 7H ₂ O
"	0	27.22	"
"	9	31.55	"
"	15	34.19	"
"	22.6	37.90	"
"	22.8	38.88	"
"	30.0	42.46	"
NiSO ₄ + 6H ₂ O	30.0	42.47	"
NiSO ₄ + 7H ₂ O	32.3	44.02	"
"	33.0	45.74	"
"	34.0	45.5	"

(Steele and Johnson, Chem. Soc. 1904, 88, 116.)

M.-pt. of NiSO₄ + 7H₂O = 98–100°. (Tilden, Chem. Soc. 45, 409.)

Tr. point from α -6H₂O salt to 7H₂O salt = 31.5°. (Steele and Johnson.)

Exists also in an unstable, more soluble modification. (Fedorow, C. C. 1903, II, 95.)

Solubility of NiSO₄·7H₂O + Na₂SO₄·10H₂O in 100 g. H₂O at t°.

t°	grams NiSO ₄	grams Na ₂ SO ₄
0	22.46	10.09
5	25.28	15.245
10	28.26	20.64

(Koppel, Z. phys. Ch. 1905, 81, 401.)
See also under NiNa₂(SO₄)₄.

100 pts. absolute methyl alcohol dissolve 46 pts. NiSO₄ + 7H₂O at 17°; 100 pts. absolute methyl alcohol dissolve 24.7 pts. NiSO₄ + 7H₂O at 4°; 100 pts. 93.5% methyl alcohol dissolve 10.1 pts. NiSO₄ + 7H₂O at 4°; 100 pts. 50% methyl alcohol dissolve 2 pts. NiSO₄ + 7H₂O at 4°. (de Bruyn, Z. phys. Ch. 20, 786.)

Solubility in methyl alcohol+Aq at 14°. Time, 5 to 6 hrs.

Alcohol % by wt.	% NiSO ₄	In 1000 mol. of solution		
		Mol. NiSO ₄	Mol. CH ₃ O	Mol. H ₂ O
100	16.8	35.7	714	250
97.5	13.9	29	734	237
95	11.6	23.6	742	234.5
92.5	8.12	16.2	760	224
90	5.78	11.2	758	231
85	1.52	3	744	253
84	1.06
83	0.985
82	0.83
81	0.665
80	0.653	1.2	687	312
60	0.805	1.3	453	546
45	1.73
40	2.78	4	264	732
35	4.55
30	6.33
20	13.7	20	105	875
0 (water)	26.4	39.5	0	960.5

(de Bruyn, R. t. c. 1903, 22. 411.)

100 g. absolute ethyl alcohol dissolve 1.3 g. NiSO₄+7H₂O at 4°, and 2.2 g. at 17°. (de Bruyn, Z. phys. Ch. 10. 786.)
Min. Morenosite.

Nickel hydrazine sulphate, NiH₂(SO₄)₂, 2N₂H₄.

1 pt. is sol. in 275.5 pts. H₂O at 18°. Sl. sol. in hot H₂O. Sol. in HNO₃ with decomp.; insol. in HCl. Sol. in NH₄OH+Aq. (Curtius, J. pr. 1894, (2) 50. 331.)

Nickel potassium sulphate, NiSO₄, K₂SO₄+6H₂O.

Sol. in 8-9 pts. H₂O. (Tupputi.)
100 pts. H₂O dissolve at:
0° 10° 14° 20° 30°
5.3 8.9 10.5 13.8 18.6 pts. anhydrous salt,
36° 49° 55° 60° 75°
20.4 27.7 32.4 35.4 45.6 pts. anhydrous salt.
(Tobler, A. 95. 193.)

Saturated solution contains at:
20° 40° 60° 80°
8.7 12.3 17.6 22.0% anhydrous salt.
(v. Hauer, J. pr. 74. 433.)

1 l. H₂O dissolves 68.8 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Nickel potassium zinc sulphate, NiSO₄, 2K₂SO₄, ZnSO₄+12H₂O.
Sol. in H₂O. (Vohl, A. 94. 51.)

Nickel rubidium sulphate, NiSO₄, Rb₂SO₄+6H₂O.

Sol. in H₂O. (Tutton, Chem. Soc. 63. 337.)
1 l. H₂O dissolves 59.8 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Nickel sodium sulphate, NiNa₂(SO₄)₂+4H₂O.

Solubility of NiNa₂(SO₄)₂.4H₂O in 100 g. H₂O at t°.

t°	grams NiSO ₄	grams Na ₂ SO ₄
20	29.31	26.87
25	27.33	25.33
30	24.64	22.58
35	23.66	21.67
40	21.88	20.65

(Koppel, Z. phys. Ch. 1905, 52. 401.)

Solubility of NiNa₂(SO₄)₂.4H₂O + NiSO₄. 7H₂O in 100 g. H₂O at t°.

t°	grams NiSO ₄	grams Na ₂ SO ₄	t°	grams NiSO ₄	grams Na ₂ SO ₄
18.5	30.70	25.805	30	34.98	19.825
20	31.59	25.355	35	36.01	16.435
25	33.11	23.07	40	37.935	14.295

(Koppel.)

Solubility of NiNa₂(SO₄)₂.4H₂O + Na₂SO₄. 10H₂O in 100 g. H₂O at t°.

t°	grams NiSO ₄	grams Na ₂ SO ₄
18.5	26.14	29.455
20	24.07	31.365
25	18.81	37.13
30	9.87	44.25

(Koppel.)

Solubility of NiNa₂(SO₄)₂.4H₂O + Na₂SO₄ (anhydrous) in 100 g. H₂O at t°.

t°	grams NiSO ₄	grams Na ₂ SO ₄
35	7.13	49.595
40	7.245	49.03

(Koppel.)

Nickel thallium sulphate, NiSO₄, Tl₂SO₄+6H₂O.

Easily sol. in H₂O. Can be recryst. from little H₂O without decomp. (Werther, J. pr. 92. 132.)

1 l. H₂O dissolves 46.1 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Nickel zinc sulphate, NiSO₄, ZnSO₄+13H₂O.

Sol. in 3-4 pts. cold H₂O. Insol. in alcohol. (Tupputi, 1811.)

Completely sol. in NH₄OH+Aq.
2NiSO₄, 2ZnSO₄, H₂SO₄. (Étard, C. R. 87. 602.)

Nickel sulphate ammonia, $\text{NiSO}_4, 6\text{NH}_3$.

Sol. in H_2O with separation of hydroxide. (Rose, Pogg. 20. 151.)

$\text{NiSO}_4, 5\text{NH}_3 + 3\frac{1}{2}\text{H}_2\text{O}$. Deliquescent. (André, C. R. 106. 936.)

$\text{NiSO}_4, 4\text{NH}_3 + 2\text{H}_2\text{O}$. Easily sol. in H_2O . Can be recrystallized out of little H_2O . Insol. even in dil. alcohol. (Erdmann.)

Nickel sulphate cupric oxide, $\text{NiSO}_4, 2\text{CuO} + 6\text{H}_2\text{O}$.

(Mailhe, Bull. Soc. 1902, (3) 27. 172.)

$2\text{NiSO}_4, 3\text{CuO} + 10\text{H}_2\text{O}$, and $+12\text{H}_2\text{O}$. (Mailhe.)

$5\text{NiSO}_4, 16\text{CuO} + x\text{H}_2\text{O}$. (Recoura, C. R. 1901, 132. 1415.)

$\text{NiSO}_4, 20\text{CuO} + x\text{H}_2\text{O}$. (Recoura.)

Nickel sulphate, hydrazine, $\text{NiSO}_4, 3\text{N}_2\text{H}_4$.

Ppt. (Curtius, J. pr. 1894, (2) 50. 343.)

Nickel sulphate hydroxylamine, $\text{NiSO}_4, 6\text{NH}_2\text{OH}$.

Decomp. by H_2O . (Uhlenhut, A. 1899, 307. 334.)

Nitrosyl sulphate, $\text{H}(\text{NO})\text{SO}_4$.

See Nitrosulphonic acid.

Osmious sulphate.

Easily sol. in H_2O and alcohol.

Osmic sulphate.

Sol. in H_2O . (Berzelius.)

Palladous sulphate, basic, $\text{PdSO}_4, 7\text{PdO} + 6\text{H}_2\text{O}$, and $10\text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in $\text{HCl} + \text{Aq}$. (Kane.)

Palladous sulphate, $\text{PdSO}_4 + 2\text{H}_2\text{O}$.

Deliquescent in moist air; very sol. in H_2O , but decomp. by much H_2O , with separation of a basic salt. (Kane.)

Phosphoryl sulphate, $(\text{PO})_2(\text{SO}_4)_2$ (?).

Possible composition of Weber's (B. 20. 86) $\text{P}_2\text{O}_5, 3\text{SO}_3$ (?).

$3\text{P}_2\text{O}_5, 2\text{SO}_3$. Immediately decomp. by H_2O . (Adie, C. N. 1891, 63. 102.)

Platinic sulphate, $\text{Pt}(\text{SO}_4)_2$.

Deliquescent. Sol. in H_2O , alcohol, or ether; also in H_3PO_4 , HCl , and $\text{HNO}_3 + \text{Aq}$. (Berzelius.)

$\text{H}_2\text{PtO}_4, \text{SO}_4$. Sol. in H_2O . (Blondel, A. ch. 1905, (8) 6. 109.)

$\text{PtO}_2, \text{SO}_3 + 4\text{H}_2\text{O}$. Ppt. Decomp. by H_2O . Sol. in H_2SO_4 .

$\text{PtSO}_4(\text{OH})_2, 4\text{Pt}(\text{OH})_4 + 3\text{H}_2\text{O}$. Ppt. (Prost, Bull. Soc. (2) 46. 156.)

$\text{Pt}_2\text{SO}_4\text{O}_{11} + 16\text{H}_2\text{O}$. As above. (Prost.)

Platinum hydroxylamine sulphate, $\text{Pt}(\text{NH}_2\text{OH})_2\text{SO}_4$.

Only sl. sol. in H_2O ; sol. in dil $\text{H}_2\text{SO}_4 + \text{Aq}$. (Uhlenhut, A. 1900, 311. 123.)

Platinic potassium sulphate, basic.

Insol. in boiling H_2O , HNO_3 , H_2SO_4 , H_3PO_4 , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{NH}_4\text{OH} + \text{Aq}$. Easily sol. in boiling $\text{HCl} + \text{Aq}$. Sl. decomp. by aqua regia. (E. Davy.)

$\text{Pt}_{10}(\text{SO}_4)_2\text{O}_{10}, 3\text{K}_2\text{SO}_4 + 34\text{H}_2\text{O}$. Insol. in H_2O . (Prost, Bull. Soc. (2) 46. 156.)

$\text{Pt}_{12}(\text{SO}_4)_2\text{O}_{12}, 5\text{K}_2\text{SO}_4 + 34\text{H}_2\text{O}$. As above. (Prost.)

Platinum rubidium sulphate, $\text{Pt}_2\text{Rb}_2(\text{SO}_4)_4 + 17\text{H}_2\text{O}$.

Sol. in H_2O . (Prost, Bull. Soc. (2) 46. 156.)

Platinum sulphate sulphocarbamide, $\text{PtSO}_4, 4\text{CS}(\text{NH}_2)_2$.

Insol. in H_2O . Sol. in conc. H_2SO_4 without decomp. (Kurnakow. J. pr. 1894, (2) 50. 489.)

Potassium sulphate, K_2SO_4 .

Not hygroscopic in the ordinary sense of the word. 100 pts. K_2SO_4 over H_2O at $14-20^\circ$ absorb 58 pts. H_2O in 22 days, and finally deliquesce completely. (Mulder.)

12 pts. K_2SO_4 mixed with 100 pts. H_2O lower the temp. 3.3° . (Rüdorff, B. 2. 68.)

100 pts. H_2O dissolve with absorption of heat at 0° :

8.36 pts. K_2SO_4 . (Gay-Lussac.)

8.46 " (Mulder.)

8.5 " (Gerardin.)

7.31 " (Möller, Pogg. 117. 386.)

7.3-7.9 " (Nordenskiöld, Pogg. 126. 314.)

100 pts. H_2O at 0° dissolve 8.36 pts. K_2SO_4 ; at 12.7° , 10.57 pts.; at 49.06° , 16.91 pts.; at 63.90° , 19.29 pts.; at 101.50° , 26.33 pts. (Gay Lussac, A. ch. (2) 11. 311.)

Solubility in 100 pts. H_2O at t° .

t°	Pts. K_2SO_4	t°	Pts. K_2SO_4
0	7.8	47.0	16.0
15.65	10.3	70.2	20.3
28.1	12.8	98.0	23.9

(Nordenskiöld, Pogg. 126. 341.)

100 pts. sat. K_2SO_4 at 101.7° contain 17.5 pts. K_2SO_4 or 100 pts. H_2O at 101.25° dissolve 21.212 pts. K_2SO_4 . (Griffiths.)

100 pts. H_2O at 102.8° dissolve 29 pts. K_2SO_4 (Penny); at 15° , 7.3-8.25 pts. (Ure's Dict.); at 100° , 20 pts. (Ure's Dict.); at 100° , 24.2 pts. (Wenzel).

Sol. in 9.081 pts. H_2O at 15° (Gerlach); in 16 pts. at 15° , and 5 pts. at 100° (Bergmann); in 18 pts. cold, and 5 pts. boiling H_2O (Fourcroy); in 15 pts. cold, and 5 pts. boiling H_2O (Reid); in 12 pts. H_2O at 0° , and 4 pts. boiling H_2O (M. R. and P.); in 12 pts. H_2O at 18.75° (Abl.).

K_2SO_4 sat. at 15° has sp. gr. = 1.0774, and contains 10.055 pts. K_2SO_4 in 100 pts. H_2O . (Michel and Krafft, A. ch. (3) 41. 478.)

100 pts. H_2O dissolve 9.26 pts. K_2SO_4 at 15.6° , and sat. solution has sp. gr. = 1.177. (Page and Keightley, Chem. Soc. (2) 10. 566.)

Solubility in 100 pts. H₂O at t°.

t°	Pts. K ₂ SO ₄	t°	Pts. K ₂ SO ₄	t°	Pts. K ₂ SO ₄
0	8.5	35	13.1	70	19.8
1	8.6	36	13.3	71	20.0
2	8.7	37	13.4	72	20.2
3	8.8	38	13.6	73	20.4
4	9.0	39	13.8	74	20.6
5	9.1	40	14.0	75	20.8
6	9.2	41	14.2	76	21.0
7	9.3	42	14.3	77	21.2
8	9.4	43	14.5	78	21.4
9	9.5	44	14.7	79	21.6
10	9.7	45	14.9	80	21.8
11	9.8	46	15.1	81	22.0
12	9.9	47	15.3	82	22.2
13	10.0	48	15.5	83	22.4
14	10.2	49	15.6	84	22.6
15	10.3	50	15.8	85	22.8
16	10.4	51	16.0	86	23.0
17	10.5	52	16.2	87	23.2
18	10.7	53	16.4	88	23.4
19	10.8	54	16.6	89	23.6
20	10.9	55	16.8	90	23.9
21	11.1	56	17.0	91	24.1
22	11.2	57	17.2	92	24.3
23	11.3	58	17.4	93	24.5
24	11.5	59	17.6	94	24.7
25	11.6	60	17.8	95	25.0
26	11.7	61	18.0	96	25.2
27	11.9	62	18.2	97	25.5
28	12.0	63	18.4	98	25.7
29	12.2	64	18.6	99	25.9
30	12.3	65	18.8	100	26.2
31	12.5	66	19.0	101	26.4
32	12.6	67	19.2	102	26.6
33	12.8	68	19.4	102.25	26.75
34	13.0	69	19.6		

(Mulder, calculated from his own and other experiments, Scheik. Verhandel. 1864. 50.)

If solubility S = pts. anhydrous salt in 100 pts. of solution, $S = 7.5 + 0.1070t$ from 0° to 163°. Solubility from 163° to 220° is constant at 25. (Étard, C. R. 106. 208.)

Solubility of K₂SO₄ in 100 pts. H₂O at t°.

t°	Pts. K ₂ SO ₄	t°	Pts. K ₂ SO ₄	t°	Pts. K ₂ SO ₄
16	9.76	39	14.21	120	26.5
20	10.30	54	17.39	143	28.8
28	12.59	98	23.91	170	32.9
36	13.28

(Tilden and Shenstone, Phil. Trans. 1864. 23.)

Solubility of K₂SO₄ in H₂O. 100 pts. H₂O dissolve at:

4.3°	18.4°	69.9°
8.16	10.8	19.7 pts. K ₂ SO ₄ .

(Andreas, J. pr. (2) 29. 456.)

100 ccm. H₂O dissolve 12.04 g. K₂SO₄ at 25°. (Trevor, Z. phys. Ch. 7. 468.)

Sat. K₂SO₄ + Aq contains at:

21°	23°	60°	99°	130°
10.1	10.3	14.5	19.1	21.1% K ₂ SO ₄ ,

130°	152°	175°	195°	220°
21.3	22.8	24.5	23.8	24.6% K ₂ SO ₄ .

(Étard, A. ch. 1894, (7) 2. 549.)

Solubility of K₂SO₄ in H₂O at t°.

G. K₂SO₄ per 100 g. H₂O

t°	K ₂ SO ₄	Sp. gr.	t°	K ₂ SO ₄	Sp. gr.
0.40	7.47	1.0589	58.95	18.01	1.1089
15.70	10.37	1.0770	74.85	20.64	1.1157
31.45	13.34	1.0921	89.70	22.80	1.1194
42.75	15.51	1.1010	101.1*	24.21	1.1207

* = b.-pt.

(Berkeley, Phil. Trans. Roy. Soc. 1904, 203. A, 189.)

100 g. H₂O dissolve 0.133 gram-equivalent K₂SO₄ at 25° (Van't Hoff and Meyerhoffer, Z. phys. Ch. 1904, 49. 315.)

1 l. sat K₂SO₄ + Aq at 25° contains 0.617 mols. K₂SO₄. (Herz, Z. anorg. 1911, 73. 274.)

Solubility of K₂SO₄ in H₂O at t°.

t°	% K ₂ SO ₄
4.78	7.82
30.05	11.43
54.20	14.77
68.90	16.40

(Le Blanc and Schmandt, Z. phys. Ch. 1911, 77. 614.)

100 g. H₂O dissolve 12.10 g. K₂SO₄ at 25°. (Amadori, Rend. Acc. Linc. 1912, (5) 21. II. 667.)

Solubility of K₂SO₄ in H₂O at various pressures. Figures denote pts. K₂SO₄ contained in 100 pts. sat. K₂SO₄ + Aq at t° and A pressure in atmospheres.

A	0°	15°	15.5°	16.2°
1	6.81	9.14	9.24	9.35
20	7.14	...	9.44	9.54
30	7.14

(Möller, Pogg. 117. 386.)

Sat. K₂SO₄ + Aq boils at 101.5°, and contains 26.33 pts. K₂SO₄ to 100 pts. H₂O (Gay-Lussac); at 101.7°, and contains 21.2 pts. K₂SO₄ to 100 pts. H₂O (Griffiths); at 102.25°, and contains 26.75 pts. K₂SO₄ to 100 pts. H₂O (Mulder); boils at 103° (Kremers).

Crust forms at 101.7°, and solution contains 25.3 pts. K₂SO₄ to 100 pts. H₂O; highest

Sp. gr. of sat. K ₂ SO ₄ +Aq at t°.		
t°	g. K ₂ SO ₄ sol. in 100 g. H ₂ O	Sp. gr.
0	7.35	1.058
10	9.22	1.069
20	11.11	1.081
30	12.97	1.089
40	14.76	1.097
50	16.50	1.106
60	18.17	1.114
70	19.75	1.121

(Tschernaj, J. Russ. phys. Chem. Soc. 1912, 44. 1565.)

Sol. in conc. acids; not pptd. by glacial HC₂H₃O₂. Insol. in KOH+Aq of 1.35 sp. gr. (Liebig, A. 11. 262.)

Solubility of K₂SO₄ in H₂SO₄+Aq at 18°.

Mols. per 100 mols. solution		Solid phase
K ₂ SO ₄	H ₂ SO ₄	
1.10	0	K ₂ SO ₄
1.59	0.95	"
2.49	2.70	"
2.75	3.17	K ₂ SO ₄ , KHSO ₄
2.75	3.74	"
2.83	5.08	"
2.80	5.79	K ₂ SO ₄ , 3KHSO ₄
2.61	5.61	K ₂ SO ₄ , 6KHSO ₄
2.25	6.19	" +KHSO ₄
1.08	7.94	KHSO ₄
0.77	9.2	"
0.44	22.7	"

(Stortenbecker, R. t. c. 1902, 21. 407.)

Solubility in H₂SO₄+Aq at 0°.

1000 g. of the solution contain		Solid phase
Mols H ₂ SO ₄	Mols K ₂ SO ₄	
...	0.393	K ₂ SO ₄
0.37	0.53	"
0.75	0.64	"
1.08	0.74	K ₂ SO ₄ +K ₂ H(SO ₄) ₂
1.13	0.73	K ₂ H(SO ₄) ₂
1.44	0.71	"
1.66	0.69	"
1.89	0.66	K ₂ H(SO ₄) ₂ +Ka
1.88	0.69	"
2.15	0.59	Ka+Kb
2.12	0.61	"
2.29	0.54	Kb
2.30	0.53	Kb+KHSO ₄
2.33	...	"
2.48	0.43	KHSO ₄
3.08	0.28	"
4.43	0.12	"
5.27	0.09	"

These results show that at 0° there exist

between K₂H(SO₄)₂ and KHSO₄, two acid sulphates, Ka and Kb. Ka is probably K₂SO₄, 3KHSO₄ and Kb is probably K₂SO₄, 6KHSO₄.

(D'Ans, Z. anorg. 1909, 63. 228.)

3.1 mols. K₂SO₄ are sol. in absolute H₂SO₄ at 25°. (Bergius, Z. phys. Ch. 1910, 73. 353.)

Solubility in H₂SO₄+Aq at 25°.

Millimols H ₂ SO ₄ in 10 cm.	Millimols K ₂ SO ₄ in 10 cm.
...	6.17
3.97	8.92
7.57	10.82
14.35	14.86

(Herz, Z. anorg. 1912, 73. 276.)

Solubility in H₂SO₄+Aq at 25°.

In 1000 g. of the solution		Solid phase
Mols SO ₃	Mols K ₂ SO ₄	
6.42	0.171	KHSO ₄
6.60	0.190	"
6.91	0.266	KHSO ₄ +KH ₂ (SO ₄) ₂ , H ₂ O
7.26	0.182	"
7.62	0.157	"
7.88	0.167	"
8.00	0.201	"
8.10	0.250	KH ₂ (SO ₄) ₂ , H ₂ O
8.15	0.352	"
8.16	0.364	KH ₂ (SO ₄) ₂ , H ₂ O+KH ₂ (SO ₄) ₂
8.29	0.341	"
8.33	0.322	"
8.45	0.325	"
8.62	0.346	"
8.57	0.384	KH ₂ (SO ₄) ₂
8.71	0.412	"
8.82	0.583	KH ₂ (SO ₄) ₂
8.65	0.880	KH ₂ (SO ₄) ₂ +KHS ₂ O ₇
8.63	0.899	KHS ₂ O ₇ (metastable solution)
8.70	0.882	"
8.96	0.561	"
9.80	0.365	"
9.78	0.430	"
9.80	0.665	KHS ₂ O ₇
9.66	0.904	"
9.66	0.937	"

(D'Ans, Z. anorg. 1913, 80. 239.)

Pptd. from K₂SO₄+Aq by NH₄OH+Aq. (Sullivan.)

Solubility of K_2SO_4 in $NH_4OH + Aq$ at 20° .

G. NH_3 in 100 ccm. H_2O	G. K_2SO_4 in 100 ccm. H_2O
0	10.804
6.08	4.100
15.37	0.828
24.69	0.140
31.02	0.042

(Girard, Bull. Soc. (2) 43. 522.)

1 l. sat. solution in H_2O contains 105.7 g. K_2SO_4 at 20° ; in $NH_4OH + Aq$ (5.2% NH_3), 45.2 g. (Konowalow, J. Russ. Phys. Chem. Soc. 1894, 31. 985.)

Solubility of K_2SO_4 in $KOH + Aq$ at 25° .

In 1000 g. of the solution	
Mols K_2SO_4	Mols $(KOH)_2$
0.617	0.0
0.433	0.258
0.280	0.433
0.137	1.13
0.035	2.86
0.009	3.42
0.0	4.809

(D'Ans and Schreiner, Z. anorg. 1910, 67. 438.)

Sol. in sat. $NH_4Cl + Aq$ without pptn. (See NH_4Cl .)

Sl. sol. in sat. $KCl + Aq$ without pptn. 100 g. sat $KCl + Aq$. at 25° dissolve 0.0167 g. equiv. K_2SO_4 at 25° . (Van't Hoff and Meyerhoffer Z. phys. Ch. 1904. 49. 315.)

Sl. sol. in sat. $KNO_3 + Aq$ without causing pptn.

1 l. of the solution contains 50.7 g. $K_2SO_4 + 216.5$ g. $KNO_3 = 267.2$ g. mixed salts at 15° . Sp. gr. $K_2SO_4 + KNO_3 + Aq = 1.165$.

1 l. of the solution contains 47.66 g. $K_2SO_4 + 308.5$ g. $KNO_3 = 356.2$ g. mixed salts at 25° . Sp. gr. of $K_2SO_4 + KNO_3 + Aq = 1.210$. (Euler, Z. phys. Ch. 1904, 49. 313.)

See also under KNO_3 .

Sol. in sat. $NaNO_3 + Aq$ without causing pptn at first, but soon KNO_3 is pptd. (Karsten.) (See $NaNO_3$.)

Sol. in $(NH_4)_2SO_4 + Aq$ with pptn. of $(NH_4)_2SO_4$. (Rüdorff, B. 6. 485.) (See $(NH_4)_2SO_4$.)

More sol. in aqueous solutions of other salts, as Na_2SO_4 , $MgSO_4$, $CuSO_4$, etc., than in pure H_2O . (Pfaff, A. 99. 227.)

Sol. in sat. $Na_2SO_4 + Aq$, $MgSO_4 + Aq$, $NaCl + Aq$. (See $MgSO_4$ and $NaCl$.)

Solubility of K_2SO_4 in $Na_2SO_4 + Aq$.

Temp. $= 34^\circ$		Temp. $= 60^\circ$	
% Na_2SO_4	% K_2SO_4	% Na_2SO_4	% K_2SO_4
0	11.9	0	15.3
7.1	10.7	6.6	13.9
31.4	4.3	27.1	8.2
33.1	0	31.3	0

(Nacken, B. A. B. 1910. 1016.)

100 g. H_2O sat. with both K_2SO_4 and Th_2SO_4 dissolve:

4.74 g. $Th_2SO_4 + 10.3$ g. K_2SO_4 at 15° .
11.5 g. " + 16.4 g. " " 62° .
18.52 g. " + 26.2 g. " " 100° .

(Rabe, Z. anorg. 1902, 31. 156.)

Sl. sol. in sat. $ZnSO_4$ or $CuSO_4 + Aq$ with separation of double salt.

100 pts. H_2O dissolve $8.5 + 0.12t$ pts. K_2SO_4 . On addition of a K salt, K_2SO_4 is pptd. The amount of K_2SO_4 remaining in solution plus the amt. of K in the salt added is a constant. (Blarez, C. R. 112. 939.)

Solubility of $K_2SO_4 + Th(SO_4)_2$ at 16° .

Solid phase, $Th(SO_4)_2$.

Pts. per 100 pts H_2O		Pts. per 100 pts H_2O	
K_2SO_4	$Th(SO_4)_2$	K_2SO_4	$Th(SO_4)_2$
0.000	1.390	1.487	0.870
0.424	1.667	1.633	0.635
1.004	2.193	1.844	0.370
1.152	3.191	2.512	0.128
1.224	2.514	3.092	0.070
1.283	2.222	4.050	0.027
1.348	1.706	4.825	0.003
1.378	1.637

(Barre, C. R. 1911, 150. 1555.)

Difficultly sol. in 20% $KC_2H_3O_2 + Aq$. (Stromeyer.)

Solubility in K acetate + Aq at 25° .

Composition of the solutions		
% K acetate	% K_2SO_4	% H_2O
6.11	6.65	87.24
8.68	5.09	86.23
11.29	3.99	84.72
15.59	2.35	82.06
20.12	1.23	78.65
29.95	0.39	69.66

The solid phase in these solutions is K_2SO_4 .

(Fox, Chem. Soc. 1909, 96. 885.)

100 g. hydroxylamine dissolve 3.5 g. K_2SO_4 at $17-18^\circ$. (de Bruyn, Z. phys. Ch. 1892, 10. 782.)

l. in liquid HF. (Franklin, Z. 46. 2)
liquid NH₃. (Franklin, Am. 20. 829.)
dissolved nor attacked by liquid
nkland, Chem. Soc. 1901, 79.

absolute alcohol.
alcohol, the sp. gr. of which is
thon.)
in dil. alcohol increases with the

alcohol of 0.939 sp. gr. (53% by
y weight) dissolve at:
8° 60°
0.21 0.92 pts. K₂SO₄.
rardin, A. ch. (4) 5. 147.)
f the sat. solution at 15° in alcohol

20 30 40% by weight,
1.46 0.56 0.21 pts. K₂SO₄.
(Schiff, A. 118. 362.)

ity in acetic acid + Aq at 25°.

tion of the Mol./Liter	Solubility of K ₂ SO ₄ Mol./Liter
100	0.6714
170	0.6619
137	0.6559
328	0.6350
578	0.6097
151	0.5556
183	0.4743

d and Wilsmore, Z. phys. Ch. 1902,
40. 620.)

bility in phenol + Aq at 25°.

tion of the Mol./Liter	Solubility of K ₂ SO ₄ Mol./Liter
	0.6714
	0.6650
	0.6614
	0.6598
	0.6555
	0.6522
	0.6502
	0.6494
	0.6310
	0.6251
	0.6042
	0.5956
	0.5834
	0.5572
	0.5480
	0.5425
	0.5389
(saturated)	0.5377

d and Wilsmore, Z. phys. Ch. 1902,
40. 619.)

95% formic acid dissolve 36.5 g.
21°. (Aschan, Ch. Ztg. 1913, 37.

Solubility in organic substances + Aq at 25°.

Organic substance	Composition of the solutions		
	% organic substance	% K ₂ SO ₄	% H ₂ O
Alcohol	1.35	9.17	89.48
	4.80	6.90	88.30
	7.80	4.96	87.24
	9.70	4.32	85.98
	12.34	3.57	84.09
	14.51	2.71	82.78
	15.26	2.66	82.08
	20.50	1.83	77.67
	26.91	0.97	72.12
	35.97	0.41	63.62
	43.90	0.22	55.88
	69.26	0.016	30.72
Pyridine	4.23	7.95	87.82
	13.90	4.77	81.33
	24.51	2.75	72.74
	34.19	1.47	64.34
	46.29	0.45	53.26
	55.93	0.12	43.95
	75.90	0.006	24.09
Ethylene glycol	3.16	9.67	87.17
	9.89	7.69	82.53
	18.47	5.74	75.79
	32.11	3.57	64.32
	49.03	1.83	49.14
Chloral hydrate	6.44	9.13	84.43
	9.09	8.41	82.50
	12.38	7.79	79.83
	13.20	7.31	79.49
	22.07	5.88	72.05
	33.15	4.54	62.31
	44.40	3.36	52.24
	47.30	2.92	49.78
	62.82	2.00	35.18
	70.28	1.75	27.97
	80.36	1.40	18.24
Glycerol	85.26	1.08	13.66
	8.96	8.87	82.17
	13.36	7.69	78.95
	20.34	6.47	73.19
	24.15	5.83	70.02
	33.73	4.44	61.83
	40.40	3.65	55.95
	43.52	3.38	53.10
	50.18	2.69	47.13
	57.22	2.07	40.71
	67.94	1.53	30.53
	78.18	0.98	20.84
	98.28	0.73	0.99
Mannitol	3.20	10.32	86.48
	5.82	10.07	84.11
	8.35	9.61	82.04
	11.26	9.19	79.55
	14.30	8.66	77.04
	17.22	8.35	74.43

Solubility in organic substances + Aq at 25°.—
Continued.

Organic substance	Composition of the solutions		
	% organic substance	% K ₂ SO ₄	% H ₂ O
Sucrose	9.56	9.65	80.79
	18.55	8.65	72.80
	28.16	7.42	64.42
	37.24	6.35	56.41
	47.55	5.21	47.24
	57.00	4.24	38.76
Acetone	4.92	7.20	87.88
	10.06	5.02	84.92
	16.23	2.96	80.81
	24.31	1.50	74.19
	37.19	0.47	62.34
	46.29	0.20	53.51
	62.40	0.03	37.57

(Fox and Gage, Chem. Soc. 1910, **97**. 381.)

Sol. in 76 pts. glycerine of 1.225 sp. gr. at ordinary temp. (Vogel, N. Repert. **16**. 557.)

Insol. in acetone. (Krug and M'Elroy; Eidman, C. C. **1899**, II. 1014.)

Insol. in CS₂. (Arctowski, Z. anorg. **1894**, **6**. 257); benzonitrile. (Naumann, B. **1914**, **47**. 1370); methyl acetate. (Naumann, B. **1909**, **42**, 3790); ethyl acetate. (Naumann, B. **1904**, **37**, 3602.)

100 g. H₂O dissolve 104 g. K₂SO₄ + 219.0 g. sugar at 31.25°, or 100 g. sat. solution contain 3.8 g. K₂SO₄ + 66.74 g. sugar. (Köhler, Z. Ver. Zuckerind. **1897**, **47**. 447.)

Min. *Glaserite*.

+½H₂O. 100 pts. H₂O dissolve 9.82 pts. (Ogier, C. R. **82**. 1055.)

Tripotassium hydrogen sulphate, K₃H(SO₄)₃.
Sol. in H₂O.

Potassium hydrogen sulphate, KHSO₄.

1.07 pts. KHSO₄ (= 1 pt. K₂S₂O₇) dissolve:

at 0° in 2.95 pts. H₂O.

" 20° " 2.08 "

" 40° " 1.59 "

" 100° " 0.88 "

(Kremers, Pogg. **92**. 497.)

Sp. gr. of KHSO₄ + Aq at 15° containing:

5 10 15 % KHSO₄
1.0354 1.0726 1.1116

20 25 27% KHSO₄
1.1516 1.1920 1.2110

(Kohlrausch, W. Ann. **1879**. 1.)

Sat. solution boils at 105.5° (Griffiths); 108° (Kremers).

Alcohol dissolves out H₂SO₄.

K₂SO₄ crystallises from dilute solutions.

100 g. 95% formic acid dissolve 14.6 g. KHSO₄ at 19.3°. (Aschan, Ch. Ztg. **1913**, **37**. 1117.)

Insol. in methyl acetate. (Naumann B. **1909**, **42**, 3790.)

Min. *Misinit*.

+5½H₂O. Deliquescent. (Senderens, Bull. Soc. (3) **2**. 278.)

Potassium dihydrogen sulphate, K₂H₂(SO₄)₂.

Sol. in H₂O. (Phillips, Phil. Mag. **1**. 429.)

Composition is 4K₂O, 7SO₃ + 3H₂O, according to Berthelot (A. ch. (4) **30**. 442).

Potassium trihydrogen sulphate, KH₃(SO₄)₃.

Sol. in H₂O with rise of temperature (Schultz, Pogg. **133**. 137.)

+1½H₂O. (Lescœur, C. R. **78**. 1044.)

Potassium disulphate (pyrosulphate), K₂S₂O₇.

When dissolved in exactly the necessary amount of hot H₂O for solution, it crystallises on cooling without decomp. Decomp. by excess of H₂O. (Jacquelin, A. ch. **70**. 311.)

Insol. in methyl acetate. (Naumann, B. **1909**. **42**. 3790.)

Potassium hydrogen disulphate, KHS₂O₇.

Sol. in fuming H₂SO₄ without decomposition.

Potassium octosulphate, K₂S₈O₂₄.

Decomp. by H₂O. (Weber.)

Potassium praseodymium sulphate, 3K₂SO₄.

Pr₂(SO₄)₃ + H₂O.

Sl. sol. in H₂O.

Sol. in conc. HCl and HNO₃. (Von Schreck Z. anorg. **1898**, **18**. 358.)

Potassium rhodium sulphate, 3K₂SO₄.

Rh₂(SO₄)₃.

Does not exist. (Leidié, C. R. **107**. 234.)

K₂SO₄, Rh₂(SO₄)₃ + 24H₂O. Very sol. in H₂O. (Piccini, Z. anorg. **1901**, **27**. 66.)

Potassium samarium sulphate, 9K₂SO₄.

2Sm₂(SO₄)₃ + 3H₂O.

Sl. sol. in H₂O.

Sl. sol. in sat. K₂SO₄ + Aq.

1 l. sat. K₂SO₄ + Aq dissolves 0.5 g. Sm₂O₃. (Cleve, Bull. Soc. (2) **43**. 166.)

Potassium scandium sulphate, 3K₂SO₄.

Sc₂(SO₄)₃.

Very slowly sol. in cold, more easily sol. in warm H₂O. Insol. in sat. K₂SO₄ + Aq.

Sol. in H₂O and in dil. K₂SO₄ + Aq. (Meyer, Z. anorg. **1914**, **86**. 279.)

2K₂SO₄, Sc₂(SO₄)₃. Sol. in K₂SO₄ + Aq. (Cleve.)

Does not exist. (Nilson.)

Potassium sodium sulphate, 3K₂SO₄, Na₂SO₄.

100 pts. H₂O dissolve 40.8 pts. at 103.5°. (Penny, Phil. Mag. (4) **10**. 401.)

5K₂SO₄, Na₂SO₄. 100 pts. H₂O at 100° dissolve 25 pts.; at 12.7°, 10.1 pts.; at 4.4°, 9.2 pts. (Gladstone, Chem. Soc. **6**. 111.)

Potassium strontium sulphate, $K_2Sr_2(SO_4)_2$.

Decomp. by $(NH_4)_2CO_3 + Aq.$ (Rose. ogg. 93. 604.)

$K_2SO_4, SrSO_4$. This is the only double salt formed from these two components between 0° and 100° .

100 pts of the sat. solution in equilibrium with $SrSO_4$ and the double salt contain at:

17.5°	50°	75°	100°
1.27	1.88	2.71	3.9 pts. K_2SO_4

(Barre, C. R. 1909, 149. 292.)

Potassium tellurium sulphate, $KHSO_4, 2TeO_2, SO_3 + 2H_2O$.

(Metzner, A. ch. 1898, (7) 15. 203.)

Potassium terbium sulphate.

Easily sol. in H_2O . Sl. sol. in $K_2SO_4 + Aq.$ Delafontaine, Zeit. Chem. (2) 2. 230.)

Potassium thallic sulphate, $KTI(SO_4)_2 + 4H_2O$.

Decomp. by H_2O . (Fortini, Gazz. ch. 1905, 35. (2) 453.)

$2K_2O, TI_2O_3, 4SO_3$. Insol. in H_2O . Very difficultly sol. in warm dil. $H_2SO_4 + Aq.$ Strecker, A. 135. 207.)

Potassium thorium sulphate, $K_2SO_4, Th(SO_4)_2 + 4H_2O$.

Sol. in hot H_2O containing a few drops HCl . (Barre, A. ch. 1911, (8) 24. 227.)

$2K_2SO_4, Th(SO_4)_2 + 2H_2O$. Slowly sol. in cold, easily and abundantly in hot H_2O , and is gradually decomp. by boiling. Easily sol. in acids. Insol. in alcohol. (Berzelius.)

$3.5K_2SO_4, Th(SO_4)_2$. Insol. in $K_2SO_4 + Aq$ of concentrations above 4.5%. (Barre.)

$4K_2SO_4, Th(SO_4)_2 + 2H_2O$. (Chydenius.)

Potassium tin (stannous) sulphate, $K_2SO_4, SnSO_4$.

(Marignac.)

Potassium tin (stannic) sulphate, $K_2Sn(SO_4)_2$.

Easily sol. in H_2O with decomp.

Sol. in HCl . (Weinland, Z. anorg. 1907, 14. 250.)

Potassium tin (stannous) sulphate chloride, $4K_2SO_4, 4SnSO_4, SnCl_2$.

Can be recrystallised from H_2O . (Marignac, Ann. Min. (5) 12. 62.)

Potassium titanium sulphate, $K_2SO_4, Ti(SO_4)_2 + 3H_2O$.

Difficultly sol. in H_2O or $HCl + Aq.$ Decomp. by much H_2O . (Wallace, Pogg. 102. 153.)

Potassium titanyl sulphate, $2K_2SO_4, 3TiO, SO_3 + 10H_2O$.

Very sol. in H_2O with decomp. Insol. in conc. H_2SO_4 . (Rosenheim, Z. anorg. 1901, 26. 251.)

$K_2SO_4, TiO, SO_3 + 7H_2O$. Very hygroscopic and sol. in H_2O . (Massuchelli and Pontanelli, C. C. 1909, II. 420.)

$K_2SO_4, (TiO)SO_4$. (Spence, C. C. 1901, II. 747.)

Potassium uranous sulphate, $K_2SO_4, U(SO_4)_2 + H_2O$.

Very sl. sol. in H_2O . (Rammelsberg.)

Potassium uranyl sulphate, $K_2SO_4, (UO_2)SO_4 + 2H_2O$.

Sol. in 9 pts. H_2O at 22° and in 0.51 pt. at 100° . Insol. in alcohol. (Ebelmen, A. ch. (3) 5. 211.)

100 pts. of aqueous solution sat. at 25° contain 10.5 pts. salt; at 70.5° contain 23.93 pts. salt. (Rimbach, B. 1904, 37. 478.)

$+ 3H_2O$. (de Coninck, Chem. Soc. 1905, 88. (2) 394.)

$2K_2SO_4, (UO_2)SO_4 + 2H_2O$. Decomp. by H_2O .

Cannot be cryst. from rather conc. H_2SO_4 , as it is completely decomp. by it. (Rimbach, B. 1905, 38. 1572.)

$K_2O, 2UO_2, 3SO_3$. Ppt. Identical with UO_2, OK, SO_3H of Scheller, (A. 1867, 144. 238.) (Kohlschütter, A., 1900, 311. 11.)

$2K_2SO_4, 3(UO_2)SO_4 + H_2O$. Sol. in H_2O . Insol. in alcohol. (Berzelius.)

Does not exist. (Ebelmen.)

Potassium vanadium sulphate, $K_2O, V_2O_5, 2SO_3 + 6H_2O = K(VO_2)SO_4 + 3H_2O$.

(Friedheim, B. 24. 1183.)

$= KVO_2, K_2SO_4, V_2O_5, 2SO_3 + 9H_2O$ of Münzing (Berlin, Dissert. 1889).

$K_2SO_4, VSO_4 + 6H_2O$. Sol. in H_2O . (Piccini, Z. anorg. 1902, 32. 61.)

$K_2V_2(SO_4)_4 + 24H_2O$. 100 pts. H_2O dissolve 198.4 pts. salt. at 10° . Sp. gr. of sat. solution at $4^\circ/20^\circ = 1.782$. (Piccini, Z. anorg. 1897, 13. 446.)

Potassium vanadyl sulphate,

$K_2SO_4, (VO)_2(SO_4)_2$.

Very slowly sol. in H_2O , still less sol. in dil. alcohol. (Gerland.)

$K_2SO_4, VOSO_4 + 3H_2O$. Easily sol. in H_2O . Sol. in alcohol + conc. H_2SO_4 . (Koppel, Z. anorg. 1903, 35. 178.)

$K_2SO_4, 2VOSO_4$. Very hygroscopic. Very sol. in H_2O but goes into solution slowly. (Koppel and Behrendt, B. 1901, 34. 3935.)

Easily sol. in H_2O . (Koppel, Z. anorg. 1903, 35. 174.)

Potassium yttrium sulphate, $4K_2SO_4, Y_2(SO_4)_3$.

Sol. in 16 pts. cold H_2O , and in 10 pts. sat. $K_2SO_4 + Aq$, and more abundantly if the latter solution contains ammonium salts or free acid. (Berlin.)

$3K_2SO_4, 2Y_2(SO_4)_3$. 100 ccm. cold sat.

$K_2SO_4 + Aq$ dissolve an amount of this salt corresponding to 4.685 g. Y_2O_3 . (Cleve.)

Potassium zinc sulphate, $K_2SO_4, ZnSO_4 + 6H_2O$.

Sol. in 5 pts. cold H_2O . (Bucholz, N. J. Pharm. 9. 2. 26.)

100 pts. H_2O dissolve at:

0° 10° 15° 25° 36°
12.6 18.7 22.5 28.8 39.9 pts. hydrous salt,

45° 50° 58° 65° 70°
51.2 54.0 67.6 81.3 87.9 pts. hydrous salt.
(Tobler, A. 95. 193.)

100 pts. H_2O at 15° dissolve 14.8 pts. $K_2SO_4, ZnSO_4 + 6H_2O$; sp. gr. of sat. H_2O solution at 15° = 1.0939. (Schiff, A. 109. 326.)

1 l. H_2O dissolves 131.9 g. anhydrous salt at 25°. (Locke, Am. Ch. J. 1902, 27. 459.)

Potassium zirconium sulphate, $2K_2O, 6ZrO_2, 7SO_3 + 9H_2O$.

Decomp. by H_2O .

$3K_2O, 3ZrO_2, 7SO_3 + 9H_2O$. Insol. in H_2O .
 $Zr_2O_3(KSO_4)_2 + 8H_2O$. Ppt. (Rosenheim, B. 1905, 38. 815.)

Potassium sulphate vanadate.

Very difficultly sol. in H_2O . Insol. in alcohol. (Berzelius.)

Potassium sulphate antimony trifluoride.

See Antimony trifluoride potassium sulphate.

Praseodymium sulphate, basic, $(PrO)_2SO_4$.

Insol. in H_2O . (Matignon, C. R. 1902, 134. 660.)

Insol. in H_2O . Nearly insol. in dil. acids. (Wöhler, B. 1913, 46. 1730.)

Praseodymium sulphate, $Pr_2(SO_4)_3$.

Sol. in H_2O ; very hygroscopic. 23.64 pts. are sol. in 100 pts. H_2O at 0° and 17.7 pts. at 20°. (von Scheele, Z. anorg. 1898, 18. 357-358.)

+5 H_2O . Sol. in H_2O . (von Scheele, Z. anorg. 1898, 18. 357.)

Difficultly sol. in H_2O . (Kraus, Zeit. Kryst. 1901, 34. 400.)

1.50 pts. $Pr_2(SO_4)_3$ are sol. in 100 pts. H_2O at 85°; 1.45 pts. at 90°; and 1.02 pts. at 95°. (Muthmann and Rölig, B. 1898, 31. 1729.)

+8 H_2O . (Kraus, Zeit. Kryst. 1901, 34. 406.)

Sol. in H_2O . (von Scheele, Z. anorg. 1898, 18. 357.)

Solubility in H_2O at t°.

t°	Pts. $Pr_2(SO_4)_3$
0	19.79
18	14.10
35	10.31
55	7.09
75	4.13

(Muthmann and Rölig, B. 1898, 31. 1727.)

+15½ H_2O . Sol. in H_2O . (von Scheele, Z. anorg. 1898, 18. 357.)

Praseodymium hydrogen sulphate, $Pr(SO_3H)_3$.

(Brauner, Z. anorg. 1904, 38. 330.)

Solubility in boiling conc. H_2SO_4 . 100 g. of the solution contain 1.02 g. of the acid sulphate. (Matignon, C. R. 1902, 134. 659.)

Radium sulphate.

Less sol. in H_2O than corresponding Ba comp. (Curie, Dissert. 1903.)

Rhodium sulphate, $Rh_2(SO_4)_3 + 12H_2O$.

Easily sol. in H_2O . (Berzelius.)

Sl. sol. in, but not decomp. by H_2O when not more than 16 pts. H_2O are present to 1 pt. salt. Decomp. by hot H_2O to—

$Rh_2(SO_4)_3, Rh_2O_3$. Insol. in H_2O . (Leidié, C. R. 107. 234.)

Rhodium rubidium sulphate, $Rh_2(SO_4)_3, Rb_2SO_4 + 24H_2O$.

Sol. in H_2O ; m.-pt., 108-109°. (Piccini, Z. anorg. 1901, 37. 65.)

Rhodium thallium sulphate, $Rh_2(SO_4)_3, Tl_2SO_4 + 24H_2O$.

Very sol. H_2O . (Piccini, Z. anorg. 1901, 37. 69.)

Rhodium sodium sulphate, $Rh_2Na_2(SO_4)_4$.

Insol. in H_2SO_4 or aqua regia. (Seubert and Kobbe, B. 23. 2560.)

Rubidium sulphate, Rb_2SO_4 .

100 pts. H_2O dissolve 42.4 pts. at 10°. (Bunsen.)

100 cc. H_2O at 17-18° dissolve 44.7 g. Rb_2SO_4 . (Tutton, Chem. Soc. 1894, 65. 632.)

Sat. $Rb_2SO_4 + Aq$ contains at:

3°	20°
27.4	32.5% Rb_2SO_4
37°	97°
37.3	43.9
	49.2% Rb_2SO_4
	(Étard, A. ch. 1894, (7) 2. 650.)

Solubility of Rb_2SO_4 in H_2O at t° .

t°	g. Rb_2SO_4 per 100 g.		t°	g. Rb_2SO_4 per 100 g.	
	H_2O	solution		H_2O	solution
0	36.4	27.3	60	67.4	40.3
10	42.6	29.9	70	71.4	41.7
20	48.2	32.5	80	75.0	42.9
30	53.5	34.9	90	78.7	44.0
40	58.5	36.9	100	81.8	45.0
50	63.1	38.7	102.4*	82.6	45.2

* B.-pt. at 742.4 mm.

(Berkeley, calc. by Seidell, Solubilities, 2nd Ed., p. 587.)

Sp. gr. of $\text{Rb}_2\text{SO}_4 + \text{Aq}$ sat. at $10^\circ = 1.2978$.
(Erdmann, Arch. Pharm. 1894, 232. 16.)G.-equiv. salt per l. at $18^\circ = 0.501$ 1.01Sp. gr. $6^\circ/6^\circ$ 1.05587 1.11047" $18^\circ/18^\circ$ 1.05496 1.10896" $30^\circ/30^\circ$ 1.05433 1.10810G.-equiv. salt per l. at $18^\circ = 2.043$ 3.168Sp. gr. $6^\circ/6^\circ$ 1.21888 1.33276" $18^\circ/18^\circ$ 1.21613 1.32912" $30^\circ/30^\circ$ 1.21443 1.32750

(Clausen, W. Ann. 1914, (4) 44. 1071.)

10 ccm. of sat. $\text{Rb}_2\text{SO}_4 + \text{absolute } \text{H}_2\text{SO}_4$ contain approx. 5.881 g. Rb_2SO_4 . (Bergius, Z. phys. Ch. 1910, 72. 355.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); acetone (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Rubidium pyrosulphate, $\text{Rb}_2\text{S}_2\text{O}_7$.Decomp. by H_2O .**Rubidium octosulphate, $\text{Rb}_8\text{S}_8\text{O}_{32}$.**Decomp. by H_2O . (Weber, B. 17. 2497.)**Rubidium hydrogen sulphate, RbHSO_4 .**Sol. in H_2O .**Rubidium tin (stannic) sulphate, $\text{Rb}_2\text{Sn}(\text{SO}_4)_2$.**Decomp. by H_2O . Sol. in HCl . (Weinland, Z. anorg. 1907, 54. 250.)**Rubidium thallic sulphate, $\text{RbTl}(\text{SO}_4)_2$.**

(Marshall, C. C. 1902, II. 1089.)

+ $4\text{H}_2\text{O}$. (Fortini, Gazz. ch. it. 1905, 35. (2) 455.)**Rubidium thorium sulphate, $\text{Rb}_2\text{SO}_4, \text{Th}(\text{S}_4\text{O}_{10})_2 + 2\text{H}_2\text{O}$.**Sl. sol. in H_2O . (Manuelli, Gazz. ch. it. 1903, 32. (2) 523.)**Rubidium titanium sulphate, $\text{Rb}_2\text{SO}_4, \text{Ti}_2(\text{SO}_4)_3 + 24\text{H}_2\text{O}$.**Sol. in H_2O acidified with H_2SO_4 . Decomp. in neutral aq. solution. (Piccini, Z. anorg. 1898, 17. 359.)Insol. in H_2O . Sol. in HCl . Insol. in H_2SO_4 . Decomp. by boiling with conc. H_2SO_4 . (Stähler, B. 1905, 38. 2623.)**Rubidium uranyl sulphate, $\text{Rb}_2(\text{UO}_2)(\text{SO}_4)_2 + 2\text{H}_2\text{O}$.**Somewhat less sol. in H_2O than K salt. (Rimbach, B. 1904, 37. 479.)**Rubidium vanadium sulphate, $\text{Rb}_2\text{V}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.**0.177 gram mols. of anhydrous salt are sol. in 1 l. H_2O . (Locke, Am. Ch. J. 1901, 26. 175.)Insol. in H_2O .Insol. in H_2SO_4 . Decomp. by boiling with conc. H_2SO_4 .Sol. in HCl . (Stähler, B. 1905, 38. 3980.)100 pts. H_2O dissolve 2.56 pts. salt at 10° .Sp. gr. of solution at $4^\circ/20^\circ = 1.915$. (Piccini, Z. anorg. 1897, 13. 446.)**Rubidium zinc sulphate, $\text{Rb}_2\text{SO}_4, \text{ZnSO}_4 + 6\text{H}_2\text{O}$.**Sol. in H_2O . (Bunsen and Kopp, Pogg. 113. 337.)1 l. H_2O dissolves 101 g. anhydrous salt at 25° . (Locke, Am. Ch. J. 1902, 27. 459.)**Rubidium zirconium sulphate, $\text{Zr}_2\text{O}_3, (\text{RbSO}_4)_2 + 15\text{H}_2\text{O}$.**

Ppt. (Rosenheim, B. 1905, 38. 815.)

Ruthenic sulphate, $\text{Ru}(\text{SO}_4)_2$.Deliquescent, and easily sol. in H_2O . (Claus, A. 59. 246.)**Samarium sulphate, basic, $(\text{SmO})_2\text{SO}_4$.**Insol. in H_2O and in cold dil. H_2SO_4 . (Matignon, C. R. 1905, 141. 1231.)**Samarium sulphate, $\text{Sm}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$.**Difficultly sol. in H_2O .Much less sol. than $\text{Di}_2(\text{SO}_4)_3 + 8\text{H}_2\text{O}$. (Cleve.)2.05 pts. anhydrous salt are sol. in 100 pts. H_2O at 25° . (Keyes and James, J. Am. Chem. Soc. 1914, 36. 635.)100 g. $\text{Sm}_2(\text{SO}_4)_3 + \text{Aq}$ sat. at 25° contain 3.426 g. anhyd. $\text{Sm}_2(\text{SO}_4)_3$. (Wirth, Z. anorg. 1912, 76. 174.)Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° . n = equiv. g. of H_2SO_4 in 1 l. of solvent. c = g Sm_2O_3 in 100 g. of solution. c_1 = g. $\text{Sm}_2(\text{SO}_4)_3$ in 100 g. of solution.

n	c	c_1	n	c	c_1
0	2.029	3.426	2.16	1.43	2.416
0.1	2.038	3.441	6.175	0.416	0.7025
0.505	1.985	3.352	12.6	0.0656	0.1107
1.1	1.821	3.075

(Wirth, Z. anorg. 1912, 76. 174.)

Solubility in $(\text{NH}_4)_2\text{SO}_4 + \text{Aq}$ at 25° .

Pts. $\text{Sm}_2(\text{SO}_4)_3$ per 100 pts. H_2O	Pts. $(\text{NH}_4)_2\text{SO}_4$ per 100 pts. H_2O	Solid phase
2.1	0.3	$\text{Sm}_2(\text{SO}_4)_3$
2.0	0.8	
2.8	1.1	
1.5	1.9	
1.2	2.7	
0.8	7.4	
0.8	9.5	
0.8	8.7	
0.8	18.8	
0.8	12.2	
0.8	12.3	$\text{Sm}_2(\text{SO}_4)_3, (\text{NH}_4)_2\text{SO}_4, 7\text{H}_2\text{O}$
0.9	32.5	
1.0	46.3	
0.9	40.3	
1.3	77.5	
0.2	77.2	
0.3	77.3	
0.6	76.8	
		$(\text{NH}_4)_2\text{SO}_4$

(Keyes and James, J. Am. Chem. Soc. 1914, **36**. 637.)

Solubility in $\text{Na}_2\text{SO}_4 + \text{Aq}$ at 25° .

Pts. Na_2SO_4 per 100 pts. H_2O	Pts. $\text{Sm}_2(\text{SO}_4)_3$ per 100 pts. H_2O	Solid phase
0.1	2.0	$\text{Sm}_2(\text{SO}_4)_3$
0.5	0.11	"
1.9	0.03	$2\text{Sm}_2(\text{SO}_4)_3, 3\text{Na}_2\text{SO}_4 + 6\text{H}_2\text{O}$
6.44	0.016	"
7.00	0.008	"
9.02	0.016	"
10.51	0.012	"
11.48	0.012	"
13.58	0.010	"
14.71	0.010	"
14.47	0.009	"
20.02	0.012	"
23.42	0.012	"
23.68	0.018	"
25.93	0.015	"
27.40	0.011	"

These results seem to indicate that there is only one double salt formed by the union of $\text{Sm}_2(\text{SO}_4)_3$ with Na_2SO_4 . Formula of this salt is $2\text{Sm}_2(\text{SO}_4)_3, 3\text{Na}_2\text{SO}_4, 6\text{H}_2\text{O}$.

(Keyes and James, J. Am. Chem. Soc. 1914, **36**. 635.)

Samarium hydrogen sulphate, $\text{Sm}(\text{HSO}_4)_3$.

Sl. sol. in H_2O . (Matignon, C. R. 1905, **141**. 1230.)

Ppt. (Brauner, Z. anorg. 1904, **33**. 331.)

Samarium sodium sulphate, $\text{Sm}_2(\text{SO}_4)_3, \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$.

Sl. sol. in sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$. (Cleve, Bull. Soc. (2) **43**. 166.)

$2\text{Sm}_2(\text{SO}_4)_3, 3\text{Na}_2\text{SO}_4 + 6\text{H}_2\text{O}$. Only double salt formed at 25° . (Keyes and James, J. Am. Chem. Soc. 1914, **36**. 365.)

Scandium sulphate, basic, $\text{Sc}_2\text{O}(\text{SO}_4)_2$.

(Crookes, Roy. Soc. Proc. 1908, **80**. A, 515.)

Scandium sulphate, $\text{Sc}_2(\text{SO}_4)_3$.

Anhydrous. Easily sol. in H_2O . $+2\text{H}_2\text{O}$.

$+5\text{H}_2\text{O}$; 54.61 g. of pentahydrate are sol in 100 cc. H_2O at 25° . (Wirth, Z. anorg. 1914, **87**. 10.)

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

$\text{H}_2\text{SO}_4 + \text{Aq}$.	g. $\text{Sc}_2(\text{SO}_4)_3$ in 100 g. of the solution
0.00	28.52
0.5-n	29.29
1.0-n	19.87
4.86-n	8.363
9.73-n	1.315

In 22.35-n H_2SO_4 the solid phase is $\text{Sc}_2(\text{SO}_4)_3, 3\text{H}_2\text{SO}_4$ and 100 g. sat. solution contain 0.484 g. $\text{Sc}_2(\text{SO}_4)_3$.

(Wirth, Z. anorg. 1914, **87**. 10.)

$+6\text{H}_2\text{O}$. Extremely sol. in H_2O , but not deliquescent.

Scandium hydrogen sulphate, $\text{Sc}_2(\text{SO}_4)_3 + 3\text{H}_2\text{O}$.

(Wirth.)

Scandium sodium sulphate, $\text{Sc}_2(\text{SO}_4)_3, 3\text{Na}_2\text{SO}_4 + 12\text{H}_2\text{O}$.

Sol. in H_2O . (Cleve.)

$+10\text{H}_2\text{O}$. Sol. in H_2O and in excess of $\text{Na}_2\text{SO}_4 + \text{Aq}$. (Meyers, Z. anorg. 1914, **88**. 279.)

Silver (argentoargentic) sulphate, $\text{Ag}_2\text{SO}_4, \text{Ag}_2\text{SO}_4 + \text{H}_2\text{O}$.

Gradually sol. in conc., but not attacked by dil. $\text{HNO}_3 + \text{Aq}$. Not attacked by hot conc. H_2SO_4 . (Lea, Sill. Am. J. **144**. 322.)

Silver sulphate, Ag_2SO_4 .

Sol. in 200 pts. cold, and less than 100 pts. boiling H_2O . (Wittstein.)

Sol. in 88 pts. boiling H_2O (Schnaubart); in 87.25 pts. boiling H_2O (Wenzel); in 68.85 pts. H_2O at 100° (Kremers).

100 pts. H_2O at 15.5° dissolve 1.15 pts. Ag_2SO_4 . (Ure's Dict.)

Sol. in 160 pts. H_2O at 18.75° . (Abl.)

1 l. H_2O dissolves 2.57×10^{-2} g.-mol. Ag_2SO_4 at 25° . (Drucker, Z. anorg. 1901, **28**. 362.)

1 l. H_2O dissolves 7.707 g. Ag_2SO_4 at 17° . (Euler, Z. phys. Ch. 1904, **49**. 314.)

1 l. H_2O dissolves 0.0267 mol. Ag_2SO_4 at 25° . (Rothmund, Z. phys. Ch. 1909, **60**. 539.)

1 l. H_2O dissolves 8.35 g. Ag_2SO_4 at 25° .

(Hill and Simmons, Z. phys. Ch. 1909, 67. 603.)

1 l. H_2O dissolves 8.344 g. Ag_2SO_4 at 25° .
Sp. gr. of solution = 1.0052. (Harkins, J. Am. Chem. Soc. 1911, 33. 1812.)

Solubility in H_2O at t° .

t°	Pts. Ag_2SO_4 in 100 pts. of the solution.
14.5	0.730
33	0.909
51.5	1.062
75	1.237
100	1.393

(Barre, A. ch. 1911, (8) 24. 211.)

More sol. in $H_2SO_4 + Aq$ than in pure H_2O .
Still more sol. in $HNO_3 + Aq$ and still more in conc. H_2SO_4 , from which it is pptd. by H_2O . (Schnaubart.)

Solubility in $H_2SO_4 + Aq$ at 25° .

$\frac{1}{2}H_2SO_4 + Aq$. Normality	Solubility of Ag_2SO_4 g.-mol. per litre
0.02	2.60×10^{-2}
0.04	2.64×10^{-2}
0.10	2.71×10^{-2}
0.20	2.75×10^{-2}

(Drucker, Z. anorg. 1901, 28. 362.)

Solubility of Ag_2SO_4 in acids + Aq at 25° .
C = concentration of acid in acid + Aq in milliequivalents per l.

S = Solubility of Ag_2SO_4 in acid + Aq in milliequivalents per l.

Acid	C	S
HNO_3	0.0	53.98
	15.89	59.86
	31.78	65.32
	63.57	75.90
H_2SO_4	0.0	53.98
	29.02	54.88
	58.02	55.64
	105.26	56.82

(Swan, J. Am. Chem. Soc. 1911, 33. 1814.)

Solubility in $HNO_3 + Aq$ at 25° .

Normality HNO_3	Sp. gr. of the solution	g. Ag_2SO_4 dissolved per l.
0.000	1.0054	8.350
1.0046	1.061	34.086
2.0452	1.1069	49.010
4.017	1.1871	71.166
4.209	1.1956	73.212
5.564	1.2456	84.609
8.487	1.3326	94.671
10.034	1.3676	90.806

(Hill and Simmons, Z. phys. Ch. 1909, 67. 603.)

Sol. in NH_4OH , and $(NH_4)_2CO_3 + Aq$.

100 pts. H_2O dissolve 0.58 pt. at 18° . 100 pts. $(NH_4)_2SO_4 + Aq$ (15%) dissolve 0.85 pt. Ag_2SO_4 at 18° . Other sulphates have little effect. (Eder, J. pr. (2) 17. 44.)

Determinations of the solubility of Ag_2SO_4 in $(NH_4)_2SO_4$ at temp. between 16.5° and 100° show that no double salt is formed by these two sulphates. (Étard, A. ch. 1911, (8) 24. 221.)

Solubility of Ag_2SO_4 in $(NH_4)_2SO_4 + Aq$.

G. per 100 g. H_2O .

$(NH_4)_2SO_4$	Ag_2SO_4	$(NH_4)_2SO_4$	Ag_2SO_4
Temp. = 33°		Temp. = 51°	
8.85	1.101	8.90	1.362
15.90	1.331	16.27	1.680
22.22	1.500	22.43	1.887
27.25	1.585	32.10	2.061
30.80	1.619	35.38	2.095
35.88	1.627	39.03	2.082
39.46	1.600	42.37	2.055
43.22	1.557	45.05	2.026
Temp. = 75°		Temp. = 100°	
8.80	1.758	9.23	2.221
15.23	2.155	15.00	2.626
22.30	2.490	22.01	3.075
28.25	2.734	27.00	3.325
32.00	2.823	34.90	3.663
35.82	2.889	38.70	3.772
41.16	2.929	44.15	3.854
46.46	2.902	47.63	3.867

(Barre, A. ch. 1911, (8) 24. 149, 202, 210.)

Solubility of Ag_2SO_4 in $K_2SO_4 + Aq$.

G. per 100 g. H_2O .

K_2SO_4	Ag_2SO_4	K_2SO_4	Ag_2SO_4
Temp. = 33°		Temp. = 51°	
3.22	0.863	3.20	1.023
5.62	0.940	5.61	1.127
8.37	1.046	8.40	1.247
10.41	1.117	10.55	1.340
11.80	1.177	13.16	1.450
...	...	14.37	1.524
Temp. = 75°		Temp. = 100°	
3.12	1.273	3.23	1.488
5.73	1.406	5.60	1.675
8.43	1.554	8.45	1.890
10.55	1.665	11.30	2.115
13.17	1.806	15.07	2.410
17.06	2.021	18.58	2.677

(Barre, A. ch. 1911, (8) 24. 149, 202, 210.)

Solubility in $K_2SO_4 + Aq$ at 25°	
$\frac{1}{2} K_2SO_4 + Aq$ Normality	Solubility of Ag_2SO_4 g.-mol. per litre
0.02	2.46×10^{-3}
0.04	2.36×10^{-3}
0.10	2.31×10^{-3}
0.20	2.32×10^{-3}

(Drucker, Z. anorg. 1901, 23. 362.)

Solubility in $Na_2SO_4 + Aq$ at t° .		
t°	Ag_2SO_4 in 100 pts. H_2O	Na_2SO_4 in 100 pts. H_2O
14.5	0.741	5.278
	0.904	10.103
	1.003	13.045
33	0.972	5.345
	1 150	10.056
	1 320	15 185
	1.448	20.093
	1.548	25.412
	1 570	29.556
	1.549	34.732
	1 462	39.447
	1 199	44.693
	0.932	46.976
51	1.173	5.407
	1.377	10.116
	1.572	15.146
	1.705	20.247
	1.787	25.196
	1.802	29.230
	1.727	34.625
	1.540	39.302
	1.188	42.914
	0 882	44.464
75	1 458	5 368
	1 697	9 813
	1 934	15.260
	2 075	19 978
	2 161	25 556
	2 138	29 662
	1 910	35 278
	1 603	38 944
	1 156	41 365
100	1 651	5.336
	2 012	10 153
	2 312	15 532
	2 351	25 451
	2 260	29 714
	2 012	34 718
	1 687	38 635
	1 158	40 160

Up to 33° , the solubility of Ag_2SO_4 in $Na_2SO_4 + Aq$ increases with the concentration of Na_2SO_4 , above 33° the solubility of Ag_2SO_4 rises to a maximum at a certain concentration of Na_2SO_4 , dependent on the temp. The

solubility curves for various temp. all end at concentration of 40% Na_2SO_4 , that is, the mixed crystals formed at this concentration are equally sol. at all temp.

(Barre, C. R. 1910, 150. 1323.)

Solubility in $Na_2SO_4 + Aq$ at t° .

t°	100 pts. H_2O dissolve	
	Na_2SO_4	Ag_2SO_4
18	0.0	0.766
	0.25	0.712
	0.51	0.682
	0.74	0.675
	1.00	0.665
	1.48	0.670
	2.01	0.673
	2.50	0.689
	3.04	0.703
	4.00	0.736
	4.99	0.768
	10.10	0.932
	13.04	1.028
33	0.0	0.917
	0.25	0.861
	0.51	0.835
	0.75	0.825
	0.98	0.816
	1.50	0.820
	2.01	0.832
	2.48	0.849
	3.00	0.867
51	0.00	1.081
	0.25	1.032
	0.49	1.010
	0.68	0.990
	1.02	0.995
	1.51	1.002
	1.90	1.017
	2.46	1.034
	2.92	1.053
	3.95	1.103
75	0.00	1.267
	0.20	1.215
	0.47	1.206
	0.80	1.206
	1.52	1.210
	1.96	1.222
	2.50	1.238
	2.98	1.269
	4.06	1.296
100	0.00	1.404
	0.50	1.341
	1.01	1.363
	1.44	1.382
	1.94	1.418
	3.02	1.494

(Barre, A. ch. 1911, (8) 26. 215.)

Solubility in salts + Aq at 25°.

C = concentration of salt in salt + Aq in milliequivalents per l.
d₁ = sp. gr. 25°/4° of salt + Aq.
S = solubility of Ag₂SO₄ in salt + Aq expressed in milliequivalents per l.
d₂ = sp. gr. 25°/4° of Ag₂SO₄ + salt + Aq.

Salt	C	d ₁	S	d ₂
none	53.52	...
KNO ₃	24.914	0.9986	57.70	1.0072
	49.774	1.0002	61.13	1.0092
	99.870	1.0034	67.93	1.0034
Mg(NO ₃) ₂	24.764	0.9985	59.44	1.0073
	49.595	0.9999	64.32	1.0094
	99.460	1.0026	72.70	1.0133
AgNO ₃	24.961	1.0007	39.09	1.0065
	49.86	1.0044	28.45	1.0084
	99.61	1.0112	16.96	1.0137
K ₂ SO ₄	25.024	0.9989	50.66	1.0064
	50.044	1.0006	49.35	1.0079
	100.0	1.0041	48.04	1.0112
	200.03	1.0110	48.30	1.0180
MgSO ₄	20.22	0.9984	52.21	1.0061
	50.069	1.0002	50.93	1.0079
	100.04	1.0032	49.95	1.0105
	200.05	1.0092	49.60	1.0164

(Harkins, J. Am. Chem. Soc. 1911, 33. 1813.)

Solubility of Ag₂SO₄ in salts + Aq at 25°.

C = concentration of salt in salt + Aq in milliequivalents per l.
S = solubility of Ag₂SO₄ in salt + Aq in milliequivalents per l.

Salt	C	S
KHSO ₄	0.0	53.98
	52.64	52.18
	105.26	51.76
K ₂ SO ₄	0.0	53.98
	27.18	50.90
	54.34	49.30

(Swan, J. Am. Chem. Soc. 1911, 33. 1814.)

Decomp. by alkali thiosulphates + Aq. (Herschell.)
100 ccm. Ag₂SO₄ + AgC₂H₃O₂ + Aq sat. at 17° contain 3.95 g. Ag₂SO₄ and 8.30 g. AgC₂H₃O₂ and solution has sp. gr. = 1.0094. (Euler, C. C. 1904, I. 1316.)
Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898 20. 829.)

Solubility in organic compds. + Aq at 25°.

Solvent	Mol. Ag ₂ SO ₄ sol. in 1 litre
Water	0.0267
0.5-N Methyl alcohol	0.0249
" Ethyl alcohol	0.0228
" Propyl alcohol	0.0218
" Tert. amyl alcohol	0.0204
" Acetone	0.0220
" Ether	0.0206
" Formaldehyde	0.0227
" Glycol	0.0259
" Glycerine	0.0263
" Mannitol	0.0297
" Glucose	0.0283
" Sucrose	0.0270
" Urea	0.0303
" Dimethylpyrone	0.0216
" Urethane	0.0227
" Formamide	0.0270
" Acetamide	0.0253
" Acetonitrile	0.0525
" Glycocoll	0.0433
" Acetic acid	0.0252
" Phenol	0.0379
" Chloral	0.0233
" Methylal	0.0205
" Methyl acetate	0.0212

(Rothmund, Z. phys. Ch. 1909, 69. 539.)

Insol. in methyl acetate (Bezold, Dissert. 1906; Naumann, B. 1909, 42. 3790); ethylacetate. (Naumann, B. 1904, 37. 3601); liquid methylamine. (Franklin, J. Am. Chem. Soc. 1906, 28. 1420); acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Very sol. in a hot mixture of H₂SO₄ and monobrombenzene, less sol. in cold. (Couper, A. ch. (3) 52. 311.)

Silver hydrogen sulphate, AgHSO₄.

Decomp. by H₂O; sol. in H₂SO₄. (Stas.)
Ag₂O, 3H₂O, 4SO₂ + 2H₂O = AgH₂(SO₄)₂ + H₂O. As above. (Schultz, Pogg. 133. 137.)
2Ag₂O, 3H₂O, 5SO₂ + 2H₂O = Ag₄H₂(SO₄)₃ + 2H₂O. As above. (Schultz.)

Silver pyrosulphate, Ag₂S₂O₇.

Decomp. by H₂O. (Weber, B. 17. 2497.)

Silver thallic sulphate, AgTl(SO₄)₂.

(Lepsius, Chem. Ztg. 1890. 1327.)

Silver tin (stannic) sulphate, Ag₂Sn(SO₄)₂ + 3H₂O.

Ppt. Decomp. by H₂O. Sol. in HCl. (Weinland, Z. anorg. 1907, 54. 250.)

Silver sulphate acetylde, Ag₂SO₄, 2Ag₂C₂.

(Plimpton, Proc. Chem. Soc. 1892, 8. 109.)

Silver sulphate ammonia, $\text{Ag}_2\text{SO}_4 \cdot 2\text{NH}_3$.

Completely sol. in H_2O . (Rose, Pogg. 20. 153.)

$\text{Ag}_2\text{SO}_4 \cdot 4\text{NH}_3$. Easily sol. in H_2O or $\text{NH}_4\text{OH} + \text{Aq}$ without decomp. (Mitscherlich.)

Silver sulphate mercuric oxide, $\text{Ag}_2\text{SO}_4 \cdot \text{HgO}$.

Insol. in H_2O , but decomp. even in the cold. Sol. in HNO_3 and H_2SO_4 . (Finci, Gazz. ch. it. 1911, 41. (2) 548.)

Silver sulphate sulphide, $\text{Ag}_2\text{SO}_4 \cdot \text{Ag}_2\text{S}$.

Decomp. by hot H_2O or cold $\text{HCl} + \text{Aq}$. Sol. in boiling $\text{HNO}_3 + \text{Aq}$. (Poleck and Thümmel, B. 16. 2435.)

Sodium sulphate, Na_2SO_4 .**Anhydrous.**

1 pt. Na_2SO_4 is sol. in 7.367 pts. H_2O at 15° (Gerlach); in 8.52 pts. H_2O at 13.3° (Poggendorf); in 10 pts. H_2O at 13° , and in 3.3 pts. H_2O at 62.2° (Wenzel).

100 pts. H_2O at 0° dissolve 5.155 pts. Na_2SO_4 (Pfaff, A. 99. 226); at 100.6° dissolve 45.985 pts. Na_2SO_4 (Griffiths).

See below for further data.

+7 H_2O . Efflorescent. Insol. in alcohol.

See below for further data.

+10 H_2O .

$\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ is sol. in H_2O with absorption of heat; 20 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ mixed with 100 pts. H_2O at 12.5° lower the temperature 6.8° . (Rüdorff, B. 2. 68.)

Sol. in 2.33 pts. H_2O at 19° , or 100 pts. H_2O at 19° dissolve 42.8 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$. (Schiff, A. 100. 326.)

100 pts. H_2O dissolve a pts. Na_2SO_4 and b pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ at t°

t°	a	b	t°	a	b
0	5.02	12.17	33.88	50.04	312.11
11.67	10.12	26.38	40.15	48.78	291.44
13.30	11.74	31.33	45.04	47.81	276.91
17.91	16.73	48.28	50.40	46.82	262.35
25.05	28.11	99.48	59.79	45.42	
28.76	37.35	161.53	70.61	44.35	
30.75	43.05	215.77	84.42	42.96	
31.84	47.37	270.22	103.17	42.65	
32.73	50.65	322.12			

(Gay-Lussac, A. ch. (2) 11. 312.)

Maximum solubility is at 33° from experiment and theoretical considerations. At this temp. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ is converted into Na_2SO_4 . (Kopp, A. 24. 271.)

100 pts. H_2O at t° dissolve pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$.

t°	Pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$	t°	Pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$	t°	Pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$
2.5	11.39	37.50	294.04	75	241.68
7.5	16.38	43.75	261.04	81.25	217.20
12.5	29.03	50	285.06	87.50	220.65
16.75	70.78	56.25	248.11	93.75	225.46
25	143.38	62.5	222.22	100	241.69
31.25	479.97	68.75	242.88		

(Brandes and Firnhaber, 1884.)

1 pt. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ is sol. in 6.1 pts. H_2O at 7.5° , 3.44 pts. at 12.5° , 2.41 pts. at 18.75° and 1.74 pt. at 20° . (Karsten.)

1 pt. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ is sol. in 2.96 pts. cold, and 0.8 pt. boiling H_2O (Bergmann), in 3 pts. cold, and 0.5 pt. boiling H_2O (Wittstein), in 4 pts. cold, and 1 pt. boiling H_2O (Fourcroy), in 3 pts. H_2O at 15.75° (Abli).

100 pts. H_2O dissolve 12.494 pts. Na_2SO_4 , or 35.492 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ at 15° , and sp. gr. of solution = 1.10847 (Michel and Krafft, A. ch. (3) 41. 478.)

100 pts. H_2O dissolve 39.4 pts. crys. salt at 15.5° , 80 pts. crys. salt at 100° . (Ure's Dict.)

100 pts. H_2O dissolve pts. Na_2SO_4 at t°

t°	Na_2SO_4	t°	Pts. Na_2SO_4
0	4.53	24.1	25.92
17.9	16.28	33	50.81

(Dracon, J. B. 1866. 61.)

Solubility of Na_2SO_4 in H_2O at various pressures and temp. Pts. Na_2SO_4 contained in 100 pts. sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ at A pressure in atmos. and t° are given.

A	t°	15°	15.4°	A	15°
1	4.40	11.32	11.4	100	10.05
20	4.53	10.78	10.74	100	10.33

(Möller, Pogg. 117. 386.)

The solubility of $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ increases with the temperature from 0 to 34° . At 34° and above, it is converted into the anhydrous salt, the solubility of which is least at 103.17° , which is the boiling point of the saturated solution, and increases by cooling from that temp. down to $18-17^\circ$. Below the latter temperature the anhydrous salt cannot exist in the presence of H_2O , but is converted into $\text{Na}_2\text{SO}_4 + 7\text{H}_2\text{O}$, or $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$. The solubility of $\text{Na}_2\text{SO}_4 + 7\text{H}_2\text{O}$ increases with the temperature from $0-26^\circ$, and at 27° it is converted into the anhydrous salt.

Thus there are two different rates of solubility for Na_2SO_4 for temperatures from $0-18^\circ$, three different rates from $18-26^\circ$, two from $26-34^\circ$, and only one above 34° .

1. By heating $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ to fumes and raising the heat until the liquid boils, placing in a closed vessel and cooling, the greater part of the anhydrous salt, which separates out on heating, redissolves on cooling, and the amount increases as the temp. falls until 18° is reached. Below 18° $\text{Na}_2\text{SO}_4 + 7\text{H}_2\text{O}$ is formed. Saturated $\text{Na}_2\text{SO}_4 + \text{Aq}$ thus obtained contains for 100 pts. H_2O at

18° 20° 26°
53.25 52.76 51.53 51.31 pts. Na_2SO_4 .

30° 33° 34° 36°
50.87 49.71 49.53 49.27 pts. Na_2SO_4 .

2. By allowing the boiling saturated solution free from undissolved salt to cool to 0°

with exclusion of air until crystals of $\text{Na}_2\text{SO}_4 \cdot 7\text{H}_2\text{O}$ are formed, then removing the greater part of the mother liquor with a warm pipette, and warming the rest of the mother liquor with the excess of crystals, the crystals dissolve in increasing quantity between 0° and $26-27^\circ$, so that at 27° the solution contains 56 pts. Na_2SO_4 to 100 pts. H_2O . The remaining undissolved crystals of $\text{Na}_2\text{SO}_4 \cdot 7\text{H}_2\text{O}$ begin to melt very slowly at 27° , more quickly at higher temperatures, and cause the separation of anhydrous crusts, and thus the strength of the solution is gradually lowered to the normal. Saturated solutions prepared in this way contain for 100 pts. H_2O at:

0°	10°	13°
19.62	30.49	34.27 pts. Na_2SO_4 ,
or 44.89	78.9	92.9 pts. $\text{Na}_2\text{SO}_4 \cdot 7\text{H}_2\text{O}$.
15°	16°	17°
37.43	38.73	39.99 pts. Na_2SO_4 ,
or 105.8	117.4	111.0 pts. $\text{Na}_2\text{SO}_4 \cdot 7\text{H}_2\text{O}$,
18°	19°	20°
41.63	43.35	44.73 pts. Na_2SO_4 ,
or 124.6	133.0	140.0 pts. $\text{Na}_2\text{SO}_4 \cdot 7\text{H}_2\text{O}$,
25°	26°	
52.94	54.97	pts. Na_2SO_4 .
or 188.5	202.6	pts. $\text{Na}_2\text{SO}_4 \cdot 7\text{H}_2\text{O}$.

3. Solutions obtained by shaking H_2O with $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ contain for 100 pts. H_2O at:

0°	10°	15°
5.02	9.00	13.20 pts. Na_2SO_4 ,
or 12.16	23.04	35.96 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$,
18°	20°	25°
16.80	19.40	28.00 pts. Na_2SO_4 ,
or 48.41	58.85	98.48 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$,
26°	30°	
30.00	40.00	pts. Na_2SO_4 ,
or 109.81	184.1	pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$,
33°	34°	
50.76	55.0	pts. Na_2SO_4 .
or 323.1	412.2	pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$.

At 34° , $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ begins to melt in its crystal H_2O . As long as there is a considerable quantity of unchanged crystals present, the solution contains 55 pts. Na_2SO_4 for 100 pts. H_2O , but as the hydrous salt decreases in amount and becomes converted into the anhydrous salt, the solution becomes weaker and contains only 49.53 pts. Na_2SO_4 for 100 pts. H_2O after warming for 6 or 8 hours at 34° . In the same way temporary solutions can be obtained at $36-40^\circ$ with 55-56 pts. Na_2SO_4 to 100 pts. H_2O , but this amount sinks to the normal even more quickly than at 34° .

Na_2SO_4 dehydrated at $100-150^\circ$, after the addition of $1\frac{1}{2}-1\frac{3}{4}$ pts. H_2O , gives a solution between 0° and 32° of the same strength as $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$, but at 34° a solution with

55 pts. Na_2SO_4 to 100 pts. H_2O cannot be obtained, but one with 49.53 pts. is formed. (Löwel, A. ch. (3) 49. 32.)

4 Solubility of anhydrous salt. Above 34° , 100 pts. H_2O dissolve at:

35°	40°	45°	50°	55°
50.2	48.8	47.7	46.7	45.9 pts. Na_2SO_4 ,
60°	65°	70°	75°	80°
45.3	44.8	44.4	44.0	43.7 pts. Na_2SO_4 ,
85°	90°	95°	100°	103.5°
43.3	43.1	42.8	42.5	42.2 pts. Na_2SO_4 .

(Mulder.)

Solubility in 100 pts. H_2O at t° .

t°	Pts. Na_2SO_4	t°	Pts. Na_2SO_4	t°	Pts. Na_2SO_4
0	4.8	35	50.2	70	44.4
1	5.1	36	49.9	71	44.3
2	5.4	37	49.6	72	44.2
3	5.7	38	49.3	73	44.2
4	6.0	39	49.1	74	44.1
5	6.4	40	48.8	75	44.0
6	6.8	41	48.5	76	44.0
7	7.3	42	48.3	77	43.9
8	7.8	43	48.1	78	43.8
9	8.4	44	47.9	79	43.7
10	9.0	45	47.7	80	43.7
11	9.7	46	47.5	81	43.6
12	10.5	47	47.3	82	43.5
13	11.4	48	47.1	83	43.5
14	12.4	49	46.9	84	43.4
15	13.4	50	46.7	85	43.3
16	14.5	51	46.6	86	43.3
17	15.7	52	46.4	87	43.2
18	16.9	53	46.2	88	43.2
19	18.2	54	46.1	89	43.1
20	19.5	55	45.9	90	43.1
21	20.9	56	45.8	91	43.0
22	22.5	57	45.7	92	43.0
23	24.1	58	45.6	93	42.9
24	25.9	59	45.4	94	42.9
25	27.9	60	45.3	95	42.8
26	30.1	61	45.2	96	42.7
27	32.4	62	45.1	97	42.6
28	35.0	63	45.0	98	42.6
29	37.8	64	44.9	99	42.5
30	40.9	65	44.8	100	42.5
31	44.2	66	44.7	101	42.4
32	47.8	67	44.6	102	42.3
32.75	50.65	68	44.5	103	42.2
33	50.6	69	44.5	103.5	42.2
34	50.4				

(Mulder, Scheik. Verhandel, 1864. 123.)

100 pts. dissolve at:

0°	34°	100°	120°
5	78.8(?)	42.7	41.95 pts. Na_2SO_4 ,

140°	160°	180°	230°
42.0	42.9	44.25	46.4 pts. Na_2SO_4 .

(Tilden and Shenstone, Lond. R. Soc. Proc. 36. 345.)

Solubility decreases above 230°. (Étard, C. R. 113. 854)

Sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ contains at:

0° 7° 13° 24° 28° 30°
4.1 6.2 9.9 19.3 25.2 29.5% Na_2SO_4

49° 62° 83° 99° 134° 150°
32.8 31.3 30.0 29.7 29.4 29.8% Na_2SO_4

190° 240° 279° 320°
29.9 30.0 24.5 17.8% Na_2SO_4

(Étard, A. ch. 1894, (7) 2. 548.)

Solubility of Na_2SO_4 in H_2O at t°.
G. per 100 g. H_2O .

t°	Na_2SO_4	Sp. gr.	t°	Na_2SO_4	Sp. gr.
0.70	4.71	1.0432	33.5	49.39	1.3307
10.25	9.21	1.0802	38.15	48.47	1.3229
15.65	14.07	1.1150	44.85	47.49	1.3136
24.90	27.67	1.2067	60.10	45.22	1.2918
27.65	34.05	1.2459	75.05	43.59	1.2728
30.20	41.78	1.2894	89.85	42.67	1.2571
31.95	47.98	1.3230	101.9*	42.18	1.2450

* B.-pt.

(Berkeley, Phil. Trans. Roy. Soc. 1904, 203. A, 189.)

Transition point from $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ to $\text{Na}_2\text{SO}_4 = 32.5^\circ$ (Berkeley); 32.383° . (Richards and Churchill, Z. phys. Ch. 1899, 28. 314.)

100 g. $\text{Na}_2\text{SO}_4 + \text{Aq}$ sat. at 15° contain 11.5 g. anhydrous Na_2SO_4 ; 21.9 g. at 25° . (Schreinemakers, Arch. Néer. Sc. 1910, (2) 15. 81.)

1 l. $\text{Na}_2\text{SO}_4 + \text{Aq}$ sat. at 25° contains 1.881 mols. Na_2SO_4 . (Herz, Z. anorg. 1911, 70. 127)

Solubility in H_2O at t°.

t°	Mol. % Na_2SO_4
62	5.39
70	5.27
72	5.25
80	5.18
120	5.04
190	5.25
192	5.27
208	5.39
241	5.39
250	5.04
279	4.12
319	2.56
252	4.9
310	3.2
340	1.8
365	0.0

(Witte, Z. phys. Ch. 1913, 86. 364.)

Supersaturated solutions of Na_2SO_4 are easily formed; when $\text{Na}_2\text{SO}_4 + \text{Aq}$ sat. at its b.-pt. is hermetically sealed, no crystals are deposited on cooling (Löwel). Supersat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ may also be obtained by cooling hot sat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ in flasks loosely stoppered with cotton wool (Schroeder, A. 100. 45), or by covering the containing vessel with a glass plate, watch-glass, card, etc., or by covering the liquid itself with a layer of oil, and then allowing to cool.

Hot $\text{Na}_2\text{SO}_4 + \text{Aq}$ containing 1 pt. H_2O to 1 pt. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ does not crystallise on slowly cooling or on being quickly cooled by immersion in cold water, if it is contained in a barometer tube freed from air by boiling, or in an exhausted well-closed vessel, or in an open vessel with a layer of oil of turpentine on it (Gay-Lussac); or in a vessel containing air, either well stoppered or furnished with a loose cover (Schweigger); or in an open vessel under a bell jar full of air and closed at the bottom with a water joint; or in open bottles placed in a quiet situation; or in an open glass enclosed in a stoppered vessel, containing air and some KOH for drying; in this case $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ effloresces from the solution, and when washed down again does not cause instant crystallisation, but redissolves.

The crystallisation of a solution cooled in this way may often be brought about instantaneously, or often again after a short time; (1) by agitation, when the solution has been cooled in an open vessel; (2) by access of air caused by opening the vessel, the crystallisation taking place the more rapidly the larger the opening. In this case the crystallisation begins at the top, where the solution, the vessel, and the air come in contact, when a particle of dust falls in the liquid the crystallisation begins a little under the surface. When the solution has been cooled in vacuo, a bubble of air, hydrogen, carbonic acid or nitrous oxide is sufficient to set up the crystallisation; (3) by contact with a solid body. The latter do not cause crystallisation when cooled in contact with the liquid, nor excepting a crystal of $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ when they are moistened or warmed before contact with the solution.

Supersat. $\text{Na}_2\text{SO}_4 + \text{Aq}$ is brought to crystallisation by addition of a crystal of $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$, or an isomorphous substance as $\text{Na}_2\text{SeO}_4 + 10\text{H}_2\text{O}$, or $\text{Na}_2\text{CrO}_4 + 10\text{H}_2\text{O}$. Other crystals, as $\text{MgSO}_4 + 7\text{H}_2\text{O}$, etc., have no action. (Thomson, Chem. Soc. 25. 199)

See also Hartley, Jones and Hutchinson, Chem. Soc. 1908, 93. 825, on "Spontaneous crystallisation of sodium sulphate solutions," and de Coppet (A. ch. 1907, (8) 10. 457) on same subject.

A more extended discussion of the phenomena and causes of supersaturation is not considered to be within the scope of this work.

$\text{Na}_2\text{SO}_4 + \text{Aq}$ sat. at 15° has sp. gr. 1.10547 (Mandel and Kraft); at 15° has sp. gr. 1.119 (Stollé), at 10°

has sp. gr. 1.1162 (Stolba); at 10° contains 29 pts. Na₂SO₄ to 100 pts. H₂O (supersaturated?), and has sp. gr. 1.1259 (Karsten).

• Sp. gr. of Na₂SO₄ + Aq at 19.5°.

% Na ₂ SO ₄	Sp. gr.	% Na ₂ SO ₄	Sp. gr.
2.894	1.0262	10.538	1.0977
5.589	1.0509	12.473	1.1162
7.995	1.0733

(Kremers, Pogg. 95. 120.)

Sp. gr. of Na₂SO₄ + Aq.

% Na ₂ SO ₄ + 10H ₂ O	Sp. gr.	% Na ₂ SO ₄ + 10H ₂ O	Sp. gr.
1.262	1.005	13.744	1.055
2.522	1.010	14.975	1.060
3.780	1.015	16.203	1.065
5.035	1.020	17.426	1.070
6.288	1.025	18.645	1.075
7.538	1.030	19.860	1.080
8.786	1.035	21.071	1.085
10.030	1.040	22.277	1.090
11.272	1.045	23.478	1.095
12.510	1.050	24.674	1.100

(Schmidt, Pogg. 132. 132.)

Sp. gr. of Na₂SO₄ + Aq at 19°.

% Na ₂ SO ₄ + 10H ₂ O	Sp. gr.	% Na ₂ SO ₄ + 10H ₂ O	Sp. gr.
1	1.0040	16	1.0642
2	1.0079	17	1.0683
3	1.0118	18	1.0725
4	1.0158	19	1.0766
5	1.0198	20	1.0807
6	1.0232	21	1.0849
7	1.0278	22	1.0890
8	1.0318	23	1.0931
9	1.0358	24	1.0973
10	1.0398	25	1.1015
11	1.0439	26	1.1057
12	1.0479	27	1.1100
13	1.0520	28	1.1142
14	1.0560	29	1.1184
15	1.0601	30	1.1226

(Schiff, A. 110. 70.)

Sp. gr. of Na₂SO₄ + Aq at 15°.

%	Sp. gr. if Na ₂ SO ₄	Sp. gr. if Na ₂ SO ₄ + 10H ₂ O	%	Sp. gr. if Na ₂ SO ₄ + 10H ₂ O	%	Sp. gr. if Na ₂ SO ₄ + 10H ₂ O
1	1.0091	1.004	11	1.044	21	1.086
2	1.0182	1.008	12	1.047	22	1.090
3	1.0274	1.013	13	1.052	23	1.094
4	1.0365	1.016	14	1.056	24	1.098
5	1.0457	1.020	15	1.060	25	1.103
6	1.0550	1.024	16	1.064	26	1.107
7	1.0644	1.028	17	1.069	27	1.111
8	1.0737	1.032	18	1.073	28	1.116
9	1.0832	1.036	19	1.077	29	1.120
10	1.0927	1.040	20	1.082	30	1.125

(Gerlach, Z. anal. 8. 287.)

Sp. gr. of Na₂SO + Aq at 24.8°. a = no. of g., equivalent to ½ mol. wt., dissolved in 1000 g. H₂O; b = sp. gr. if a is Na₂SO₄ + 10H₂O, ½ mol. wt. = 161; c = sp. gr. if a is Na₂SO₄, ½ mol. wt. = 71.

a	b	c	a	b	c
1	1.054	1.059	4	1.163	1.213
2	1.098	1.114	5	1.188	...
3	1.134	1.165	6	1.209	...

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of Na₂SO₄ + Aq at 18°.

% Na ₂ SO ₄	Sp. gr.	% Na ₂ SO ₄	Sp. gr.
5	1.0450	15	1.1426
10	1.0915

(Kohlrausch, W. Ann. 1879. 1.)

Sp. gr. of Na₂SO₄ + Aq at 20° containing 0.5 mol. Na₂SO₄ to 100 mols. H₂O = 1.03466; 1.0 mol. Na₂SO₄ to 100 mols. H₂O = 1.06744. (Nicol, Phil. Mag. (5) 16. 122.)

Sp. gr. of Na₂SO₄ + Aq at 25°.

Concentration of Na ₂ SO ₄ + Aq	Sp. gr.
1-normal	1.0606
1/5 " "	1.0309
1/6 " "	1.0156
1/8 " "	1.0079

(Wagner, Z. phys. Ch. 1890, 5. 39.)

Sp. gr. at 16°/4° of Na₂SO₄ + Aq containing 9.4043% Na₂SO₄ = 1.08655. (Schönrock, Z. phys. Ch. 1893, 11. 781.)

Na₂SO₄ + Aq containing 25.51% Na₂SO₄ has sp. gr. 20°/20° = 1.2527. Na₂SO₄ + Aq containing 10.14% Na₂SO₄ has sp. gr. 20°/20° = 1.0938. (Le Blanc and Rohland, Z. phys. Ch. 1896, 19. 278.)

Sp. gr. of Na₂SO₄ + Aq at 17.5°, when p = per cent strength of solution; d = observed density; and w = volume conc. in grs. per cc. ($\frac{pd}{100} = w$)

p.	d.	w.
13.06	1.1226	0.14662
11.75	1.1094	0.13043
10.68	1.0990	0.11737
8.544	1.0784	0.09214
6.762	1.0615	0.07178
4.015	1.0358	0.04159
2.599	1.0225	0.02658
2.375	1.0204	0.02423
1.818	1.0154	0.01846
1.349	1.0109	0.01364
0.5204	1.0037	0.00522
0.2921	1.0014	0.00293

(Barnes, J. phys. Chem. 1898, 2. 543.)

Sp. gr. of Na₂SO₄+Aq at 20°.

Normality of Na ₂ SO ₄ +Aq	% Na ₂ SO ₄	Sp. gr.
0.97	12.36	1.1138
0.48	6.41	1.0570

(Forchheimer, Z. phys. Ch. 1900, 34. 23.)

Sp. gr. of sat. Na₂SO₄, 10H₂O+Aq at t°.

t°	wt. of 1 ccm. of the solution	100 g. H ₂ O dissolve g. Na ₂ SO ₄ +10H ₂ O
0	1.040	12.16
5	1.058	...
10	1.078	21.04
15	1.109	35.96
18	1.137	48.41
20	1.156	58.35
25	1.209	98.48
26	1.222	109.81
30	1.287	184.1
33	1.312	323.1
34	1.317	413.2
35	1.317	...

(Tschernaj, J. Russ. Phys. Chem. Soc. 1914, 46. 8.)

Sp. gr. and b.-pt. of Na₂SO₄+Aq. Na₂SO₄+Aq containing P pts. Na₂SO₄+10H₂O for every 100 pts. H₂O has given sp. gr. and b.pt.

P	Sp. gr.	B.-pt.	P	Sp. gr.	B.-pt.
1	1.005	100.5°	16	1.064	101.25°
2	1.008	100.62	17	1.067	101.25
3	1.014	100.62	18	1.070	101.37
4	1.020	100.75	19	1.072	101.37
5	1.021	100.75	20	1.074	101.37
6	1.028	100.87	21	1.076	101.37
7	1.030	100.87	22	1.078	101.5
8	1.032	101.0	23	1.080	101.5
9	1.036	101.0	24	1.082	101.5
10	1.040	101.0	25	1.084	101.5
11	1.043	101.12	26	1.090	101.5
12	1.050	101.12	27	1.092	101.63
13	1.055	101.25	28	1.095	101.63
14	1.060	101.25	29	1.098	101.63
15	1.062	101.25	30	1.100	101.75

(Brandes and Gruner, 1837.)

Saturated solution boils at 103.17° (Löwel), 103.5° (Mulder), 105° (Kremers), 100.5° (Griffiths), 100.8° (Gerlach).

Crust forms at 102.9°; highest temp., 103.2°, and solution contains 43.9 pts. Na₂SO₄ to 100 pts. H₂O. (Gerlach, Z. anal. 26. 426.)

B.-pt. of Na₂SO₄+Aq containing pts. Na₂ to 100 pts. H₂O.

B.-pt.	Pts. Na ₂ SO ₄	B.-pt.	Pts. Na ₂ SO ₄
100.5°	9.5	102.5°	39.1
101.0	18.0	103.0	44.3
101.5	26.0	103.2	46.7
102.0	33.0

(Gerlach, Z. anal. 26. 430.)

M.-pt. of Na₂SO₄+10H₂O=34°. (Tik Chem. Soc. 45. 409.)

Sol. with decomp. in HCl+Aq.

Solubility in H₂SO₄+Aq at 25°.

1000 g. of the solution contain		Solid phase
Mols H ₂ SO ₄	Mols Na ₂ SO ₄	
...	1.539	Na ₂ SO ₄ , 10H ₂ O
0.286	1.671	"
0.338	1.742	"
0.884	2.256	Na ₂ SO ₄ , 10H ₂ O+Na ₂ SO ₄
1.576	2.363	Na ₂ SO ₄ +Na ₂ H(SO ₄) ₂
1.666	2.437	"
2.611	2.091	Na ₂ H(SO ₄) ₂ +Na ₂ H(SO ₄) ₂ .l

(D'Ans, Z. anorg. 1906, 49. 356.)

Solubility of Na₂SO₄ in H₂SO₄+Aq at

1000 g. of the solution contain		Solid phase
Mol. Na ₂ SO ₄	Mol. H ₂ SO ₄	
1.55	0.08	Na ₂ SO ₄
1.59	0.147	"
1.85	0.60	Na ₂ SO ₄ , 10H ₂ O
2.00	0.763	"
0.77	4.23	NaHSO ₄ . H ₂ O
0.47	4.96	"
0.32	6.61	Na ₂ HSO ₄
0.305	6.87	"
0.07	7.18	Na ₂ H(SO ₄) ₂
0.79	8.78	"

(D'Ans, Z. anorg. 1909, 61. 92.)

10 ccm. of sat. Na₂SO₄+absolute H₂SO₄ contain approx. 2.999 g. Na₂SO₄. (Berl Z. phys. Ch. 1910, 72. 355.)

Solubility in H ₂ SO ₄ + Aq at 25°.	
Solid Phase, Na ₂ SO ₄ + 10H ₂ O.	
Millimols H ₂ SO ₄ in 10 ccm.	Millimols Na ₂ SO ₄ in 10 ccm.
...	18.81
5.10	22.38
7.79	24.65

(Herz, Z. anorg. 1912, 73. 276.)

1000 g. of the solution contain		Solid phase
Mols SO ₃	Mols Na ₂ SO ₄	
5.91	0.409	NaHSO ₄
6.30	0.332	"
6.64	0.297	NaHSO ₄ + NaH ₂ (SO ₄) ₂ , H ₂ O
6.90	0.173	NaH ₂ (SO ₄) ₂ , H ₂ O
7.36	0.071	"
7.74	0.047	"
7.82	0.044	"
8.12	0.037	"
8.29	0.042	"
8.40	0.046	"
8.70	0.076	"
8.86	0.156	"
8.93	0.259	"
8.93	0.269	"
8.93	0.273	"
8.84	0.527	"
8.73	0.681	"
8.70	0.808	NaH ₂ (SO ₄) ₂ , H ₂ O
8.62	0.834	metastable solutions
8.62	0.844	"
8.61	0.899	"
8.87	0.445	NaH ₂ (SO ₄) ₂ , H ₂ O + Na ₂ SO ₄ , 4.5H ₂ SO ₄
8.93	0.437	Na ₂ SO ₄ , 4.5H ₂ SO ₄
9.08	0.394	"
9.36	0.425	Na ₂ SO ₄ , 4.5H ₂ SO ₄ + NaHS ₂ O ₇
9.18	0.567	NaHS ₂ O ₇
9.42	0.728	"
9.48	0.760	"
9.55	0.775	"
9.48	0.953	NaHS ₂ O ₇ + ?
9.85	0.787	?
9.98	0.908	?
(9.77)	(1.03)	Metastable
10.16	0.797	"
10.78	0.302	?

(D'Ans, Z. anorg. 1913, 80. 236.)

Sl. sol. in conc. HC₂H₃O₂. (Ure's Dict.)
Not pptd. by addition of glacial HC₂H₃O₂ to
Na₂SO₄ + Aq. (Persoz.)

Solubility in NaOH + Aq at 25°.		
1000 g. of the solution contain		Solid phase
Mols (NaOH),	Mols Na ₂ SO ₄	
0	1.54	Na ₂ SO ₄ , 10H ₂ O
0.074	1.41	"
0.70	1.08	"
1.47	0.90	Na ₂ SO ₄ , 10H ₂ O + Na ₂ SO ₄
2.02	0.59	Na ₂ SO ₄
2.82	0.24	"
3.52	0.126	"
5.83	0.013	"
6.62	...	NaOH, H ₂ O

(D'Ans and Schreiner, Z. anorg. 1910, 67. 437.)

Sol. in sat. NH₄Cl + Aq.
Rapidly and abundantly sol. in sat. KCl +
Aq with pptn. of K₂SO₄.
Na₂SO₄ + 10H₂O is sol. in sat. NaCl + Aq
without pptn. If effloresced Na₂SO₄ is used,
a ppt. of NaCl is caused at first, and subse-
quently of Na₂SO₄ + 10H₂O. (Karsten.)
Sol. in boiling sat. NaCl + Aq with pptn. of
NaCl, but from cold solutions the Na₂SO₄
separates out first. (Vauquelin.)
Less sol. in NaCl + Aq than in H₂O. (Hunt,
Am. J. Sci. (2) 25. 368.)

Solubility in NaCl + Aq at t°.		
t°	g. NaCl per 100 g. H ₂ O	g. Na ₂ SO ₄ per 100 g. H ₂ O
10	0.00	9.14
	4.28	6.42
	9.60	4.76
	15.65	3.99
	21.82	3.97
	28.13	4.15
	30.11	4.34
	32.27	4.59
	33.76	4.75
21.5	0.00	21.33
	9.05	15.48
	17.48	13.73
	20.41	13.62
	26.01	15.05
	26.53	14.44
	27.74	13.39
	31.25	10.64
	31.80	10.28
	32.10	8.43
	33.69	4.73
	34.08	2.77
	35.46	0.00
25	0.00	28.74
	2.74	26.57
	8.15	23.15
	19.86	20.52
	24.58	14.86
	31.21	9.95
	32.02	9.61

Solubility in NaCl + Aq at t° —Continued.

t°	g. NaCl per 100 g. H ₂ O	g. Na ₂ SO ₄ per 100 g. H ₂ O
27	0.00	31.10
	2.66	28.73
	5.29	27.17
	7.90	26.02
	16.13	24.83
	18.91	21.39
	19.64	20.11
	20.77	19.29
	32.33	9.53
30	0.00	39.70
	2.45	38.25
	5.61	36.50
	7.91	35.96
	10.61	31.64
	12.36	29.87
	15.65	25.02
	18.44	21.30
	20.66	19.06
	32.43	9.06
33	0.00	48.48
	1.22	46.49
	1.99	45.16
	2.64	44.09
	3.47	42.61
	12.14	29.32
	21.87	16.83
	32.84	8.76
	33.99	4.63
	34.77	2.75
35	0.00	47.94
	2.14	43.75
	13.57	26.26
	18.78	19.74
	31.91	8.28
	35.63	0.00

At 33° and above the values represent the solubility of Na₂SO₄ in NaCl + Aq. At 10° the solid phase in contact with the solution is probably Na₂SO₄ · 7H₂O. Between 17° and 33° the solid phase is either Na₂SO₄ · 10H₂O or Na₂SO₄. An inversion of Na₂SO₄ · 10H₂O to Na₂SO₄ takes place at various temp. below 33°, depending on the amount of NaCl contained in the solution in contact with the solid sodium sulphate.

(Seidell, Am Ch J. 1902, 27, 55.)

Solubility in NaCl + Aq at 15°

Composition of the sat. solution		Solid Phase
% by wt. Na ₂ SO ₄	% by wt. NaCl	
11.5	0	Na ₂ SO ₄ + 10H ₂ O
7.86	5.42	"
5.87	11.51	"
5.23	15.97	"
5.26	21.03	"
5.64	23.39	Na ₂ SO ₄ · 10H ₂ O + NaCl
2.26	25.21	NaCl
0	26.3	"

(Schreinemakers and de Baat, Z. phys. Ch. 1909, 67, 554.)

Sol. in sat. NH₄NO₃ + Aq. (Marguerite, C. R. 38, 307.)

Sol. in sat. KNO₃ + Aq with pptn after several hours. (Karsten.)

Na₂SO₄ + 10H₂O is sol. in sat. NaNO₃ + Aq without pptn, but if effloresced Na₂SO₄ is used, NaNO₃ is pptd. at first, and subsequently Na₂SO₄ + 7H₂O.

The presence of CaSO₄ does not affect the solubility of Na₂SO₄ in H₂O to any great extent. (Barre, A. ch. 1911, (8) 24, 160.)

More sol. in K₂SO₄, CuSO₄, MgSO₄ + Aq than in H₂O. (Pfaff, A. 99, 226.)

100 pts. H₂O dissolve 20.7 pts. CuSO₄ and 15.9 pts. Na₂SO₄. (Rüdorff, B. 6, 484.)

Sol. in sat. MgSO₄, K₂SO₄, CuSO₄ + Aq, but if more Na₂SO₄ than can be dissolved is added to the CuSO₄ + Aq, a large quantity of a double sulphate separates out. (Karsten.)

The solubility of Na₂SO₄ in K₂SO₄ + Aq has been determined at 15°, 25°, 40°, 50°, 60°, 70° and 80°. From the results the conclusion is drawn that sodium and potassium sulphates form a double salt of the formula K₂Na₂(SO₄)₃. (Okada, Chem Soc. 1915, 106, (2) 344.)

See also under CuSO₄, MgSO₄, and K₂SO₄.

Slowly but abundantly sol. in sat. ZnSO₄ + Aq, with separation of a double salt after a few days.

Solubility of ZnSO₄ · 7H₂O + Na₂SO₄ · 10H₂O in 100 g. H₂O at t° .

t°	grams ZnSO ₄	grams Na ₂ SO ₄
0	41.815	7.905
5	42.285	9.515

(Koppel, Z. phys. Ch. 1905, 52, 409.)

See also under Na₂Zn(SO₄)₂ + 4H₂O.

Solubility of $\text{Na}_2\text{SO}_4 + \text{Th}(\text{SO}_4)_2$ at 16° .
Solid phase $\text{Th}(\text{SO}_4)_2$.

Pts. per 100 pts. H_2O		Pts. per 100 pts. H_2O	
Na_2SO_4	$\text{Th}(\text{SO}_4)_2$	Na_2SO_4	$\text{Th}(\text{SO}_4)_2$
1.094	1.743	5.79	2.136
1.960	2.387	9.35	1.379
2.84	3.800	12.24	1.169
2.98	3.962	15.36	1.048
4.11	3.375

(Barre, C. R. 1911, 150. 155.)

Solubility in Na acetate + Aq at 25° .
Solid phase, $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$.

Composition of the solutions		
% Na acetate	% Na_2SO_4	% H_2O
0	21.9	78.10
4.10	17.72	78.18
7.71	16.48	75.81
12.58	13.50	73.92
16.26	11.50	72.24
20.63	8.10	71.27

(Fox, Chem. Soc. 1909, 95. 888.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Alcohol precipitates $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$ from the cold saturated aqueous solution. (Brandes and Firnhaber.)
Insol. in alcohol of from 0.817 to 0.90 sp. gr. (Kirwan.)
1000 pts. alcohol of 0.872 sp. gr. dissolve 0.7 pt. Na_2SO_4 at $12.5-15^\circ$; of 0.905 sp. gr. dissolve 3.8 pts. Na_2SO_4 at $12.5-15^\circ$.
Insol. in alcohol of 0.83-0.85 sp. gr. (Anthon.)

From supersaturated solution in alcohol, crystals with $7\text{H}_2\text{O}$ are formed. (Schiff, A. 106. 11.)

100 pts. 10% alcohol at 15° contain 14.35 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$; 20% alcohol at 15° contain 5.6 pts. $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$; 40% alcohol at 15° contain 1.3% $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$. (Schiff, A. 118. 365.)

Very sl. sol. in abs. alcohol at ord. temp.; somewhat more, though still exceedingly sparingly, sol. in abs. alcohol acidulated with H_2SO_4 . (Fresenius.)

Alcohol does not affect crystal H_2O of $\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$.

Solubility of Na_2SO_4 in alcohol + Aq at t° .

t°	% alcohol	g. per 100 g. solution			solid phase
		H_2O	alcohol	Na_2SO_4	
15	0.7	88.7	0.0	11.3	$\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$
	9.2	85.1	8.6	6.3	"
	19.4	78.6	18.9	2.9	"
	39.7	60.0	39.5	0.5	"
	58.9	41.1	58.8	0.1	"
	72.0	28.0	72.0	0.0	"
	0.0	72.8	0.0	27.2	$\text{Na}_2\text{SO}_4 + 7\text{H}_2\text{O}$
	11.2	76.5	9.5	14.0	"
	20.6	74.3	19.2	6.5	"
	30.2	68.4	29.6	2.0	"
25	0.0	78.1	0.0	21.9	$\text{Na}_2\text{SO}_4 + 10\text{H}_2\text{O}$
	10.6	78.5	9.3	12.2	"
	24.0	72.8	22.9	4.3	"
	54.0	45.6	54.0	0.4	"
36	0.0	67.0	0.0	33.0	Na_2SO_4
	8.8	70.0	6.8	22.6	"
	12.8	71.2	10.5	18.3	"
	17.9	71.1	15.5	13.4	"
	18.1	71.0	15.7	13.3	"
	28.9	66.5	28.4	5.1	"
	48.7	50.9	48.3	0.8	"
45	0.0	67.6	0.0	32.4	"
	9.0	71.3	7.1	21.6	"
	14.5	71.8	12.1	16.1	"
	20.6	70.6	18.4	10.0	"
	31.0	65.6	29.5	4.9	"

Between certain concentrations of alcohol, the liquid separates into two layers at 25° , 36° and 45° , of the following composition.

t°	Upper Layer			Lower Layer		
	% H_2O	% alcohol	% Na_2SO_4	% H_2O	% alcohol	% Na_2SO_4
25	66.5	27.3	6.2	67.4	5.1	27.5
	68.1	23.9	8.0	68.5	6.0	25.5
	68.3	23.1	8.6	68.3	6.7	25.0
36	66.6	4.1	29.3
	57.7	38.4	3.9
	65.0	28.3	6.7	68.8	5.9	25.3
45	68.1	21.2	10.7	68.9	9.4	21.7
	61.8	32.9	5.3
	65.8	25.3	8.9	68.4	8.8	22.8
	66.0	24.0	10.0	68.6	10.1	21.3

(de Bruyn, Z. phys. Ch. 1900, 32. 101.)

Solubility in alcohol + Aq at 25°.

Composition of the sat. solution			Solid phase
% by wt. H ₂ O	% by wt. alcohol	% by wt. Na ₂ SO ₄	
63.41	34.84	1.75	Na ₂ SO ₄ · 10H ₂ O
49.0	50.5	0.5	
46.6	53.0	0.4	Na ₂ SO ₄ · 10H ₂ O + Na ₂ SO ₄
34.9	64.95	0.15	Na ₂ SO ₄

(Schreinemakers, Z. phys. Ch. 1909, 67. 552.)

Solubility data for solution of NaCl in ethyl alcohol + Aq. at 15°, 25°, and 30° are given by Schreinemakers (Z. phys. Ch. 1909, 67. 556).

Solubility in propyl alcohol + Aq at 20°.

% propyl alcohol	g. Na ₂ SO ₄ per 100 g. solution	% propyl alcohol	g. Na ₂ SO ₄ per 100 g. solution
42.20	1.99	56.57	0.55
49.77	1.15	60.64	0.44
55.65	0.72	62.81	0.38

(Linebarger, Am. Ch. J. 1892, 14. 380.)

Sol. in glycerine

Insol. in acetone. (Naumann, B. 1904 37. 4329; Eidmann, C. C. 1899, II, 1014); benzonitrile. (Naumann, B. 1914, 47. 1370) methyl acetate (Naumann, B. 1909, 42. 7790); ethyl acetate. (Naumann, B. 1910, 43. 314.)

100 g. H₂O dissolve 183.7 g. sugar + 30.5 g. Na₂SO₄ at 31.25°, or 100 g. sat. solution contain 52.2 g. sugar + 9.6 g. Na₂SO₄. (Köhler, Z. Ver. Zuckerind, 1897, 47. 447.)

Min. Anhydrous, *Thenardite*. +10H₂O, *Mirabilite*.

Sodium hydrogen sulphate, NaHSO₄.

Not deliquescent. Very sol. in H₂O with decomposition.

Sol. in 2 pts. H₂O at 0° (Link); 1 pt. H₂O at 100° (Schubarth). 100 pts. H₂O at 15.5° dissolve 92.72 pts. Sol. in 2 pts. H₂O at 18.75° (Abl); decomp. by alcohol.

Insol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

+H₂O. Deliquescent, and decomp. by the H₂O which it takes up

NaHSO₄ · 2H₂O. Decomp. by H₂O. (Schultz)

Trisodium hydrogen sulphate, Na₃H(SO₄)₂.

Sol. in H₂O with decomp.

+H₂O. (Rose)

Sodium pyrosulphate, Na₂S₂O₇.

Sol. in fuming H₂SO₄ without decomp.

Sodium thallic sulphate, Na₂SO₄ · Tl₂(SO₄)₃.

Sol. in H₂O. (Strecker, A. 135. 207.)

Sodium thorium sulphate, Na₂SO₄ · Th(SO₄)₂ · 6H₂O.

Sol. in H₂O. 100 pts. cold sat. Na₂SO₄ + 4g dissolve 4 pts. of this salt. (Cleve.)

See also under Na₂SO₄ + ThSO₄.

Sodium titanium sesquisulphate, Na₂Ti₂(SO₄)₃ · 5H₂O.

Sol. in H₂O. (Spence, Chem. Soc. 1904, 86. (2) 412.)

Insol. in alcohol. (Knecht, B. 1903, 36. 169.)

Sodium titanyl sulphate, Na₂TiO₂(SO₄)₂ · 10H₂O.

Hygroscopic. (Massuchelli and Pantanelli, C. C. 1909, II. 420.)

Sodium uranyl sulphate, Na₂(UO₂)(SO₄)₂ · 3H₂O.

(de Coninck, C. C. 1905, I. 919.)

Sodium vanadium sulphate, Na₂V₂(SO₄)₃ · 24H₂O.

Very sol. in H₂O. (Piccini, Z. anorg. 1897, 13. 444.)

Sodium vanadyl sulphate, Na₂SO₄ · VOSO₄ · 4H₂O.

Easily sol. in H₂O and alcohol + conc. H₂SO₄. (Koppel, Z. anorg. 1903, 35. 177.) Na₂SO₄ · 2VOSO₄ · 2½H₂O. Slowly sol. in H₂O.

Can be cryst. from H₂SO₄ at 100°. (Koppel.)

Sodium yttrium sulphate, Na₂SO₄ · Y₂(SO₄)₃ · 2H₂O.

Quite sol. in H₂O. (Cleve.)

The only double salt capable of existing at 25°. (James and Holden, J. Am. Chem. Soc. 1913, 35. 562.)

Sodium zinc sulphate, Na₂SO₄ · ZnSO₄ · 4H₂O.

Deliquescent in moist air.

Decomp. into constituents on dissolving in H₂O. (Graham, Phil. Mag. 13. 417.)

Solubility of Na₂Zn(SO₄)₂ · 4H₂O in 100 g. H₂O at t°.

t°	grams ZnSO ₄	grams Na ₂ SO ₄
25	26.32	23.40
30	26.475	23.445
35	26.365	23.525
40	26.68	23.63

(Koppel, Z. phys. Ch. 1905, 31. 499.)

solubility of $\text{Na}_2\text{Zn}(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{ZnSO}_4$,
7H₂O in 100 g. H₂O at t°.

t°	grams ZnSO ₄	grams Na ₂ SO ₄	t°	grams ZnSO ₄	grams Na ₂ SO ₄
0	45.79	11.24	25	56.155	7.215
5	48.81	10.175	30	60.55	6.34
0	52.34	8.625	35	65.25	5.64

(Koppel.)

solubility of $\text{Na}_2\text{Zn}(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{ZnSO}_4$,
6H₂O in 100 g. H₂O at t°.

t°	grams ZnSO ₄	grams Na ₂ SO ₄
38	66.64	4.98
40	64.89	4.71

(Koppel.)

solubility of $\text{Na}_2\text{Zn}(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{Na}_2\text{SO}_4$,
10H₂O in 100 g. H₂O at t°.

t°	grams ZnSO ₄	grams Na ₂ SO ₄
10	43.495	12.35
15	36.925	16.71
20	28.77	21.98
25	19.935	29.875
30	10.67	42.515

(Koppel.)

solubility of $\text{Na}_2\text{Zn}(\text{SO}_4)_2 \cdot 4\text{H}_2\text{O} + \text{Na}_2\text{SO}_4$,
(anhydrous) in 100 g. H₂O at t°.

t°	grams ZnSO ₄	grams Na ₂ SO ₄
35	8.725	46.61
40	9.16	43.835

(Koppel.)

strontium sulphate fluoride, Na_2SO_4 , NaF.

Cryst from H₂O without decomp. (Marignac, Ann. Min. (5) 15. 236.)

strontium sulphate antimony trifluoride.

See Antimony trifluoride sodium sulphate.

strontium sulphate, SrSO_4 .

Very sl. sol. in cold, and still less in boiling O.

1 l. H₂O at 11–15° dissolves 0.066 g. SrSO_4 (Brandes and Silber); 0.145 g. SrSO_4 (Fresenius); 0.154–0.167 g. SrSO_4 (Marignac); 87 g. SrSO_4 (Kremers); 0.278 g. SrSO_4 (Andrews).

1 l. boiling H₂O dissolves 0.104 g. SrSO_4 (Fresenius); 0.282 g. SrSO_4 (Brandes and Silber).

When a Sr salt is precipitated by H₂SO₄, 1 g. SrSO_4 remains dissolved in 700 pts. H₂O. (Marignac.)

Sol. in about 8000 pts. H₂O. (Schweitzer, J. B. 1877. 1054.)

Calculated from electrical conductivity of the solution, SrSO_4 is sol. in 10,070 pts. H₂O at 16.1° and 10,090 pts. at 20.1° (Holleman, Z. phys. Ch. 12. 131.)

1 l. H₂O dissolves 107 mg. SrSO_4 at 18° and not much more at higher temp. (Kohlrausch and Rose, Z. phys. Ch. 12. 241.)

100 g. H₂O dissolve

at t° 0°–5° 10°–12° 20° 30°
g. SrSO_4 0.0983 0.0994 0.1479 1.0600

at t° 50° 80° 90° 95–98°
g. SrSO_4 0.1629 0.1688 0.1727 0.1789

(Wolfmann, C. C. 1897, I. 632.)

1 l. H₂O dissolves 114 mg. SrSO_4 at 18°. (Kohlrausch, Z. phys. Ch. 1904, 50. 356.); 114.3 mg at 18°. (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)

Sol. in 6895 pts. cold, and 9638 pts. boiling H₂O; in 11,000–12,000 pts. H₂O containing H₂SO₄; in 474 HCl+Aq containing 8.5% HCl; in 432 pts. HNO₃+Aq containing 4.8% N₂O₅; in 7843 pts. HC₂H₃O₂+Aq containing 15.6% HC₂H₃O₂. (Fresenius.)

Or, 1 l. cold HCl+Aq of 8.5% dissolves 2.11 g. SrSO_4 ; 1 l. cold HNO₃+Aq of 4.8% N₂O₅ dissolves 2.31 g. SrSO_4 ; 1 l. cold HC₂H₃O₂+Aq of 15.6% HC₂H₃O₂ dissolves 0.1275 g. SrSO_4 . (Fresenius.)

Solubility of SrSO_4 in HCl+Aq.

No. cc HCl + Aq containing 1 mg. equiv. HCl	g. per 100 cc. solution	
	HCl	SrSO_4
0.2	18.23	0.161
0.5	7.29	0.207
1.0	3.65	0.188
2.0	1.82	0.126
10.0	0.36	0.048

(Banthisch, J. pr. 1884, (2) 29. 54.)

Solubility of SrSO_4 in HNO₃+Aq.

No. cc. HNO ₃ + Aq containing 1 mg. equiv. HNO ₃	g. per 100 cc. solution	
	HNO ₃	SrSO_4
0.2	31.52	0.381
0.5	12.61	0.307
1.0	6.30	0.217
2.0	3.15	0.138
10.0	0.63	0.049

(Banthisch, J. pr. 1884, (2) 29. 54.)

Sol. in conc H₂SO₄. See under $\text{SrH}_2(\text{SO}_4)_2$. Insol. in NH₄Cl+Aq or conc. (NH₄)₂SO₄+Aq. (Rose.)

Slowly but completely sol. in NaCl+Aq. (Wackenroder.)

H₂O containing Na₂SO₄ dissolves less SrSO_4 .

than pure H₂O; H₂O containing H₂SO₄ still less. (Andrews, Phil. Mag. Ann. 7. 406.)
 Insol. in Na₂S₂O₃+Aq.
 Insol. in boiling conc. (NH₄)₂SO₄+Aq. (Rose, Pogg. 110. 292.)
 Sol. in 16.949 pts. (NH₄)₂SO₄+Aq (1:4). (Fresenius, Z. anal. 32. 195.)
 Pptn. is hindered by alkali metaphosphates and citrates, but not by citric acid.
 Decomp. at ord. temp., and more rapidly on boiling by alkali carbonates+Aq.
 Sol. in MgCl₂ or KCl+Aq, solubility increasing with strength of solution; sol. in NaCl or CaCl₂+Aq, maximum solubility occurring when the solutions are of a medium concentration. The numerical results are as follows:

100 pts. of the salt solutions containing given
 pts. salt dissolve pts. SrSO₄.

Salt	Pts. salt	Pts. SrSO ₄
NaCl	22.17	0.1811
	15.54	0.2186
	8.44	0.1653
KCl	18.08	0.2513
	12.54	0.1933
	8.22	0.1925
MgCl ₂	13.63	0.2419
	4.03	0.2057
	1.59	0.1986
CaCl ₂	33.70	0.1706
	16.51	0.1853
	8.67	0.1756

(Virck, C. C. 1862. 402.)

Solubility in H₂O, and in solutions of the sulphates chlorides and nitrates of the alkalies and alkaline earths and in solutions of salts of the alkalies with strong organic acids has been determined. No data in abstract. (Wolfmann, Chem. Soc. 1898, (2) 74. 220.)
 Solubility in H₂O is considerably decreased by the presence of K₂SO₄. (Barre, A. ch. 1911, (8) 24. 175.)

Solubility of SrSO₄ in Ca(NO₃)₂+Aq at ord. temp.

G. per 100 cc. sat. solution.

Ca(NO ₃) ₂	SrSO ₄	Ca(NO ₃) ₂	SrSO ₄
0.5	0.0483	4	0.1489
1	0.0619	5	0.1689
2	0.1081	6	0.1955
3	0.1275

(Raffo and Rossi, Gazz. ch. it. 1915, 45. (1) 45.)

Insol. in liquid NH₃. (Franklin, Am. Ch J. 1898, 20. 829.)
 100 g. 95% formic acid dissolve 0.02 g. SrSO₄ at 18.5°. (Aschan, Chem. Ztg. 1913, 37. 1117.)
 Insol. in absolute alcohol; scarcely sol. in dil. alcohol.
 Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790); acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)
 Min. *Celestite*.

Strontium hydrogen sulphate, SrH₂(SO₄)₂

100 pts. H₂SO₄ dissolve 2.2 pts. SrSO₄ (Lec-Bodart and Jacquemin); 100 pts. H₂SO₄ dissolve 5.68 pts. (Struve, Z. anal. 9. 34); 100 pts. fuming H₂SO₄ dissolve 9.77 pts (Struve).
 1 g. SrSO₄ dissolves in 1256 g. 91% H₂SO₄+Aq (Varenne and Pauleau, C. R. 93. 1016; boiling H₂SO₄ dissolves about 15% SrSO₄ and still more at 100°. (Schultz, Pogg. 133. 147
 Sol. in 1519 pts. 91% H₂SO₄. (Varenne and Pauleau, C. R. 93. 1016.)
 100 pts. H₂SO₄ (sp. gr. 1.843) dissolve 14 pts. SrSO₄ at 70°. (Garside, C. N. 31. 245
 Decomp. by H₂O.
 100 pts. hot conc. H₂SO₄ dissolve about 9.0 pts. SrSO₄. (Rohland, Z. anorg. 1910, 66. 206.)
 10 ccm. of sat. SrSO₄+absolute H₂SO₄ contain approx. 2.17 g. SrSO₄. (Bergius, Z. phys. Ch. 1910, 72. 355.)
 +H₂O. Decomp. by H₂O.

Strontium tin (stannic) sulphate, SrSO₄. Sn(SO₄)₂+3H₂O.

Decomp. by H₂O. Sol. in HCl. (Weinland and Kühl, Z. anorg. 1907, 54. 249.)

Strontium titanium sulphate, SrSO₄. Ti(SO₄)₂

Ppt; decomp. by H₂O giving titanil acid (Weinland and Kühl, Z. anorg. 1907, 54. 254.)

Tantalum sulphate, 3Ta₂O₅. SO₃+9H₂O. (Hermann, J. pr. 70. 201.)

Tellurium sulphate, basic, TeO₂. SO₃.

Sol. in cold dil. H₂SO₄. Decomp. by hot H₂O. (Klein, C. R. 99. 326.)

Terbium sulphate, Tr₂(SO₄)₃+8H₂O.

Sol. in H₂O.
 Sol. in H₂O; pptd. by alcohol. (Urthain, C. R. 1908, 146. 127.)

Thallous sulphate, Tl₂SO₄.

1 pt. dissolves at t° in pta. H₂O, according to C=Crookes; L=Lamy:
 15° 18° 62° 100° 101.2°
 21.1 20.8 8.7 5.4 5.22 pta. H₂O.
 C L L C L

Solubility of Th_2SO_4 in H_2O .

t°	% Th_2SO_4	t°	% Th_2SO_4
0	2.63	60	9.85
10	3.57	70	11.31
20	4.64	80	12.75
30	5.80	90	14.19
40	7.06	99.7*	15.57
50	8.44		

B.-pt. at 748 mm.
Berkeley, Phil. Trans. Roy. Soc. 1904, **203**.
1, 189; calc. by Landolt-Börnstein.)

100 g. H_2O dissolve 3.36 g. Th_2SO_4 at 6.5° ;
3 g. at 12° ; 19.14 g. at 100° . (Tutton, Proc.
Roy. Soc. 1907, **79**, A, 351.)

1 l. H_2O dissolves 0.1928 equivalents
 Th_2SO_4 at 20° , or 48.59 g. in 1 l. of solution.
Noyes, J. Am. Chem. Soc. 1911, **33**, 1657.)

1 l. H_2O dissolves 0.1083 g. equiv. Th_2SO_4
at 25° , or 27.28 g. in 1 l. of solution. (Noyes.)

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

Strength of $\text{H}_2\text{SO}_4 + \text{Aq}$ g. mols. per l.	g. mols. Th_2SO_4 per l.
0.0494	0.1172
0.0987	0.1249

(Noyes, J. Am. Chem. Soc. 1911, **33**, 1662.)

1 l. $\text{ThClO}_4 + \text{Aq}$. containing 0.1058 equiv-
alents ThClO_4 , dissolves 0.1366 equivalents
 Th_2SO_4 at 20° . (Noyes.)

See also ThClO_4 .

Solubility in salts + Aq at 25° .

g. mols. per l.	g. mols. per l. Th_2SO_4 dissolved
0.0996 TiNO_3	0.08365
0.0497 Na_2SO_4	0.1080
0.1988 Na_2SO_4	0.1173
0.1010 NaHSO_4	0.1161

(Noyes.)

Thallous hydrogen sulphate, ThHSO_4 .

Sl. hygroscopic.

Solubility in $\text{H}_2\text{SO}_4 + \text{Aq}$ at 25° .

1000 g. of the solution contain:

Mols. H_2SO_4	Mols. Th_2SO_4
4.55	0.56
4.79	0.55
4.89	0.59
4.92	0.66
4.78	0.75
4.26	1.01
4.03	1.08

(D'Ans, Z. anorg. 1910, **65**, 232.)

Th_2SO_4 , ThHSO_4 . Sol. in H_2O . (Storten-
becker, R. t. c. 1902, **21**, 90.)

Thallous pyrosulphate, $\text{Th}_2\text{S}_2\text{O}_7$.

Decomp. by H_2O . (Weber, B. 17, 2502.)

Thallous octosulphate, $\text{Th}_2\text{S}_8\text{O}_{32}$.

Decomp. by H_2O . (Weber, B. 17, 2502.)

Thallic sulphate, basic, $\text{Th}_2\text{O}_3 \cdot 2\text{SO}_3 + 3\text{H}_2\text{O}$.

Sol. in H_2O .

+ $5\text{H}_2\text{O}$. As above. (Willm, A. ch. (4) 5.
5.)

Thallic sulphate, $\text{Th}_2(\text{SO}_4)_3 + 7\text{H}_2\text{O}$.

Decomp. by cold H_2O with separation of
 $\text{ThO}(\text{OH})$. (Crookes.)

Thallothallic sulphate, $2\text{Th}_2\text{O} \cdot 3\text{Th}_2\text{O}_3 \cdot 12\text{SO}_3 + 25\text{H}_2\text{O}$.

Gradually efflorescent. (Willm.)

$\text{Th}_2(\text{SO}_4)_2$. (Lepsius, Chem. Ztg. 1890.
1327.)

$\text{ThH}(\text{SO}_4)_2$. (Lepsius.)

Thallous uranyl sulphate, $\text{Th}_2(\text{UO}_2)(\text{SO}_4)_2 + 3\text{H}_2\text{O}$.

Sl. sol. in cold H_2O .

Easily forms supersat. solutions. (Kohn,
Z. anorg. 1908, **59**, 112.)

Thallium vanadium sulphate, $\text{Th}_2\text{V}_2(\text{SO}_4)_4 + 24\text{H}_2\text{O}$.

100 pts. H_2O dissolve 11.06 pts. salt at 10° .

Sp. gr. of solution at $4^\circ/20^\circ = 2.342$.

Very sol. in hot H_2O . (Piccini, Z. anorg.
1897, **13**, 446.)

256 g. anhydrous, or 433 g. hydrated salt,
or 0.573 g. mols. of anhydrous salt are sol. in
1 l. H_2O at 25° .

Melts in crystal H_2O at 48° . (Locke, Am.
Ch. J. 1901, **26**, 175.)

Thallous zinc sulphate, $\text{Th}_2\text{Zn}(\text{SO}_4)_2 + 6\text{H}_2\text{O}$.

1 l. H_2O dissolves 86 g. anhydrous salt
at 25° . (Locke, Am. Ch. J. 1902, **27**, 459.)

Thallothallic sulphate bromide, $\text{Th}_2\text{Br}_2\text{SO}_4$.

Very sol. in cold H_2O . Decomp. by much
 H_2O . (Meyer and Goldschmidt, B. 1903, **36**.
242.)

Thorium sulphate, basic, $3[\text{Th}(\text{SO}_4)_2 + 2\text{H}_2\text{O}]$,
 $\text{Th}(\text{SO}_4)_2\text{O} + 2\text{H}_2\text{O}$.

Insol. in H_2O ; very slowly attacked by dil.
acids. (Demarcay.)

$\text{ThO}(\text{SO}_4) + 2\text{H}_2\text{O}$. Stable in aq. solution
at 100° . (Hauser, B. 1910, **43**, 2776.)

+ $5\text{H}_2\text{O}$. Somewhat sol. in hot conc.
 $\text{MgSO}_4 + \text{Aq}$. (Halla, Z. anorg. 1912, **79**, 260.)

Thorium sulphate, $\text{Th}(\text{SO}_4)_2$.

Anhydrous. Easily sol. if brought into a
large amount of H_2O , but very slowly sol. if
only a little H_2O is added to the salt.

Titanyl sulphate, (TiO)SO₄.

Decomp. by H₂O. Slowly sol. in cold, rapidly in warm HCl+Aq. (Mers, J. pr. 99. 157.)

+2H₂O. Sol. in H₂O. (Blondel, Bull. Soc. 1899, (3) 21. 262.)

2TiO₂, 3SO₃+3H₂O. Sol. in H₂O acidified with HCl. (Blondel, Bull. Soc. 1899, (3) 21. 262.)

5TiO₂, SO₃+5H₂O. (Blondel.)

7TiO₂, 2SC₂+7H₂O. (Blondel.)

2TiO₂, SO₃+2H₂O. (Blondel.)

Uranous sulphate, basic, U(OH)₂SO₄+H₂O.

Insol. in H₂O. H₂O dissolves out H₂SO₄. (Ebelmen, A. ch. (3) 5. 217.)

+5H₂O. Sol. in alcohol.

Pptd. by ether. (Rosenheim, Z. anorg. 1901, 26. 251.)

Uranous sulphate, U(SO₄)₂+4H₂O.

Sol. in H₂O with immediate decomp. Easily sol. in dil. H₂SO₄+Aq. (Kohl-schütter, B. 1901, 34. 3629.)

Decomp. by H₂O into insol. basic, and sol. acid salt. Sol. in dil. H₂SO₄ or HCl+Aq. Difficultly sol. in conc. acids. (Ebelmen, A. ch. (3) 5. 215.)

Solubility of U(SO₄)₂+4H₂O in H₂O at t°.

t°	% U(SO ₄) ₂	t°	% U(SO ₄) ₂
29	9.8	48.2	8.1
37	8.3	63	7.3

(Giolitti and Bucci, Gazz. ch. it. 1905, 35. (2) 162.)

1 pt. is sol. in 4.23 pts. H₂O at 13°; 4.3 pts. at 11.3°; 4.4 pts. at 9.1°. (de Coninck, A. ch. 1903, (7) 28. 12.)

+8H₂O.

Solubility of U(SO₄)₂+8H₂O in H₂O at t°.

t°	% U(SO ₄) ₂	t°	% U(SO ₄) ₂
18	10.17	48.2	28.72
25.6	13.32	62	36.8
37	19.98	93	63.2

(Giolitti and Bucci, Gazz. ch. it. 1905, 35. (2) 162.)

Sp. gr. of U(SO₄)₂+Aq at t°.

t°	% salt	Sp. gr.
16	1	1.0058
16.8	2	1.0107
16	3	1.0165
17.8	4	1.0218
17.2	5	1.0272
18	6	1.0320
18.3	7	1.0379
17.4	8	1.0429
15.2	9	1.0485
15.6	10	1.0539

(de Coninck, A. ch. 1903, (7) 28. 11.)

Solubility in acids+Aq.

1 pt. U(SO₄)₂ is sol. in pts. acid at t°

t°	Acid	Concentration of acid	Pts. acid
9.7 9.2	HCl	1:4	5.74 5.8
11.2 10.3	HNO ₃	1:4	5.4 5.33
11.4 10.7	H ₂ SeO ₄ (sp. gr. 1.4)	1:4	4.57 4.66
15 14.2	HBr	1:4	4 4.23
15.5 14.4	HC ₂ H ₃ O ₂	1:4.5	4.1 4.3
16.5 15.9	"	1:2	3.72 3.85
11.7 10.9 10.1 9	H ₂ SO ₄	1:4	6.36 6.42 6.45 6.5

(de Coninck, Chem. Soc. 1902, 82. (2) 439)

Sp. gr. of U(SO₄)₂+HCl (d = 1.046) at t°

d₁ = Sp. gr. referred to H₂O.

d₂ = Sp. gr. referred to HCl

t°	% salt	d ₁	d ₂
16	1	1.0525	1.0063
17	2	1.0572	1.0109
18	3	1.0619	1.0154
18.4	4	1.0667	1.0199
17.6	5	1.0714	1.0243

(de Coninck, A. ch. 1903, (7) 28. 11.)

Sp. gr. of U(SO₄)₂+H₂SO₄ (d = 1.14) at t°

d₁ = Sp. gr. referred to H₂O.

d₂ = Sp. gr. referred to H₂SO₄.

t°	% salt	d ₁	d ₂
18.7	1	1.1442	1.0038
18.3	2	1.1494	1.0063
17.4	3	1.1539	1.0123
17.6	4	1.1583	1.0162
18.1	5	1.1626	1.0204

(de Coninck, A. ch. 1903, (7) 28. 11.)

1 pt. is sol. in 8 pts. alcohol (94°)+Aq (1:4) at 10.4°. (de Coninck.)

Solubility in glycol at 14.8° = 3.15% (de Coninck, C. C. 1906, II. 883.)

Min. Johannite. Sl. sol. in H₂C.

Uranous hydrogen sulphate, U(SO₄)₂. H₂SO₄. +10H₂O.

(Giolitti, C. C. 1906, II. 1226.)

Uranyl sulphate, basic, $3\text{UO}_2, \text{SO}_3 + 2\text{H}_2\text{O}$.

(Athanasesco.)

+14H₂O. Sol. in H₂O. (Ordway, Sill. Am. J. (2) **26**. 208.)4UO₂, SO₃ + 7H₂O. (Athanasesco, C. R. **103**. 271.)UO₂, 4UOSO₄ + 8H₂O. Less sol. in min. acids, especially dil. H₂SO₄ + Aq, than UOSO₄ + 2H₂O. (de Coninck, C. C. **1901**, II. 1038.)**Uranyl sulphate, (UO₂)SO₄.**+H₂O. (de Coninck.)+3H₂O. Efflorescent. Very sol. in H₂O and alcohol.1 pt. is sol. in 0.6 pt. cold H₂O; in 0.45 pt. boiling H₂O; in 25 pts. cold absolute alcohol; in 20 pts. boiling absolute alcohol. (Bucholz.)Sol. in 0.47 pt. H₂O at 21°, and 0.28 pt. boiling H₂O. (Ebelmen.)100 pts. H₂O at 15.5° dissolve 160 pts., and at 100°, 220 pts. (Ure's Dict.)1 pt. is sol. in 5.3 pts. H₂O at 13.2°; 5.16 pts. at 14.1°; 4.96 pts. at 15.1°; 4.88 pts. at 15.5°. (de Coninck, A. ch. **1903**, (7) **28**. 8.)Sp. gr. of (UO₂)SO₄ + Aq at t°.

t°	% (UO ₂)SO ₄	Sp. gr.
14	1	1.0062
15.5	2	1.0113
11.3	3	1.0172
10.2	4	1.0229
10.2	5	1.0280
10	6	1.0338
14	7	1.0389
15.6	8	1.0442
11	9	1.0503
10.3	10	1.0557
11.4	11	1.0612
11.6	12	1.0669

(de Coninck, A. ch. **1903**, (7) **28**. 7.)

Solubility in acids + Aq.

1 pt. (UO₂)SO₄ is sol. in:

3.4 pts. conc. HCl at 12.8°

3.25 " " " " 13.6°

5.9 " HBr(d=1.21) " 12.9°

6.1 " " " " 11.2°

10.8 " conc. HNO₃ " 12.3°

11.2 " " " " 10.8°

4.3 " H₂SO₄(d=1.38) " 12.7°

4.1 " " " " 14.0°

5.6 " aqua regia (equal vol. HCl + HNO₃) at 15.4°5.47 pts. aqua regia (equal vol. HCl + HNO₃) at 16.4°

3.7 pts. selenic acid (d=1.4) at 15.3°.

(de Coninck, A. ch. **1903**, (7) **28**. 8.)Sp. gr. of (UO₂)SO₄ + H₂SO₄(d=1.168) at t°.d₁ = Sp. gr. referred to H₂O.d₂ = Sp. gr. referred to H₂SO₄.

t°	% salt	d ₁	d ₂
20.6	1	1.1738	1.0050
22.2	2	1.1775	1.0082
21.1	3	1.1880	1.0129
22.7	4	1.1872	1.0165
22.3	5	1.1918	1.0204

(de Coninck, A. ch. **1903**, (7) **28**. 7.)1 pt. is sol. in 37.9 pts. alcohol (85°) at 16.7°; 38.6 pts. at 15.8°. (de Coninck, A. ch. **1903**, (7) **28**. 8.)Very sl. sol. in formic and glacial acetic acids. (de Coninck, A. ch. **1903**, (7) **28**. 9.)Completely pptd. from (UO₂)SO₄ + Aq by HC₂H₃O₂. (Persoz.)**Uranyl hydrogen sulphate, (UO₂)SO₄, H₂SO₄.**

Very deliquescent. (Schultz-Sellack.)

2(UO₂)SO₄, H₂SO₄ + 5H₂O. Very deliquescent. (Wyruboff, Bull. Soc. Min. **1909**, **32**. 351.)**Uranyl pyrosulphate, (UO₂)S₂O₇.**Very deliquescent. Hisses with H₂O. (Schultz-Sellack.)**Uranouranyl sulphate, USO₄, (UO₂)SO₄.**Sol. in H₂O. (Ebelmen.) Decomp. by boiling. (Berzelius.)Min. *Voglianite*.**Uranyl sulphate ammonia, (UO₂)SO₄, 2NH₃.**(v. Unruh, Dissert. **1909**.)(UO₂)SO₄, 3NH₃. (v. Unruh.)(UO₂)SO₄, 4NH₃. (v. Unruh.)**Vanadous sulphate, V₂O₃, 4SO₃ + 9H₂O.**Sol. in H₂O. (Brierley, Chem. Soc. **49**. 882.)**Vanadium sulphate, V₂O₅, 2SO₃ = (VO₂)₂S₂O₇.**Deliquescent. Easily sol. in H₂O.V₂O₅, 3SO₃. Deliquescent. Sol. in H₂O and alcohol.+3H₂O. Deliquescent. Very sol. in H₂O, but decomp. by boiling. Sol. in alcohol. (Ditte, C. R. **102**. 757.)VSO₄ + 7H₂O. Decomp. by air; very unstable; sol. in H₂O. (Piccini, Z. anorg. **1899**, **19**. 204.)**Vanadium sesquisulphate, V₂(SO₄)₃.***Anhydrous.*Insol. in H₂O.Insol. in conc. H₂SO₄, but slowly sol. in boiling dil. H₂SO₄. Sol. in HCl.Insol. in alcohol and ether. (Stähler, B. **1905**, **38**. 3979.)

Vanadium hydrogen sulphate,Sol. in H_2O .Sol. in HCl . Insol. in 60% $H_2SO_4 + Aq$, alcohol, ether and acetic acid. (Stähler, B. 1905, **38**. 3978.)**Divanadyl sulphate, $V_2O_5(SO_4)_2$.**Insol. in H_2O , HCl , or $H_2SO_4 + Aq$, but on heating to 400° becomes sol. in H_2O if heated to 130° therewith. (Gerland.)+ $4H_2O$. Very slowly sol. in H_2O at 10° , quickly at 60° , and still more rapidly at 100° . Deliquesces in warm moist air more quickly than it dissolves in H_2O at 10° . Insol. in absolute alcohol. Very sol. in alcohol of 0.833 sp. gr. (Berzelius.)+ $5H_2O$. (Koppel and Behrendt, Z. anorg. 1903, **35**. 168.)+ $7H_2O$, and $10H_2O$.+ $13H_2O$. Efflorescent. (Gerland.) $2V_2O_5, 5SO_3 + 18H_2O$. (Gain, C. R. 1906, **143**. 1154.) $V_2O_5, 3SO_3 + 10H_2O$. (G.) $2V_2O_5, 7SO_3 + 20H_2O$. (G.) $2V_2O_5, 9SO_3 + 22H_2O$. (G.) $V_2O_5, 5SO_3 + 12H_2O$. (G.)**Divanadyl hydrogen sulphate,**+ $2H_2O$.+ $3H_2O$. Deliquescent. Very slowly sol. in cold H_2O or alcohol. Easily sol. in hot H_2O . (Gerland.)+ $5H_2O$. Deliquescent. Insol. in ether. Scarcely sol. in alcohol. Slowly sol. in cold, easily in hot H_2O . (Crow.)+ $14H_2O$. Easily sol. in cold H_2O or dil. alcohol. (Gerland.) $2VO_3, 3SO_3$. Sl. sol. in H_2O . (Koppell and Behrendt, Z. anorg. 1903, **35**. 163.) $2VOSO_4, H_2SO_4 + H_2O$. Very slowly sol. in H_2O . (Koppel and Behrendt, Z. anorg. 1903, **35**. 163.) $2VOSO_4, 3H_2SO_4 + 15H_2O$. (Gain, C. R. 1906, **143**. 1156.) $2VOSO_4, 4H_2SO_4 + 16H_2O$. (G.) $2VOSO_4, 5H_2SO_4 + 15H_2O$. (G.) $2VOSO_4, 7H_2SO_4 + 15H_2O$. (G.) $2VOSO_4, 8H_2SO_4 + 16H_2O$. (G.)**Ytterbium sulphate, $Yb_2(SO_4)_3 + 8H_2O$.**Quite slowly sol. in H_2O even at 100° . Anhydrous salt is easily sol. in much H_2O , but if little H_2O is used the hydrous salt is formed, which only slowly dissolves. Sol. in $K_2SO_4 + Aq$.100 pts. H_2O dissolve at:

0°	15.5°	35°	55°	60°
44.2	34.6	19.1	11.5	10.4

pts. $Yb_2(SO_4)_3$,

70°	80°	90°	100°
7.22	6.93	5.83	4.67

pts. $Yb_2(SO_4)_3$.

(Cleve, Z. anorg. 1902, **32**. 143.)**Yttrium sulphate, basic, $Y_2O_3, SO_3 = (YO)_2SO_4$.**Insol. in H_2O . (Berzelius.) $2Y_2O_3, SO_3 + 10H_2O$. (Cleve.)**Yttrium sulphate, $Y_2(SO_4)_3$.***Anhydrous*. More sol. in H_2O than the hydrous salt, and more sol. in cold than hot H_2O . Solution sat. at 0° separates $Y_2(SO_4)_3 + 8H_2O$ at 50° . 100 pts. H_2O dissolve 15.2 pts. anhydrous salt. at ord. temp.5.38 pts. are sol. in 100 pts. H_2O at 25° . (James and Holden, J. Am. Chem. Soc. 1913, **35**. 561.)Easily sol. in large amount of sat. $K_2SO_4 + Aq$, from which $3K_2SO_4, 2Y(SO_4)_3$ is pptd. on warming. (Cleve and Höglund, Sv. V. A. H. Bih. 1. No. 8.)Solubility of $Y_2(SO_4)_3$ in $Na_2SO_4 + Aq$ at 25° .

Pts. $Y_2(SO_4)_3$ per 100 pts. H_2O	Pts. Na_2SO_4 per 100 pts. H_2O	Solid phase
5.61	1.29	$Y_2(SO_4)_3$
6.38	3.85	
7.40	6.21	
8.43	8.53	
5.86	7.57	
4.75	7.72	
3.42	10.14	
2.36	11.36	
2.02	13.42	
1.90	14.89	
1.79	16.51	$Y_2(SO_4)_3, Na_2SO_4 + 2H_2O$
1.86	18.44	
2.99	19.96	
3.04	21.05	
2.27	27.14	
1.52	28.22	
1.61	28.13	$Na_2SO_4 + 10H_2O$
5.38	0.0	

(James and Holden, J. Am. Chem. Soc. 1913, **35**. 560.)+ $8H_2O$. 100 pts. H_2O dissolve 9.3 pts. of cryst. salt at ord. temp., and 4.8 pts. at 100° . (Cleve, Bull. Soc. (2) **21**. 344.)Less sol. in H_2O containing H_2SO_4 than in pure H_2O . (Berzelius.)Completely pptd. by $HC_2H_3O_2 + Aq$. Insol. in alcohol.**Yttrium hydrogen sulphate, $Y_2(SO_4H)_3$.**(Brauner, Z. anorg. 1904, **33**. 332.)**Zinc sulphate, basic, $8ZnO, SO_3 + 2H_2O$.**Insol. in H_2O . (Schindler, Mag. Pharm. **31**. 181.) $6ZnO, SO_3 + 10H_2O$. Insol. in H_2O . (Kane, A. ch. **72**. 310.) $4ZnO, SO_3 + 2H_2O$. Scarcely sol. in hot or cold H_2O . Sol. in $ZnSO_4 + Aq$. (Kühn, Schw. J. **60**. 337.)

+3H₂O. (Werner, B. 1907, 40. 4443.)
 +5H₂O. Nearly insol. in H₂O. (Habermann, M. 5. 432.)
 +6H₂O. (Kraut, Z. anorg. 1897, 13. 5.)
 +7H₂O. (Athanasesco, C. R. 103. 271.)
 +8H₂O. Extremely slowly decomp. by H₂O. (Reindel, J. pr. 1869, (1) 103. 373.)
 +10H₂O. (Schindler.)
 3ZnO, SO₃. Insol. in cold, sl. sol. in hot H₂O. (Vogel.)
 2ZnO, SO₃. (Athanasesco.)
 5ZnO, SO₃. (Pickering, Chem. Soc. 1907, 91. 1986.)
 +4H₂O. (Moody, Am. J. Sci. 1906; [4] 22. 184.)
 9ZnO, 2SO₃+12H₂O. (Reindel, J. pr. 1869, (1) 103. 374.)

Zinc sulphate, ZnSO₄.

Sol. in H₂O with evolution of heat.
 Sol. in HCl+Aq.
 +H₂O. (Étard.)
 +2H₂O. Insol. in alcohol. (Kühn.)
 +3½H₂O. (Anthon.)
 +5H₂O. Insol. in boiling alcohol of 0.86 sp. gr. (Kühn.)
 +6H₂O. (Marignac.)
 +7H₂O. Slowly efflorescent.
 M.-pt. of ZnSO₄+7H₂O = 50°. (Tilden, Chem. Soc. 45. 409.)
 For solubility data on hydrated salts, see below.

Sol. in 2 + pts. H₂O at ord. temp., and in less at 100°. (Bergmann.)
 100 pts. H₂O at 104.4° dissolve 81.81 pts. ZnSO₄. (Griffiths.)
 100 pts. H₂O at ord. temp. dissolve 140 pts. ZnSO₄+7H₂O. (Dumas.)
 Sol. in 2.29 pts. H₂O at 18.75°. (Abl.)
 100 pts. H₂O at 15.56° dissolve 140 pts. ZnSO₄+7H₂O. (Ure's Dict.)
 100 pts. H₂O at 15° dissolve 140.53 pts. ZnSO₄+7H₂O, and has sp. gr. = 1.4442. (Michel and Krafft.)

1 pt. of the crystals dissolves in 0.923 pt. H₂O at 17.5°, and forms a solution of 1.4353 sp. gr. (Karsten.)

100 pts. ZnSO₄+Aq sat. at 18-20° contain 35.36 pts. ZnSO₄. (v. Hauer, J. B. 1866. 59.)

100 pts. H₂O dissolve at:
 0° 20° 50° 75°
 41.3 53.0 66.9 80.4 pts. ZnSO₄.
 (Tobler, J. B. 1855. 309.)

100 pts. H₂O at 20.5° dissolve 163.2 pts. ZnSO₄+7H₂O. (Schiff, A. 109. 336.)

100 pts. H₂O at t° dissolve pts. anhydrous ZnSO₄, and pts. ZnSO₄+7H₂O.

t°	Pts. ZnSO ₄	Pts. ZnSO ₄ +7H ₂ O	t°	Pts. ZnSO ₄	Pts. ZnSO ₄ +7H ₂ O
0	43.02	115.22	60	74.20	313.48
10	48.36	138.21	70	79.25	369.36
20	53.13	161.49	80	84.60	442.62
30	58.40	190.90	90	89.78	533.02
40	63.52	224.05	100	95.03	653.59
50	68.75	263.84

(Poggiale, A. ch. (3) 8. 467.)

Solubility of ZnSO₄ in 100 pts. H₂O at t°.

t°	Pts. ZnSO ₄	t°	Pts. ZnSO ₄	t°	Pts. ZnSO ₄
0	44.0	14	52.8	27	62.1
1	44.6	15	53.5	28	62.8
2	45.2	16	54.2	29	63.6
3	45.8	17	54.9	30	64.3
4	46.4	18	55.6	31	65.1
5	47.0	19	56.3	32	65.8
6	47.6	20	57.0	33	66.6
7	48.3	21	57.7	34	67.3
8	48.9	22	58.4	35	68.1
9	49.5	23	59.2	36	78.8
10	50.2	24	59.9	37	69.3
11	50.8	25	60.7	38	70.4
12	51.5	26	61.4	39	71.2
13	52.2

Decomp. into basic salt above 40°.

(Mulder, Scheik. Verhandel. 1864. 74.)

If solubility S represents number of pts. anhydrous salt in 100 pts. of solution, S = 27.6+0.2604t from -5° to +81°; S = 50.0-0.2244t from 81° to 175°. (Étard, C. R. 103. 207.)

Sat. ZnSO₄+Aq contains at:

1°	13°	20°	41°	49°
29.1	32.6	34.8	40.2	40.9% ZnSO ₄ ,
55°	62°	70°	77°	100°
43.4	45.0	47.0	46.5	44.7% ZnSO ₄ ,
111°	12.°	137°	144°	169° 171°
43.0	40.7	38.0	37.4	30.0 29.0% ZnSO ₄ .

(Étard, A. ch. 1894, (7) 2. 551.)

Transition point from +6H₂O to +1H₂O is 70°. (Étard.)

Solubility of ZnSO₄+6H₂O in H₂O at t°.

t°	g. ZnSO ₄ in 100 g. H ₂ O
-5.0	47.08
+0.1	49.48
9.1	54.20
15.0	57.15
25.0	63.74
30.0	65.82
35.0	67.99
39.0	70.08

(Cohen, Z. phys. Ch. 1900, 34. 182.)

Solubility of the hepta- and hexa-hydrates of ZnSO_4 at t° .

p = wt. of salt expressed in percent of solution.

t°	p
0	29.43
0	29.53
0	29.49
15.00	33.66
15.88	33.85
30.70	38.46
39.92	41.36
39.95	41.37
40.73	41.43
41.49	41.70
46.40	42.68
49.97	43.51
49.99	43.41
50.00	43.50
50.02	43.51

Transition point from $+7\text{H}_2\text{O}$ to $+6\text{H}_2\text{O}$ is 39° .

The formula representing the change of solubility between 0° and 39° is

$p = 29.5 + 0.270t + 0.00068t^2$
while the expression for the hexahydrate above 40° is

$$p = 41.35 + 0.210t + 0.00070t^2.$$

(Barnes, J. phys. Chem. 1900, 4, 19.)

Solubility of $\text{ZnSO}_4 + 7\text{H}_2\text{O}$ in H_2O at t° .

t°	g. ZnSO_4 in 100 g. H_2O
-5	39.30
+0.1	41.93
9.1	47.09
15.0	50.88
25.0	57.90
35.0	66.61
39.0	70.05

(Cohen, Z. phys. Ch. 1900, 34, 182.)

Solubility in H_2O at high pressures:

Pressure in atm	t°	g. ZnSO_4 in 100 g. H_2O	Solubility at 25°
1	26	57.95	57.95
500	25.8	58.43	57.92
1000	25.8	58.32	57.91
1000	25.8	57.95	57.55

(Cohen and Sinnige, Z. phys. Ch. 1909, 67, 444.)

Liabile to form supersaturated solutions.

Sat. $\text{ZnSO}_4 + \text{Aq}$ at 8° has sp. gr. (Anthon.)

Sp. gr. of $\text{ZnSO}_4 + 7\text{H}_2\text{O}$ at 20°

$$\% = \% \text{ZnSO}_4 + 7\text{H}_2\text{O}$$

%	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.0057	21	1.1288	41	1.258
2	1.0115	22	1.1355	42	1.268
3	1.0173	23	1.1423	43	1.275
4	1.0231	24	1.1491	44	1.280
5	1.0289	25	1.1560	45	1.285
6	1.0348	26	1.1629	46	1.290
7	1.0407	27	1.1699	47	1.295
8	1.0467	28	1.1770	48	1.300
9	1.0527	29	1.1842	49	1.305
10	1.0588	30	1.1914	50	1.311
11	1.0649	31	1.1987	51	1.316
12	1.0710	32	1.2060	52	1.320
13	1.0772	33	1.2134	53	1.325
14	1.0835	34	1.2209	54	1.330
15	1.0899	35	1.2285	55	1.335
16	1.0962	36	1.2362	56	1.340
17	1.1026	37	1.2439	57	1.345
18	1.1091	38	1.2517	58	1.350
19	1.1156	39	1.2595	59	1.355
20	1.1222	40	1.2674	60	1.360

(Schiff, A. 110, 72.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$ at 15° .

$$\% = \% \text{ZnSO}_4 + 7\text{H}_2\text{O}.$$

%	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.006	21	1.130	41	1.280
2	1.013	22	1.137	42	1.288
3	1.019	23	1.143	43	1.295
4	1.024	24	1.150	44	1.304
5	1.0288	25	1.1574	45	1.3100
6	1.035	26	1.164	46	1.320
7	1.041	27	1.171	47	1.330
8	1.047	28	1.179	48	1.337
9	1.053	29	1.185	49	1.346
10	1.0593	30	1.1933	50	1.3532
11	1.066	31	1.200	51	1.362
12	1.073	32	1.209	52	1.380
13	1.079	33	1.216	53	1.370
14	1.085	34	1.224	54	1.390
15	1.0905	35	1.231	55	1.3966
16	1.097	36	1.240	56	1.406
17	1.103	37	1.246	57	1.416
18	1.110	38	1.255	58	1.425
19	1.116	39	1.263	59	1.435
20	1.1236	40	1.2709	60	1.4451

(Gerlach, Z. anal. 8, 268.)

r. of $\text{ZnSO}_4 + \text{Aq}$ at 23.5° . a=no. of g., equivalent to $\frac{1}{2}$ mol. wt., dissolved in 1000 g. H_2O ; b=sp. gr. if a is $\text{ZnSO}_4 + 7\text{H}_2\text{O}$, $\frac{1}{2}$ mol. wt. = 143.5; c=sp. gr. if a is ZnSO_4 , $\frac{1}{2}$ mol. wt. = 80.5.

b	c	a	b
1.077	1.084	7	1.368
1.143	1.162	8	1.400
1.199	1.236	9	1.428
1.249	1.307	10	1.453
1.294	1.376	11	1.476
1.333	1.443

(Favre and Valson, C. R. 79. 968.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$ at 18° .

Sp. gr.	% ZnSO_4	Sp. gr.	% ZnSO_4	Sp. gr.
1.0509	15	1.1675	32	1.3045
1.1369	20	1.2313	50	1.3788

(Kohlrausch, W. Ann. 1879. 1.)

p. gr. of $\text{ZnSO}_4 + \text{Aq}$ at room temp. con-
ing:

7.12 16.64 23.09% ZnSO_4 .
1.1064 1.1953 1.2814

(Wagner, W. Ann. 1883, 18. 271.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$ at 19.5° .

Mass of salt per unit mass of solution	Density of solution. (g. per cc.)
0.00186	1.00179
0.00371	1.00356
0.00556	1.00530
0.00740	1.00711
0.01106	1.01065
0.01469	1.01410
0.01829	1.01753
0.02187	1.02112
0.02542	1.02446
0.02895	1.02798

(McGregor, C. N. 1887, 55. 4.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$ at 25° .

Concentration of $\text{ZnSO}_4 + \text{Aq}$	Sp. gr.
1—normal	1.0792
$\frac{1}{2}$ —"	1.0402
$\frac{1}{4}$ —"	1.0198
$\frac{1}{8}$ —"	1.0094
$\frac{1}{16}$ —"	1.0047

(Wagner, Z. phys. Ch. 1890, 5. 40.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$.

g. salt in 100 g. solution	Sp. gr.
24.7170	1.3152
21.4444	1.2665
17.7573	1.2145
14.0307	1.1645
9.7426	1.1106
5.1110	1.0565

(Charpy, A. ch. 1893, (6) 29. 27.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$.

g. equivalents ZnSO_4 per liter	t°	Sp. gr. t°/t°
0.001309	13.59	1.0001126
0.002616	13.575	1.0002258
0.005212	13.573	1.0004451
0.01039	13.585	1.000886
0.09818	13.621	1.008189
0.18842	13.642	1.015587
0.1890	16.11	1.01550
2.493	15.88	1.19385

(Kohlrausch and Hallwachs, W. Ann. 1894, 53. 27.)

Sp. gr. of $\text{ZnSO}_4 + \text{Aq}$ at 18.5° , when p =
percent strength of solution; d = ob-
served density; and w = volume conc.
in grams per cc. ($\frac{pd}{100} = w$)

p	d	w
29.22	1.3718	0.40057
25.14	1.3091	0.32910
21.28	1.2528	0.26659
17.08	1.1957	0.20422
11.20	1.1220	0.12567
8.44	1.0894	0.09195
6.65	1.0696	0.07112
3.82	1.0387	0.03968
3.18	1.0318	0.03281
1.46	1.0138	0.01480
0.577	1.0045	0.00580

(Barnes, J. phys. Chem. 1898, 2. 542.)

Sp. gr. of dil. $\text{ZnSO}_4 + \text{Aq}$ at 20.004° .
Conc. = g. equiv. per l. at 20.004° .
Sp. gr. compared with H_2O at $20.004^\circ = 1$.

Conc.	Sp. gr.
0.0000	1.000,000,0
0.0001	1.000,008,6
0.0002	1.000,017,2
0.0005	1.000,043,2
0.0010	1.000,086,3
0.0020	1.000,172,3
0.0050	1.000,429,1
0.0060	1.000,514,3
1.0100	1.000,853,9

(Lamb and Lee, J. Am. Chem. Soc. 1913, 35. 1690.)

Sulphurous acid, anhydrous, SO₂.

See Sulphur dioxide.

Sulphurous acid, H₂SO₃.

Known only in aqueous solution, from which SO₂ is given off upon heating. Crystallizes in cold, with various amounts of water, forming compounds which approximate H₂SO₃+8H₂O (Pierre, A. 68. 228); H₂SO₃+10H₂O (Döpping, Bull. Ac. St. Pétersb. 7. 100); H₂SO₃+14H₂O (Schönfeld, A. 95. 22); H₂SO₃+6H₂O (Roozeboom, R. t. c. 3. 29, 59, 75, 84; Geuther, A. 224. 218). Crystals are sol. in 2 pts. H₂O at 10°. (Pierre.)

For sp. gr. of solutions, etc., see sulphur dioxide.

Sulphites.

Normal. Only the alkali sulphites are sol. in H₂O, and they are insol. or only sl. sol. in alcohol.

Insol. in liquid NH₃. (Franklin, Am. ch. J. 1898, 20. 824.)

Acid. All the acid sulphites are sol. in H₂O.

In general it is rarely possible to determine whether the compd. described is a pure chemical compound or not. It is probable that many substances described by Svenssen and others are isomorphic mixtures whose composition depends upon the temp. and conc. of the solution in which it was pptd. (Rosenheim, Z. anorg. 1900, 25. 72.)

Aluminum sulphite, basic, Al₂O₃, SO₂+4H₂O.

Insol. in H₂O; sol. in H₂SO₃+Aq. (Fourcroy and Vauquelin.)

6Al(OH)₃, Al₂(SO₃)₃+9H₂O. Ppt. (Seubert, Z. anorg. 1893, 4. 66.)

Ammonium sulphite, basic, (NH₄)₂SO₃, NH₃+¹/₂H₂O.

Sol. in H₂O. Pptd. from aqueous solution by alcohol. (Muspratt.)

Does not exist. (Marignac.)

Ammonium sulphite, (NH₄)₂SO₃.

Very hygroscopic. (Divers, Chem. Soc. 1900, 77. 336.)

Insol. in acetone. (Eidmann, C.C. 1899, II. 1014.)

+H₂O. Slowly sol. in H₂O. (Muspratt, A. 50. 268.)

Sol. in 1 pt. H₂O at 12°. (Fourcroy and Vauquelin, Crell. Ann. 1800. 2. 415.)

More sol. in hot H₂O with evolution of NH₃. Sl. sol. in absolute alcohol. (Muspratt.)

Much more sol. in alcohol than K₂SO₃. (Pierre.)

Loses NH₃ in the air.

Sol. in H₂O. Conc. solution charged with NH₃ will deposit salt on evaporation over KOH. Dil. solution decomp. on evaporation. (Divers, Chem. Soc. 1900, 77. 335.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Ammonium hydrogen sulphite, NH₄HSO₃.

Insol. in acetone. (Eidmann, C.C. 1899, II. 1014; Naumann, B. 1904, 37. 4329.)

Ammonium pyrosulphite, (NH₄)₂S₂O₃.

Deliquescent. Very sol. in H₂O and alcohol. Insol. in ether. (Fock and Klm. B. 23. 3149.)

Very sol. in H₂O; very hygroscopic. Aq. solution is sl. decomp. on evaporation. (Divers, Chem. Soc. 1900, 77. 336.)

Ammonium cadmium sulphite, (NH₄)₂SO₃, CdSO₃.

Nearly insol. in H₂C. Partly sol. in excess of H₂SO₃+Aq, but separates out on boiling. (Schuler, A. 87. 34.)

Ammonium cobaltous sulphite, (NH₄)₂SO₃, CoSO₃+xH₂O.

Decomp. on air. (Berglund, B. 7. 469.)

Ammonium cobaltocobaltic sulphite.

See Cobaltisulphite, ammonium cobalt.

Ammonium cuprous sulphite, (NH₄)₂SO₃, 2Cu₂SO₃+2H₂O.

(Böttlinger, A. 51. 411.)

(NH₄)₂SO₃, Cu₂SO₃. Insol. in cold. decomp. by boiling H₂O. (Rogojski, J. B. 1861. 366.)

Decomp. by warming with H₂O, in which it is insol. Sol. in acids with evolution of SO₂. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 99.)

+2H₂O. (Commaille, J. B. 1867. 300.)
2(NH₄)₂SO₃, Cu₂SO₃+3H₂O. Very sol. in H₂O. Solution decomp. on standing. Decomp. by acids. (Rosenheim and Steinhäuser.)

5(NH₄)₂SO₃, Cu₂SO₃+2H₂O. Decomp. on air. Sol. in H₂O with decomp. (Svensson.)

6(NH₄)₂SO₃, Cu₂SO₃+4H₂O. Easily decomp. (Rosenheim and Steinhäuser.)

7(NH₄)₂SO₃, Cu₂SO₃+4H₂O. Very sol. in H₂O. Solution soon decomp. (Rosenheim and Steinhäuser.)

+10H₂O. Decomp. on air. Sl. sol. in warm, less sol. in cold H₂O. (de Saint-Gilles.)

+14H₂O. Decomp. on air. Sol. in H₂O, but solution decomp.

Very easily sol. in mother liquor. (Svensson, Acta Lund. 1899. 13.)

Ammonium cuprocupric sulphite, (NH₄)₂SO₃, 2Cu₂SO₃, CuSC₃+5H₂O.

Insol. in H₂O and weak acids. Sol. in NH₄OH+Aq. (de Saint-Gilles, A. ch. (3) 42. 31.)

+6¹/₂H₂O. Ppt. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 98.)

Ammonium glucinum sulphite,
 $(\text{NH}_4)_2\text{O}, 2\text{GIO}, 3\text{SO}_2 + 4\text{H}_2\text{O}.$

Ppt. Very unstable in the air. (Rosenheim, Z. anorg. 1897, **15**. 310.)

Ammonium gold (aurous) sulphite,
 $3(\text{NH}_4)_2\text{SO}_3, \text{Au}_2\text{SO}_3.$

Very easily sol. in H_2O . Insol. in alcohol. (Haase, Z. Ch. 1869. 535.)

Ammonium gold (aurous) sulphite ammonia,
 $(\text{NH}_4)_2\text{SO}_3, 3\text{Au}_2\text{SO}_3, 6\text{NH}_3 + \text{H}_2\text{O}.$

Decomp. by H_2O . Sol. in warm $\text{NH}_4\text{OH} + \text{Aq}$, but decomp. by boiling.

$(\text{NH}_4)_2\text{Au}_2(\text{SO}_3)_3, 3\text{NH}_3 + 4\text{H}_2\text{O}.$ Decomp. by H_2O . (Rosenheim, Z. anorg. 1908, **59**. 201.)

Ammonium iridium sulphite.

See Iridosulphite, ammonium.

Ammonium iron (ferrous) sulphite,
 $(\text{NH}_4)_2\text{SO}_3, \text{FeSO}_3 + x\text{H}_2\text{O}.$

(Berglund.)

Ammonium iron (ferric) sulphite sulphate,
 $\text{FeSO}_3\text{SO}_4\text{NH}_4 + \text{H}_2\text{O}.$

Sl. sol. in cold H_2O . Decomp. by cold dil. HCl . (Hofmann, Z. anorg. 1897, **14**. 287.)

Ammonium magnesium sulphite,
 $(\text{NH}_4)_2\text{Mg}_2(\text{SO}_3)_4 + 18\text{H}_2\text{O}.$

Very sl. sol. in H_2O . (Fourcroy and Vauquelin.)

Sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$.

$+ 5\text{H}_2\text{O}$. Much more sol. in H_2O than MgSO_3 . (Rammelsberg.)

Ammonium manganous sulphite, $(\text{NH}_4)_2\text{SO}_3,$
 $\text{MnSO}_3.$

Relatively easily decomp. by H_2O . (Berglund, Bull. Soc. (2) **21**. 213.)

Not easily decomp. (Gorgeu, C. R. **96**. 376.)

Ammonium mercuric sulphite, $(\text{NH}_4)_2\text{SO}_3,$
 $\text{HgSO}_3.$

Very easily sol. in H_2O , but H_2O solution gradually decomp., even in the cold.

Ammonium nickel sulphite, $(\text{NH}_4)_2\text{SO}_3,$
 $3\text{NiSO}_3 + 18\text{H}_2\text{O}.$

Sol. in H_2O . (Berglund, B. **7**. 469.)

Ammonium platinous sulphite.

See Platosulphite, ammonium.

Ammonium potassium sulphite, $10(\text{NH}_4)_2\text{SO}_3,$
 $\text{K}_2\text{SO}_3 + 11\text{H}_2\text{O}.$

Decomp. by H_2O , etc. (Hartog, C. R. **109**. 221.)

Ammonium scandium sulphate,
 $(\text{NH}_4)_2\text{SO}_3, \text{Sc}_2(\text{SO}_3)_3 + 7\text{H}_2\text{O}.$

Insol. in H_2O . Difficulty sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Meyer, Z. anorg. 1914, **86**. 281.)

Ammonium silver sulphite, $(\text{NH}_4)_2\text{SO}_3,$
 $\text{Ag}_2\text{SO}_3.$

Insol. in H_2O , but gradually decomp. thereby. (Svensson, B. **4**. 714.)

$6(\text{NH}_4)_2\text{SO}_3, \text{Ag}_2\text{SO}_3 + 19\text{H}_2\text{O}.$ Sol. in H_2O without decomp. (Svensson.)

$3(\text{NH}_4)_2\text{SO}_3, 4\text{NH}_4\text{HSO}_3, \text{Ag}_2\text{SO}_3 + 18\text{H}_2\text{O}.$ Easily sol. in H_2O , but decomp. by warming.

Ammonium sodium hydrogen sulphite,
 $\text{NH}_4\text{Na}_2\text{H}(\text{SO}_3)_2 + 4\text{H}_2\text{O}.$

Not deliquescent. (Marignac, Ann. Min. (5) **12**. 29.)

100 pts. H_2O dissolve 42.3 pts. salt at 12.4° , and 48.5 pts. at 15° . (Schwicker, B. **22**. 1732.)

$+ 5\text{H}_2\text{O} = 2\text{Na}_2\text{SO}_3, (\text{NH}_4)_2\text{S}_2\text{O}_3 + \text{H}_2\text{O}.$ (Tauber, Techn. J. B. 1888. 444.)

Ammonium tellurium sulphite, $(\text{NH}_4)_2\text{SO}_3,$
 $\text{TeSO}_3 + x\text{H}_2\text{O}.$

Sol. in H_2O . (Berglund, B. **7**. 469.)

Ammonium uranyl sulphite,
 $\text{NH}_4(\text{UO}_2)(\text{OH})\text{SO}_3.$

Insol. in pure H_2O . More sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$ than the K salt, and less than the Na salt. (Scheller, A. **144**. 240.)

$(\text{NH}_4)_2\text{O}, 2\text{UO}_3, 3\text{SO}_3.$

$(\text{NH}_4)_2\text{O}, 4\text{UO}_3, 5\text{SO}_3.$

$(\text{NH}_4)_2\text{O}, 3\text{UO}_3, 2\text{SO}_3.$

$(\text{NH}_4)_2\text{O}, \text{UO}_3, 2\text{SO}_3.$ (Kohlschütter, A. 1900, **311**. 10.)

Ammonium vanadium sulphite.

See Vanadiousulphite, ammonium.

Ammonium vanadyl sulphite,
 $(\text{NH}_4)_2\text{SO}_3, \text{VOSO}_3 + 2\text{H}_2\text{O}.$

Sol. in H_2O with decomp. (Koppel, Z. anorg. 1903, **35**. 184.)

$(\text{NH}_4)_2\text{O}, 3\text{VO}_3, 2\text{SO}_2 + \text{H}_2\text{O}.$ Sol. in cold H_2O without decomp.

Easily sol. in mineral acids and alkalis.

Sl. sol. in alcohol and ether. (Koppel Z. anorg. 1903, **35**. 182.)

Ammonium zinc sulphite, $(\text{NH}_4)_2\text{SO}_3, \text{ZnSO}_3.$
 Sol. in H_2O . (Berglund, B. **7**. 469.)

Ammonium sulphite mercuric chloride,
 $2(\text{NH}_4)_2\text{SO}_3, \text{HgCl}_2.$

Sl. sol. in cold, decomp. by boiling H_2O . (de St-Giles, A. ch. (3) **36**. 95.)

Antimony sulphite, $\text{Sb}_2\text{O}_3, 3\text{SO}_3(?)$.

Insol. in H_2O . (Berzelius.)

Could not be obtained. (Röhrig, J. pr. (2) **37**. 241.)

(Rosenheim and Steinhäuser, Z. anorg. 1890, 25. 96.)

Cu_2SO_3 , $2\text{K}_2\text{SO}_3$. (Chevreul, Graham, etc.)

Does not exist. (Svensson.)

Cu_2O , $3\text{K}_2\text{O}$, $6\text{SO}_2 + 7\text{H}_2\text{O} = 4\text{KHSO}_3$, K_2SO_3 , $\text{Cu}_2\text{SO}_3 + 5\text{H}_2\text{O}$. Decomp. by H_2O . (Svensson, B. 4. 713.)

Could not be obtained. (Rosenheim and Steinhäuser.)

Cu_2O , $4\text{K}_2\text{O}$, $8\text{SO}_2 + 3\text{H}_2\text{O} = 6\text{KHSO}_3$, K_2SO_3 , Cu_2SO_3 . Decomp. by H_2O . (Svensson.)

Could not be obtained. (Rosenheim and Steinhäuser.)

Cu_2SO_3 , $8\text{K}_2\text{SO}_3 + 16\text{H}_2\text{O}$. Sol. in H_2O with decomp. (Rammelsberg, Pogg. 57. 391.)

Does not exist, according to Svensson.

Cuprocupric potassium sulphite, $3\text{Cu}_2\text{SO}_3$, 3CuSO_3 , K_2SO_3 .

Properties as cuprous potassium sulphite. (Rogojski, J. B. 1851. 367.)

$2\text{Cu}_2\text{SO}_3$, CuSO_3 , $\text{K}_2\text{SO}_3 + 5\text{H}_2\text{O}$. Insol. in H_2O and weak acids. (de St-Gilles.)

Cu_2SO_3 , 4CuSO_3 , $\text{K}_2\text{SO}_3 + 16\text{H}_2\text{O}$. Decomp. by H_2O . (Rosenheim and Steinhäuser.)

Cuprous sodium sulphite, Cu_2SO_3 , $\text{Na}_2\text{SO}_3 + 2\text{H}_2\text{O}$. Decomp. by H_2O . (Svensson, 1870.)

$+11\text{H}_2\text{O}$. Insol. in cold H_2O , but decomp. by excess. (Étard, C. R. 95. 138.)

$2\text{Cu}_2\text{SO}_3$, $3\text{Na}_2\text{SO}_3 + 29\text{H}_2\text{O}$. Insol. in H_2O . (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 94.)

Cu_2SO_3 , $5\text{Na}_2\text{SO}_3 + 38\text{H}_2\text{O}$. Decomp. by H_2O . (Svensson.)

Cu_2SO_3 , $7\text{Na}_2\text{SO}_3 + 19\text{H}_2\text{O}$. Completely sol. in H_2O , but solutions decomp. on standing. (Svensson.)

"Cuprous sodium octosulphite,"

$(\text{Cu}_2)_2\text{H}_{10}\text{Na}_{16}\text{S}_8\text{O}_{32} + 43\text{H}_2\text{O}$. (Étard.)

$5\text{Cu}_2\text{SO}_3$, $2\text{Na}_2\text{SO}_3 + 30\text{H}_2\text{O}$. Easily decomp. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 94.)

Cuprocupric sodium sulphite,

Cu_2SO_3 , 2CuSO_3 , $2\text{Na}_2\text{SO}_3 + 6\text{H}_2\text{O}$.

Nearly insol. in cold, decomp. by hot H_2O . (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 95.)

$+8\text{H}_2\text{O}$. Decomp. by H_2O . (Rosenheim and Steinhäuser.)

Cuprocupric sodium hydrogen sulphite,

$\text{Na}_2\text{Cu}^{\text{II}}(\text{Cu}^{\text{I}})_2(\text{SO}_3)_2 \cdot 6\text{H}_2(\text{SO}_3) + 5\text{H}_2\text{O}$.

Insol. in H_2O . (Étard, C. R. 94. 1422.)

$(\text{Cu}^{\text{I}})_2\text{Cu}^{\text{II}}\text{Na}_2\text{H}_{12}(\text{SO}_3)_2$. (Étard.)

Copper sodium sulphites.

Doubtless many of the compds. described in this class are in reality isomorphous mixtures whose composition depends upon the temp. and conc. of the solution in which pptd.

(Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 92-95.)

Didymium sulphite, $\text{Di}_2(\text{SO}_3)_3 + 3\text{H}_2\text{O}$, or $6\text{H}_2\text{O}$.

Precipitate. Insol. in H_2O . Sol. in $\text{H}_2\text{SO}_4 + \text{Aq}$, from which it is reprecipitated by heating, redissolving on cooling. (Marignac, A. ch. (3) 38. 167.)

Erbium sulphite, $\text{Er}_2(\text{SO}_3)_3 + 3\text{H}_2\text{O}$.

Precipitate.

Glucinum sulphite, basic, 2GlSO_3 , $9\text{Gl}(\text{OH})_2 + 6\text{H}_2\text{O}$.

Ppt. (Seubert, Z. anorg. 1893, 4. 52.)

GlSO_3 , GlO . Decomp. by H_2O or alcohol (K. and M.)

3GlSO_3 , GlC . Sol. in alcohol. (K. and M.)

Glucinum sulphite, GlSO_3 .

Decomp. by H_2O or alcohol. (Krüger and Moraht, B. 23. 734.)

Glucinum potassium sulphite, 2GlSO_3 , $\text{K}_2\text{SO}_3 + 9\text{H}_2\text{O}$.

Unstable in the air. (Rosenheim, Z. anorg. 1897, 15. 310.)

Gold (aurous) potassium sulphite, Au_2SO_3 , $3\text{K}_2\text{SO}_3$.

Very sol. in H_2O ; insol. in alcohol. (Haas.)

Gold (auric) potassium sulphite, Au_2O_3 , $5\text{K}_2\text{O}$, $8\text{SO}_2 + 5\text{H}_2\text{O} = 5\text{K}_2\text{SO}_3$, $\text{Au}_2(\text{SO}_3)_3 + 5\text{H}_2\text{O}$.

Sol. in H_2O with decomp.

Decomp. by acids; insol. in alkalis. (Freymy, A. 79. 46.)

Gold (auric) potassium sulphite, $\text{Au}_2(\text{SO}_3)_3$, $5\text{K}_2\text{SO}_3 + 10\text{H}_2\text{O}$.

(Rosenheim and Hertzmann, Z. anorg. 1906, 59. 199.)

Gold (auric) potassium sulphite ammonia, $\text{Au}_2(\text{SO}_3)_3$, $3\text{K}_2\text{SO}_3$, $4\text{NH}_3 + 4\text{H}_2\text{O}$.

As the corresponding NH_4 salt. (Rosenheim and Hertzmann, Z. anorg. 1906, 59. 202.)

Gold (aurous) sodium sulphite, Au_2SO_3 , $3\text{Na}_2\text{SO}_3 + 3\text{H}_2\text{O}$.

Sol. in less than 1 pt. H_2O . Insol. in alcohol (Hasse.)

$+5\text{H}_2\text{O}$. (Himly.)

Gold (auric) sodium sulphite,

$\text{Au}_2(\text{SO}_3)_3$, $5\text{Na}_2\text{SO}_3 + 28\text{H}_2\text{O}$.

As K salt. (Rosenheim and Hertzmann, Z. anorg. 1906, 59. 199.)

Gold (aurous) sulphite ammonia, $3\text{Au}_2\text{O}$, 4SO_2 , $8\text{NH}_3 + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O with decomp. Decomp. by acids.

Sl. sol. in cold, more easily in hot $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by boiling. (Hasse, Zeit. Ch. 1899. 535.)

Gold (auric) sulphite ammonia, $\text{Au}_2(\text{SO}_3)_3$, $4\text{NH}_3 + 4\text{H}_2\text{O}$.

Ppt. Decomp. in moist air and in neutral solution (Herzmann, Z. anorg. 1908, 59. 198.)

Indium sulphite, $2\text{In}_2\text{O}_3$, $3\text{SO}_2 + 8\text{H}_2\text{O}$.

Insol. in H_2O . (Bayer, A. 158. 372.)

Iridium sulphite, $\text{Ir}_2(\text{SO}_3)_3 + 6\text{H}_2\text{O}$.

Scarcely sol. in H_2O ; easily sol. $\text{HCl} + \text{Aq}$. (Birnbaum, A. 136. 179.)

Iridyl sulphite, $(\text{IrO})\text{SO}_3 + 4\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$. (Birnbaum.)

Iridous potassium sulphite, IrO , $3\text{K}_2\text{O}$, $5\text{SO}_2(?)$.

Sl. sol. in H_2O , more sol. in $\text{KOH} + \text{Aq}$. Easily sol. in $\text{HCl} + \text{Aq}$. (Claus, J. pr. 42. 359.)

Iridous sulphite potassium chloride.

See Iridosulphite, potassium.

Iridium sulphite with M_2SO_3 .

See Iridosulphite, M.

Iron (ferrous) sulphite, $\text{FeSO}_3 + 2\frac{1}{2}\text{H}_2\text{O}$.

Very sl. sol. in H_2O . Easily sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. Insol. in alcohol, but sol. therein in presence of SO_2 . (Muspratt.)

Iron (ferric) sulphite, Fe_2O_3 , $\text{SO}_3 + 6\text{H}_2\text{O}$.

Very sl. sol. in H_2O . Sol. in acids. (Koene.) $2\text{Fe}_2\text{O}_3$, 3SO_2 . Deliquescent; decomp. by H_2O into SO_2 and above comp. $3\text{Fe}_2\text{O}_3$, $\text{SO}_2 + 7\text{H}_2\text{O}$. Ppt.

Iron (ferroferric) potassium sulphite, FeSO_3 , $(\text{FeO})_2\text{SO}_3$, $2\text{K}_2\text{SO}_3$.

Ppt. (Berglund.)

Iron (ferric) potassium sulphite, K_2O , Fe_2O_3 , $3\text{SO}_2 + 2\text{H}_2\text{O}$.

Sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Koene, Pogg. 63. 453.)

Fe_2O_3 , $2\text{K}_2\text{O}$, $3\text{SO}_2 + 5\text{H}_2\text{O}$. Ppt. (Muspratt, Phil. Mag. (3) 30. 414.)

Iron (ferric) potassium sulphite sulphate, $\text{FeSO}_3\text{SO}_4\text{K}$.

Sl. sol. in cold H_2O .

Sol. in 20% HCl ; decomp. on boiling. (Hofmann, Z. anorg. 1897, 14. 286.)

$\text{Fe}(\text{SO}_3)_2\text{SO}_4\text{K}$. Almost insol. in cold H_2O . Decomp. by boiling with dil. acids. (Hofmann.)

$\text{Fe}_2(\text{SO}_3)_3\text{SO}_4\text{K}_4 + 5\text{H}_2\text{O}$. Insol. in cold H_2O ; sol. in cold 20% $\text{HCl} + \text{Aq}$; decomp. on boiling with H_2O . (Hofmann.)

Iron (ferric) sodium sulphite sulphate, $\text{Fe}(\text{SO}_3)_2\text{SO}_4\text{Na}_3 + 6\text{H}_2\text{O}$.

Almost insol. in H_2O .

Decomp. by boiling with dil. acids. (Hofmann, Z. anorg. 1897, 14. 289.)

Iron (ferric) sodium hydrogen sulphite sulphate, $\text{FeSO}_3(\text{SO}_3)_2\text{H}_2\text{Na}_2 + 2\text{H}_2\text{O}$.

Only very sl. sol. in H_2O . (Hofmann.)

Lanthanum sulphite, $\text{La}_2(\text{SO}_3)_3 + 4\text{H}_2\text{O}$.

Precipitate. (Cleve.)

Lead sulphite, PbSO_3 .

Insol. in H_2O . Decomp. by acids. Sl. sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Röhrig, J. pr. (2) 37. 233.)

Lithium sulphite, $\text{Li}_2\text{SO}_3 + 6\text{H}_2\text{O}$.

Sol. in H_2O ; precipitated from aqueous solution by abs. alcohol. (Danson, Chem. Soc. 2. 205.) Sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$.

+ H_2O . Sl. sol. in alcohol, and still less sol. in ether. (Röhrig, J. pr. (2) 37. 225.) + $2\text{H}_2\text{O}$. (Röhrig.)

Lithium potassium sulphite, $\text{LiKSO}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Easily sol. in H_2O . (Röhrig, J. pr. (2) 37. 251.)

Lithium sodium sulphite, $6\text{Li}_2\text{SO}_3$, $\text{Na}_2\text{SO}_3 + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Röhrig.)

Magnesium sulphite, $\text{MgSO}_3 + 6\text{H}_2\text{O}$.

Sol. in 20 pts. cold, and in less hot H_2O . (Fourcroy and Vauquelin.)

Sol. in 80 pts. cold, and in 120 pts. boiling H_2O . (Hager, C. C. 1875. 135.)

More easily sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 828.)

Precipitated from aqueous solution by alcohol.

+ $3\text{H}_2\text{O}$. (Röhrig, J. pr. (2) 37. 234.)

Manganous sulphite, $\text{MnSO}_3 + 2\text{H}_2\text{O}$.

Insol. in H_2O , alcohol, or ether. Easily sol. in acids, also in $\text{H}_2\text{SO}_3 + \text{Aq}$.

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

+ $2\frac{1}{2}\text{H}_2\text{O}$. (Rammelsberg.)

+ $3\text{H}_2\text{O}$. Sol. in 10,000 pts. cold, and 5000 pts. hot H_2O ; more sol. in conc. Mn salts + Aq ; sol. in 1000 pts. $\text{H}_2\text{CO}_3 + \text{Aq}$. 100 pts. $\text{H}_2\text{SO}_3 + \text{Aq}$ dissolve 15–17 pts. (Gorgeu, C. R. 96. 341.)

Salt with $2\frac{1}{2}\text{H}_2\text{O}$ is the only one which exists. (Röhrig, J. pr. (2) 37. 2.)

Manganous potassium sulphite, $2\text{MnSO}_3, \text{K}_2\text{SO}_3$.

Insol. in H_2O , even when boiling. (Gorgeu, C. R. 96. 376.)

$\text{MnSO}_3, \text{K}_2\text{SO}_3$. Sol. in H_2O . (Gorgeu.)

Manganous sodium sulphite, $\text{MnSO}_3, \text{Na}_2\text{SO}_3 + \text{H}_2\text{O}$.

Insol. in hot H_2O , but decomp. by cold H_2O . (Gorgeu.)

$4\text{MnSO}_3, \text{Na}_2\text{SO}_3$. Insol. in H_2O . (Gorgeu.)

Mercuric sulphite, $2\text{HgO}, \text{SO}_3$.

Insol. in H_2O . Sol. in HCl , alkali sulphites with subsequent decomp., and in $\text{KCN} + \text{Aq}$. (de St-Gilles, A. ch. (3) 36. 80.)

HgSO_3 . Decomp. by cold H_2O . (de St-Gilles.)

Does not exist. (Divers and Shimidzu, Chem. Soc. 49. 553.)

$\text{HgO}, 2\text{SO}_3 + \text{H}_2\text{O}$. Sol. in H_2O , but decomp. by boiling. (de St-Gilles.) Exists only in aqueous solution. (Divers and Shimidzu.)

Mercuromercuric sulphite, $\text{Hg}_2(\text{SO}_3)_2 + 2\text{H}_2\text{O} = \text{Hg}_2\text{SO}_3, \text{HgSO}_3$.

Very efflorescent. Insol. in H_2O . Decomp. by hot H_2O . Insol. in dil. HNO_3 or $\text{H}_2\text{SO}_4 + \text{Aq}$.

$+4\text{H}_2\text{O}$. Very efflorescent.

Hypomercurosic sulphite, $\text{Hg}_4(\text{SO}_3)_2 + \text{H}_2\text{O}$.

Insol. in H_2O , but easily decomp. on standing therewith. Almost absolutely insol. in dil. HNO_3 or $\text{H}_2\text{SO}_4 + \text{Aq}$. (Divers and Shimidzu.)

Mercuric oxysulphite, $\text{Hg}(\text{SO}_3\text{OHgO}), \text{Hg} + \text{H}_2\text{O}$.

Insol. in H_2O . Decomp. by hot H_2O . Insol. in dil. HNO_3 or $\text{H}_2\text{SO}_4 + \text{Aq}$. Sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Divers and Shimidzu.)

Mercuric potassium sulphite, basic, $\text{K}_2\text{O}, 2\text{HgO}, 2\text{SO}_3$.

(Barth, Z. phys. Ch. 1892, 9. 210.)

$\text{K}_2\text{O}, 3\text{HgO}, 3\text{SO}_3$. Insol. in H_2O . Partly sol. in $\text{KOH} + \text{Aq}$. (Barth.)

Mercuric potassium sulphite, $\text{HgSO}_3, \text{K}_2\text{SO}_3 + \text{H}_2\text{O}$.

Sl. sol. in cold H_2O . Decomp. on boiling. (de St-Gilles, A. ch. (3) 36. 90.)

Mercuric potassium sulphite mercuric chloride, $\text{K}_2\text{Hg}(\text{SO}_3)_2, \text{HgCl}_2$.

Decomp. by H_2O . (Barth, Z. phys. Ch. 1892, 9. 206.)

Mercuric silver sulphite, $\text{HgSO}_3, \text{Ag}_2\text{SO}_3 + 2\text{H}_2\text{O}$.

Decomp. rapidly; insol. in H_2O . (Barth, Z. phys. Ch. 9. 195.)

Mercuric sodium sulphite, $\text{HgSO}_3, \text{Na}_2\text{SO}_3 + \text{H}_2\text{O}$.

Sol. in H_2O . (de St-Gilles.)

Sol. in 25 pts. cold H_2O , and decomp. on heating. (Divers and Shimidzu.)

$+2\text{H}_2\text{O} = \text{Na}_2(\text{SO}_3)_2\text{Hg} + 2\text{H}_2\text{O}$. (Barth, Z. phys. Ch. 9. 193.)

$2\text{HgSO}_3, \text{Na}_2\text{SO}_3 + \text{H}_2\text{O}$. Much more sol. in H_2O than the above comp. especially on heating. (de St-Gilles.)

Does not exist. (Divers and Shimidzu.)

Mercuric strontium sulphite, $\text{HgSO}_3, \text{SrSO}_3 + 2\text{H}_2\text{O}$.

Ppt. (Barth.)

Mercuric sulphite ammonium bromide, $\text{HgSO}_3, \text{NH}_4\text{Br}$.

As NH_4Cl comp. (Barth, Z. phys. Ch. 1892, 9. 215.)

Mercuric sulphite ammonium chloride, $\text{HgSO}_3, \text{NH}_4\text{Cl}$.

As K salt. (Barth.)

Mercuric sulphite potassium chloride, $\text{HgSO}_3, \text{KCl}$.

Sol. in H_2O . (Barth.)

Mercuric sulphite sodium chloride, $\text{HgSO}_3, \text{NaCl} + \text{H}_2\text{O}$.

Sol. in H_2O . (Barth.)

Nickel sulphite, basic, $2\text{NiSO}_3, \text{Ni}(\text{OH})_2 + 6\text{H}_2\text{O}$.

Ppt. (Seubert and Elten, Z. anorg. 1893. 4. 91.)

Nickel sulphite, $\text{NiSO}_3 + 4\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$. with evolution of SO_3 . (Muspratt, A. 60. 259.)

$+6\text{H}_2\text{O}$. Insol. in H_2O . Sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Rammelsberg, Pogg. 67. 391.)

Nickel sulphite ammonia, $\text{NiSO}_3, 3\text{NH}_3 + 3\text{H}_2\text{O}$.

Sol. in little H_2O . Decomp. by much H_2O or heat. (Rammelsberg, Pogg. 67. 245.)

Osmious sulphite, OsSO_3 .

Insol. in H_2O . Easily sol. in $\text{HCl} + \text{Aq}$ without evolution of SO_3 . Very slowly decomp. by $\text{KOH} + \text{Aq}$. (Claus.)

Osmious potassium sulphite, $\text{OsSO}_3, 2\text{K}_2\text{SO}_3, 2\text{KHSO}_3 + 4\text{H}_2\text{O}$.

Nearly insol. in H_2O .

Osmious potassium sulphite chloride, $\text{OsO}_2 \cdot 2\text{SO}_2 \cdot 6\text{KCl}$.

Easily sol. in H_2O .

Palladous sodium sulphite, $\text{PdSO}_3 \cdot 3\text{Na}_2\text{SO}_3 + 2\text{H}_2\text{O} = \text{Na}_4\text{Pd}(\text{SO}_3)_4 + 2\text{H}_2\text{O}$.

Sol. in hot H_2O . Sol. in $\text{NaOH} + \text{Aq}$ or $\text{H}_2\text{SO}_3 + \text{Aq}$. (Wöhler and Frerichs, A. 174. 199.)

Platinous sulphite, $\text{PtO}_2 \cdot 2\text{SO}_2$.

Easily sol. in H_2O or alcohol. (Döbereiner, J. pr. 15. 315.)

Formula is PtSO_3 . (Gmelin.)

PtSO_3 , H_2SO_3 . (Birnbaum, A. 139. 172.)

Platinic potassium sulphite, $\text{PtO}_2 \cdot \text{SO}_2 \cdot \text{K}_2\text{SO}_3 + \text{H}_2\text{O}$.

Sol. in $\text{KOH} + \text{Aq}$. (Birnbaum, A. 139. 173.)

Platinic sodium sulphite, $\text{PtO}_2 \cdot \text{SO}_2 \cdot 2\text{Na}_2\text{SO}_3 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Birnbaum.)

Platinous sulphite with M_2SO_3 .

See **Platosulphite, M**.

Platinum sulphite ammonium chloride.

See **Chloroplatosulphite, ammonium**.

Potassium sulphite, $\text{K}_2\text{SO}_3 + 2\text{H}_2\text{O}$.

Somewhat deliquescent. Sol. in 1 pt. cold, and still less hot H_2O . (Fourcroy and Vauquelin, A. ch. 24. 254.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Very slightly soluble in alcohol. Insol. in ethyl acetate. (Casaseca, C. R. 30. 821.)

Potassium hydrogen sulphite, KHSO_3 .

Sol. in H_2O . Insol. in absolute alcohol.

Potassium pyrosulphite, $\text{K}_2\text{S}_2\text{O}_5$.

Slowly sol. in H_2O . Very sl. sol. in alcohol; insol. in ether. (Muspratt, A. 50. 259.)

Potassium rhodium sulphite, $3\text{K}_2\text{SO}_3 \cdot \text{Rh}_2(\text{SO}_3)_3 + 6\text{H}_2\text{O}$.

See **Rhodosulphite, potassium**.

Potassium ruthenium sulphite,

$\text{O}[\text{Ru}(\text{SO}_3)_4\text{K}_3]_2 + 2\text{H}_2\text{O}$.

Ppt. (Miolati, C. C. 1901, I. 501.)

Potassium sodium sulphite, KNaSO_3 .

Sol. in H_2O . (Spring, B. 7. 1161.)

+1, and $2\text{H}_2\text{O}$. (Schwicker, B. 22. 1731.)

Isomeric salts, KSO_3Na and NaSO_3K . (Barth, Z. phys. Ch. 9. 176.)

Potassium sodium hydrogen sulphite, $\text{KNa}_2\text{H}(\text{SO}_3)_2 + 4\text{H}_2\text{O}$.

Easily sol. in H_2O ; 100 pts. H_2O dissolve 69 pts. salt at 15° . (Schwicker, B. 22. 1731.)

$\text{K}_2\text{NaH}(\text{SO}_3)_2 + 3\text{H}_2\text{O}$. (Schwicker.)

Potassium uranyl sulphite, $\text{K}(\text{UO}_2)(\text{OH})\text{SO}_3$.

Insol. in H_2O , but sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$. (Scheller.)

K_2O , 2UO_2 , 3SO_2 . (Kohlschütter, A. 1900, 311. 10 *et seq.*)

K_2O , 4UO_2 , 5SO_2 . (K.)

K_2O , 3UO_2 , 2SO_2 . (K.)

K_2O , UO_2 , 2SO_2 . (K.)

Potassium vanadium sulphite.

See **Vanadidosulphite, potassium**.

Potassium vanadyl sulphite, $\text{K}_2\text{SO}_3 \cdot \text{VO}\text{SO}_3 + 5\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O without decomp. and can be recryst. therefrom. (Koppel and Behrendt, B. 1901, 34. 3932.)

K_2O , 3VO_2 , 2SO_2 . Sol. without decomp. in cold and hot H_2O .

Insol. in alcohol and ether. (Koppel, Z. anorg. 1903, 35. 182.)

Potassium zinc sulphite, $\text{K}_2\text{SO}_3 \cdot 3\text{ZnSO}_3 + 7\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Berglund, Acta Lund. 1872.)

Rhodium sulphite, $\text{Rh}_2(\text{SO}_3)_3 + 6\text{H}_2\text{O}$.

Sol. in H_2O . Insol. in alcohol. (Claus.)

Rhodium sodium sulphite.

See **Rhodosulphite, sodium**.

Ruthenium sulphite, $\text{Ru}_2(\text{SO}_3)_3$.

Colloidal substance, sol. in a large quantity of H_2O . (Lucchesi, Gazz. ch. it. 1900, 30. (2) 71.)

Ruthenium sodium sulphite, $\text{Na}_7\text{Ru}(\text{SO}_3)_6 + 2\text{H}_2\text{O}$.

Ppt. (Miolati, C. C. 1901, I. 501.)

Samarium sulphite, $\text{Sm}_2(\text{SO}_3)_3$.

Amorphous precipitate. (Cleve.)

Scandium sulphite, $\text{Sc}_2(\text{SO}_3)_3$.

Insol. in cold H_2O . Sl. sol. in hot H_2O .

Sol. in excess of sodium sulphite when heated. (Crookes, Phil. Trans. 1910, 210. A. 363.)

+ $6\text{H}_2\text{O}$. Very sl. sol. in H_2O .

Decomp. by boiling with H_2O with separation of H_2SO_3 . (R. J. Meyer, Z. anorg. 1914, 36. 281.)

Selenium sulphite, SeSO_3 .

Correct composition for "selenium sulphoxide." (Divers, Chem. Soc. 49. 583.)

Silver sulphite, Ag_2SO_3 .

Very sl. sol. in cold H_2O . Decomp. on heating.

Solubility in H_2O is $<1:20,000$. (Baubigny, C. R. 1909, 149. 858.)

Easily sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ and alkali sulphites + Aq. Insol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ Decomp. by strong acids, but not by acetic acid. (Berthier, A. ch. (3) 7. 82.)

Easily sol. in alkali thiosulphates + Aq. (Herschel.)

Cold $\text{NaHSO}_3 + \text{Aq}$ dissolves a considerable amount of Ag_2SO_3 . (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 78.)

Practically insol. in $\text{HNO}_3 + \text{Aq}$ or dil. $\text{AgNO}_3 + \text{Aq}$, also in $\text{H}_2\text{SO}_3 + \text{Aq.}$ (Divers, Chem. Soc. 49. 579.)

Silver sodium sulphite, $\text{Ag}_2\text{SO}_3, \text{Na}_2\text{SO}_3 + \text{H}_2\text{O}$.

Decomp. by H_2O . (Svensson, B. 4. 714.)

Sodium sulphite, Na_2SO_3 .

100 pts. dissolve at 0° , 14.1 pts.; at 20° , 25.8 pts.; at 40° , 49.5 pts. Na_2SO_3 . (Kremers, Pogg. 99. 50.) Maximum solubility is at 33° (Mitscherlich.)

Solubility in 100 pts. H_2O at t° .

t°	Pts. Na_2SO_3
60.4	28.29
59.8	28.29
59.8	28.65
59.8	28.75
37.0	28.01
37.0	28.07
47.0	28.19
47.0	28.07
55.6	28.21
84.0	28.26

The temp. at which $\text{Na}_2\text{SO}_3 + 7\text{H}_2\text{O}$ changes into Na_2SO_3 is about 21.6° .

(Hartley and Barrett, Chem. Soc. 1909, 95. 1183.)

See also $+7\text{H}_2\text{O}$.

Sp. gr. of sat. solution at $15^\circ = 1.21$. (Greenish and Smith, Pharm. J. 1901, 66. 774.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 829.)

Insol. in alcohol.

Insol. in ethyl acetate. (Casaseca, C. R. 30. 821.); methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in benzonitrile. (Naumann, B. 1914, 47. 1370.)

$+7\text{H}_2\text{O}$. Decomp. slowly on air.

Sol. in 4 pts. H_2O at 15° with absorption of heat (Dumas), and in 1 pt. boiling H_2O (Fourcroy).

Solubility in 100 pts. H_2O at t° .

t°	Pts. Na_2SO_3
37.2	44.08
33.5	39.64
29.0	34.99
23.5	29.92
18.2	25.31
10.6	20.01
5.9	17.61
2.0	14.82
-1.9	13.09

Supersolubility curves have also been plotted for ice and $\text{Na}_2\text{SO}_3 + 7\text{H}_2\text{O}$.

(Hartley and Barrett, Chem. Soc. 1909, 95. 1181.)

$+10\text{H}_2\text{O}$. Efflorescent. Somewhat less sol. than above salt. (Muspratt.)

Sodium hydrogen sulphite, NaHSO_3 .

More difficulty sol. in H_2O than NaHCO_3 , and is precipitated by alcohol from aqueous solution. (Muspratt.)

Insol. in acetone. (Eidmann, C. C. 1899, II. 1014; Naumann, B. 1904, 37. 4329; methyl acetate. (Naumann, B. 1909, 42. 3790.)

$+4\text{H}_2\text{O}$. (Clark.)

Sodium pyrosulphite, $\text{Na}_2\text{S}_2\text{O}_3$.

Decomp. gradually on the air.

Sodium uranyl sulphite, $\text{Na}(\text{UO}_2)(\text{OH})\text{SO}_3$.

Sl. sol. in H_2O . More sol. in $\text{H}_2\text{SC}_2 + \text{Aq}$ than the K salt. (Scheller.)

$\text{Na}_2\text{O}, 2\text{UO}_3, 3\text{SO}_3$.

$\text{Na}_2\text{O}, 3\text{UO}_3, 2\text{SO}_3$. (Kohlschütter, A. 1900, 311. 10 et seq.)

Sodium vanadyl sulphite, $\text{Na}_2\text{O}, 2\text{VO}_3, \text{VO}_3 + 5\text{H}_2\text{O}$.

Sol. in H_2O with decomp.

$\text{Na}_2\text{O}, 2\text{SO}_3, 3\text{VO}_3 + 4\text{H}_2\text{O}$. Sol. in cold H_2O ; decomp. on heating. (Koppel, B. 1901, 34. 3933.)

Sodium zinc sulphite, $\text{Na}_2\text{SO}_3, 3\text{ZnSO}_3 + 7\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O with decomp. (Berglund, Acta Lund, 1872.)

Sodium sulphite silver chloride, $3\text{Na}_2\text{SO}_3, \text{AgCl} + 21\text{H}_2\text{O}$.

Sol. in H_2O . (Svensson.)

Strontium sulphite, SrSO_3 .

Precipitate. Almost insol. in H_2O . Sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ (Muspratt.)

Sol. in about 30,000 pts. H_2O at $16-18^\circ$. (Autenrieth, Z. anal. 1898, **37**. 293.)

Abundantly sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ (Röhrig.)

Tellurium sulphite, TeSO_3 .

Correct composition of "tellurium sulphoxide." (Divers, Chem. Soc. **49**. 583.)

Thallous sulphite, Tl_2SO_3 .

Sl. sol. in cold, easily in hot $\text{H}_2\text{SO}_3 + \text{Aq.}$ (Röhrig, J. pr. (2) **37**. 229.)

100 pts. H_2O dissolve 3.34 pts. at 15.5° . Easily sol. in hot H_2O ; insol. in alcohol. (Seubert and Elten, Z. anorg. **2**. 434.)

Thallous vanadyl sulphite, $2\text{Tl}_2\text{SO}_3 \cdot \text{V}_2\text{O}_5 \cdot \text{SO}_3 + 4\text{H}_2\text{O}$.

(Gain, A. ch. 1908, (8) **14**. 278.)

$\text{Tl}_2\text{SO}_3 \cdot 3\text{V}_2\text{O}_5 \cdot \text{SO}_3 + 8\text{H}_2\text{O}$. (Gain.)

Thorium sulphite, $\text{Th}(\text{SO}_3)_2 + \text{H}_2\text{O}$.

Precipitate. (Cleve.)

Tin (stannous) sulphite, $5\text{SnO} \cdot 2\text{SO}_3 + x\text{H}_2\text{O}$.

Ppt. Partly sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ (Röhrig, J. pr. (2) **37**. 249.)

$+20\text{H}_2\text{O}$. (Röhrig.)

$8\text{SnO} \cdot 2\text{SO}_3 + 20\text{H}_2\text{O}$.

$11\text{SnO} \cdot 2\text{SO}_3 + 20\text{H}_2\text{O}$. (Röhrig.)

Uranous sulphite, basic, $\text{U}(\text{OH})_2\text{SO}_3 + \text{H}_2\text{O}$.

Insol. in H_2O . Easily sol. in acids. Sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ but is soon decomp. (Rammelsberg.)

Uranyl sulphite, basic, $3\text{UO}_2(\text{OH})_2 \cdot 5(\text{UO}_2)_2\text{SO}_3 + 10\text{H}_2\text{O}$.

(Seubert and Elten, Z. anorg. 1893, **4**. 80.)

Uranyl sulphite, $(\text{UO}_2)_2\text{SO}_3 + 4\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ or alcoholic solution of SO_2 . (Röhrig, J. pr. (2) **37**. 240.)

Vanadyl sulphite, $3\text{VO}_2 \cdot 2\text{SO}_3 + 4\frac{1}{2}\text{H}_2\text{O}$.

Decomp. slowly on standing.

Sol. in H_2O without apparent decomp. (Koppel, Z. anorg. 1903, **35**. 186.)

$2\text{V}_2\text{O}_4 \cdot 3\text{SO}_3 + 10\text{H}_2\text{O}$. Sol. in H_2O ; aq. sol. decomp. on boiling giving off SO_2 and forming $\text{V}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$. (Gain, C. R. 1906, **143**. 824.)

Vanadyl zinc sulphite, $\text{ZnO} \cdot 3\text{VO}_2 \cdot 2\text{SO}_3$.

Decomp. slowly in the air.

Sol. in H_2O without decomp. (Koppel, Z. anorg. 1903, **35**. 183.)

Ytterbium sulphite, $\text{Yb}_2(\text{SO}_3)_3 + 9\text{H}_2\text{O}$.

Insol. in H_2O . (Cleve, Z. anorg. 1902, **32**. 143.)

Yttrium sulphite, $\text{Y}_2(\text{SO}_3)_3 + 3\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Cleve.)

Zinc sulphite, basic, $2\text{ZnSO}_3 \cdot 3\text{Zn}(\text{OH})_2$.

(Seubert, Arch. Pharm. **229**. 321.)

$\text{ZnSO}_3 \cdot \text{Zn}(\text{OH})_2 + \text{H}_2\text{O}$. (Seubert.)

Zinc sulphite, $\text{ZnSO}_3 + 2$, and $2\frac{1}{2}\text{H}_2\text{O}$.

Very sl. sol. in H_2O . 100 pts. H_2O dissolve 0.16 pt. $\text{ZnSO}_3 + 2\text{H}_2\text{O}$. (Henston and Tichborne, Brit. Med. J. **1890**. 1063.)

Easily sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ (Koene.)

Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$

Insol. in alcohol.

Decomp. into basic salt by boiling H_2O . (Seubert, Arch. Pharm. **229**. 1.)

Zinc sulphite ammonia, $\text{ZnSO}_3 \cdot \text{NH}_3$.

Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Rammelsberg, Pogg. **67**. 255.)

Zirconium sulphite.

Insol. in H_2O . Somewhat sol. in $\text{H}_2\text{SO}_3 + \text{Aq.}$ from which it is repptd. on boiling. Sol. in $(\text{NH}_4)_2\text{SO}_3 + \text{Aq.}$ from which Zr hydroxide is pptd. on boiling. (Berzelius.)

$\text{Zr}(\text{SO}_3)_2 + 7\text{H}_2\text{O}$. Ppt. (Venable, J. Am. Chem. Soc. 1895, **17**. 449.)

Sulphuryl bromide, SO_2Br_2 .

(Odling, Chem. Soc. **7**. 2.)

Does not exist. (Sestini, Bull. Soc. **10**. 226; Melsens, C. R. **76**. 92; Michaelis.)

Sulphuryl chloride, SO_2Cl_2 .

Decomp. by H_2O and alcohol.

Decomp. by moist air, water, or abs. alcohol; more rapidly by alkalies, HCl , SO_2 , etc. (Schiff, A. **102**. 111.)

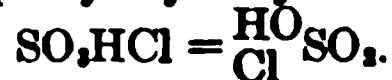
$+ \text{H}_2\text{O}$. Only sl. sol. in H_2O at 0° with slow decomp. (Baeyer, B. 1901, **34**. 737.)

$+ 15\text{H}_2\text{O}$. Sl. sol. in H_2O at 0° and stable therein for several hours. (Baeyer.)

Disulphuryl chloride (Pyrosulphuryl chloride), $\text{S}_2\text{O}_5\text{Cl}_2$.

Decomp. slowly with H_2O . (Rose, Pogg. **44**. 291.)

Sol. in CCl_4 and CHCl_3 ; miscible with liquid SO_2 .

Sulphuryl hydroxyl chloride,

Decomp. on moist air, and violently with H_2O . Not miscible with CS_2 . Decomp. with alcohol.

Sulphuryl titanium chloride, $\text{SO}_2 \cdot \text{TiCl}_4 = \text{TiCl}_3\text{OSO}_2\text{Cl}$.

Slowly deliquescent. (Clausnitzer, B. **11**. 2011.)

Disulphuryl chloride stannic oxychloride,
 $5\text{S}_2\text{O}_2\text{Cl}_2, 4\text{SnOCl}_2.$

Sol. in a little H_2O , but decomp. by more H_2O . (Rose, Pogg. 44. 320.)

Sulphuryl fluoride, SO_2F_2 .

1 pt. is sol. in 10 pts. H_2O at 9° . 3 vol. are sol. in 1 vol. alcohol at 9° ; insol. in conc. H_2SO_4 at 66° ; sol. in aq. solution of KOH , $\text{Ca}(\text{OH})_2$, $\text{Ba}(\text{OH})_2$, and in alcoholic solution of alkalis. (Moissan, C. R. 1901, 132. 377.)

Sulphuryl hydroxyl fluoride, HSO_3F .

Violently decomp. by H_2O . (Thorpe and Kirwan, Z. anorg. 3. 63.)

Sulphuryl peroxide, SO_4 .

See Sulphur heptoxide.

Sulphydric acid.

See Hydrogen Sulphide.

Sulphydroxyl.

See Sulphhydroxyl.

Tantalalic acid, $\text{H}_4\text{Ta}_2\text{O}_7$ (?).

Sol. in HF (Rose), and $\text{KH}_2(\text{C}_2\text{O}_4)_2 + \text{Aq}$ (Gahn, Schw. J. 16. 437). At the instant of precipitation is sol. in various acids. (Rose.)

Aluminum tantalate.

Insol. in H_2O . (Berzelius.)

Ammonium hexatantalate, $(\text{NH}_4)_2\text{H}_7\text{Ta}_7\text{O}_{19} + \text{H}_2\text{O}$.

Somewhat sol. in H_2O . (Rose, Pogg. 102. 57.)

Barium hexatantalate, $\text{Ba}_4\text{Ta}_6\text{O}_{19} + 6\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Rose.)

Cæsium tantalate, $4\text{Cs}_2\text{O}, 3\text{Ta}_2\text{O}_5 + 14\text{H}_2\text{O}$.

Completely sol. in a small amount of hot H_2O . (E. F. Smith, J. Am. Chem. Soc. 1908, 30. 1666.)

$7\text{Cs}_2\text{O}, 6\text{Ta}_2\text{O}_5 + 38\text{H}_2\text{O}$. Pptd. from its aqueous solution by alcohol. (Smith.)

Ferrous tantalate, $\text{Fe}(\text{TaO}_4)_2$.

Min. *Tantalite*.

$5\text{FeO}, 4\text{Ta}_2\text{O}_5$. Min. *Tapiolite*.

Magnesium hexatantalate, $\text{Mg}_4\text{Ta}_6\text{O}_{19} + 9\text{H}_2\text{O}$.

Ppt. (Rose, Pogg. 102. 61.)

$4\text{MgO}, \text{Ta}_2\text{O}_5$. Insol. in H_2O . (Joly, C. R. 81. 266.)

Mercurous tantalate, $5\text{Hg}_2\text{O}, 4\text{Ta}_2\text{O}_5 + 5\text{H}_2\text{O}$.

Decomp. by warm $\text{HNO}_3 + \text{Aq}$ (1.21 sp. gr.) with separation of Ta_2O_5 . (Rose, Pogg. 102. 64.)

Potassium tantalate, KTaO_4 .

Insol. in H_2O . Sol. in $\text{KOH} + \text{Aq}$. (Marignac, A. ch. (4) 9. 249.)

Potassium hexatantalate, $\text{K}_2\text{Ta}_6\text{O}_{19} + 16\text{H}_2\text{O}$.

Sol. without decomp. in moderately warm H_2O . Decomp. by boiling. (Marignac, A. ch. (4) 9. 259.)

Rubidium tantalate, $4\text{Rb}_2\text{O}, 3\text{Ta}_2\text{O}_5 + 14\text{H}_2\text{O}$.

Sol. in H_2O . (E. F. Smith, J. Am. Chem. Soc. 1908, 30. 1666.)

Silver tantalate, $4\text{Ag}_2\text{O}, 3\text{Ta}_2\text{O}_5$.

Completely sol. in $\text{NH}_4\text{OH} + \text{Aq}$. $\text{HNO}_3 + \text{Aq}$ dissolves Ag_2O , and Ta_2O_5 separates out. (Rose, Pogg. 102. 64.)

Sodium tantalate, NaTaO_4 .

Insol. in H_2O . (Rose.)

Sodium hexatantalate, $\text{Na}_6\text{Ta}_6\text{O}_{19} + 25\text{H}_2\text{O}$.

1 pt. salt. dissolves in 493 pts. H_2O at 13.5° , and in 162 pts. at 100° . Very slightly sol. in alcohol. Insol. in alkaline solutions. (Rose.)

Pertantalalic acid.

See Pertantalalic acid.

Tantalum, Ta .

Not attacked by HCl , HNO_3 , aqua regia, or hot conc. H_2SO_4 . Easily sol. in a mixture of HNO_3 and HF (Berzelius, Pogg. 4. 6; Rose). Also sol. in HF alone (Berzelius.)

Not attacked by alkali hydrates + Aq .

Insol. in single acids and in aqua regia. Oxidized by a mixture of HF and aqua regia. (Moissan, C. R. 1902, 134. 211.)

Pure Ta is insol. in boiling H_2SO_4 , HNO_3 , HCl , aqua regia or mixtures of these acids; slowly sol. in $\text{HF} + \text{Aq}$. (v. Bolton, Zeit. Elektrochem. 1905, 11. 45.)

Tantalum bromide, TaBr_5 .

Decomp. by H_2O . (Rose.)

Tantalum dichloride, $\text{TaCl}_5 + 2\text{H}_2\text{O}$.

Sol. in H_2O when freshly prepared. (Chabrie, C. R. 1907, 144. 805.)

Tantalum pentachloride, TaCl_5 .

Takes up H_2O from the air without deliquescing. Decomp. by H_2O . Sol. in H_2SO_4 . Sol. in cold $\text{HCl} + \text{Aq}$ to a cloudy liquid, which gelatinises after a time. Not completely sol. in boiling $\text{HCl} + \text{Aq}$, and the solution does not gelatinise by the subsequent addition of water, but all goes into solution. Partly sol. in $\text{KOH} + \text{Aq}$. Insol. in $\text{K}_2\text{SO}_4 + \text{Aq}$. Sol. in absolute alcohol.

Tantalum pentafluoride, TaF₅.

Very hygroscopic; sol. in H₂O. (Ruff, B. 1909, 42. 494.)

Tantalum fluoride with MF.

See Fluotantalate, M.

Tantalum hydroxide, Ta₂O₅, xH₂O.

See Tantallic acid.

Tantalum nitride, TaN.

Not sol. in any acids, except a mixture of HF and HNO₃. (Rose, Pogg. 100. 146.)

Ta₃N₅. (Joly, Bull. Soc. (2) 25. 506.)

Tantalum dioxide, Ta₂O₅(?).

Sol. in HF with evolution of hydrogen. (Hermann, J. pr. (2) 5. 69.)

Tantalum tetroxide, Ta₂O₄.

Not attacked by any acid, not even a mixture of HNO₃ and HF. (Berzelius, Pogg. 4. 20.)

Decomp. by HCl. (Smith, Z. anorg. 1894, 7. 98.)

Tantalum pentoxide, Ta₂O₅.

Insol. in any acid, even boiling H₂SO₄ or in HF. (Berzelius.)

Sol. in fused KHSO₄, 10 pts. being necessary to dissolve 1 pt. Ta₂O₅.

Tantalum silicide, TaSi₂.

Insol. in most inorganic acids. Sol. in HF and in HF + HNO₃.

Decomp. by fused alkali hydroxides. (Hönigschmid, M. 1907, 28. 1027.)

Tantalum sulphide, Ta₂S₄.

Not attacked by HCl + Aq. Oxidised by boiling with HNO₃ + Aq, more rapidly with aqua regia. Attacked by H₂SO₄ on heating. Not completely sol. in HF or a mixture of HF and HNO₃.

Telluretted hydrogen, TeH₂.

See Hydrogen telluride.

Telluric acid, H₂TeO₄.

Insol. in H₂O, cold conc. HCl, hot HNO₃, or boiling KOH + Aq, but when heated with H₂O is gradually converted into H₂TeO₄ + 2H₂O and dissolved.

+ 2H₂O. Very slowly sol. in cold H₂O, but sol. in hot H₂O in every proportion. Insol. in absolute alcohol; sol. in dil. alcohol according to the amount of H₂O present. Sol. in acids and alkalies. Insol. in alcohol or ether.

Insol. in alcohol; sol. in NaOH + Aq. (Mylius, B. 1901, 34. 2216.)

Stable in the air.

Sol. in H₂O; pptd. by HNO₃. (Staudenmaier, Z. anorg. 1895, 10. 191.)

Solubility in H₂O.

Solid phase	Temp.	% H ₂ TeO ₄	Mols H ₂ O to 1 mol H ₂ TeO ₄	Mols H ₂ TeO ₄ to 100 mols H ₂ O
H ₂ TeO ₄ + 6H ₂ O	0°	13.92	66.2	1.51
"	5°	17.84	49.2	2.03
"	10°	26.21	30.2	3.31
"	15°	32.79	21.9	4.55
H ₂ TeO ₄ · 2H ₂ O	10°	25.29	31.7	3.15
"	18°	28.90	26.2	3.82
"	30°	33.36	21.4	4.67
"	40°	36.38	18.8	5.33
"	60°	43.67	14.2	7.04
"	80°	51.55	10.07	9.93
"	100°	60.84	6.89	14.52

(Mylius, B. 1901, 34. 2211.)

+ 6H₂O. Obtained from solutions at 0°. (Staudenmaier, Z. anorg. 1895, 10. 191.)

Allotelluric acid, H₂TeO₄.

Miscible with H₂O.

Sol. in alcohol; pptd. by NaOH + Aq but sol. in excess. (Mylius, B. 1901, 34. 2216.)

Tellurates.

Neutral alkali salts are sol. in H₂O; the acid salts are only sl. sol. therein, but dissolve in HCl + Aq.

Aluminum tellurate.

Ppt. Sol. in excess of aluminum salts + Aq. (Berzelius.)

Ammonium tellurate, (NH₄)₂TeO₄.

Slowly but completely sol. in H₂O. Sl. sol. in NH₄OH + Aq or NH₄Cl + Aq. Sl. sol. in alcohol. (Berzelius.)

(NH₄)₂O, 2TeO₃. Sl. sol. in H₂O, but more sol. than the corresponding K salt.

(NH₄)₂O, 4TeO₃. Very sl. sol. in H₂O. Insol. in alcohol. (Berzelius.)

Barium tellurate, BaTeO₄ + 3H₂O.

Sl. sol. in cold, more in boiling H₂O. Easily sol. in HNO₃ + Aq. (Berzelius.)

BaH₂(TeO₄)₂ + 2H₂O. More sol. in H₂O than BaTeO₄. Decomp. by H₂O. (Berzelius.)

BaO, 4TeO₃. More sol. in H₂O than either BaTeO₄ or BaH₂(TeO₄)₂. (Berzelius.)

Bismuth tellurate, Bi₂TeO₆ + 2H₂O.

Min. *Montanite*. Sol. in HCl + Aq with evolution of Cl.

Cadmium tellurate, CdTeO₄.

Ppt. Sol. in HCl + Aq. (Oppenheim.)

Cæsium hydrogen tellurate, CsHTeO₄ + ½H₂O.

1 pt. is sol. in 30 pts. H₂O. (Norris, Am. Ch. J. 1901, 28. 321.)

- Calcium tellurate, CaTeO_4 .**
Ppt. Sol. in hot H_2O . (Berzelius.)
- Chromic tellurate, $\text{Cr}_2(\text{TeO}_4)_3$.**
Ppt. Sol. in excess of Cr salts + Aq.
- Cobaltous tellurate.**
Ppt. (Berzelius.)
- Cupric tellurate, CuTeO_4 .**
Ppt. (Berzelius.)
 CuO , 2TeO_3 . Ppt. (B.)
 Cu_2TeO_6 . Insol. in H_2O .
Sol. in HCl , HNO_3 , NH_4OH , KCN and acetic acid. (Hutchins, J. Am. Chem. Soc. 1905, 27. 1181.)
- Glucinum tellurate, GlTeO_4 .**
Insol. in H_2O .
- Iron (ferrous) tellurate, FeTeO_4 .**
Ppt. Min. *Ferrotellurate*.
- Iron (ferric) tellurate, $\text{Fe}_2(\text{TeO}_4)_3$.**
Ppt. Sol. in ferric salts + Aq. (Berzelius.)
- Lead tellurate, basic.**
Not completely insol. in H_2O .
- Lead tellurate, PbTeO_4 .**
Somewhat sol. in H_2O .
 PbO , 2TeO_3 . More sol. than PbTeO_4 .
 PbO , 4TeO_3 . Sl. sol. in H_2O . Sol. in HNO_3 + Aq, less sol. in $\text{HC}_2\text{H}_3\text{O}_2$ + Aq. (Berzelius.)
- Lithium tellurate, $\text{Li}_4\text{TeO}_6 + x\text{H}_2\text{O}$.**
Sl. sol. in H_2O with decomp. (Mylus, B. 1901, 34. 2209.)
- Magnesium tellurate, MgTeO_4 .**
Ppt. More sol. in H_2O than the Ba, Sr. or Ca salts.
 MgTe_2O_7 . More sol. in H_2O than MgTeO_4 .
- Manganous tellurate.**
Ppt.
- Mercurous tellurate, basic, $3\text{Hg}_2\text{O}$, 2TeO_3 .**
Ppt. (Hutchins, J. Am. Chem. Soc. 1905, 27. 1178.)
- Mercurous tellurate, Hg_2TeO_4 .**
Ppt. Min. *Magnolite*.
- Mercuric tellurate, HgTeO_4 .**
Ppt. Very easily decomp. by H_2O . (Hutchins, J. Am. Chem. Soc. 1905, 27. 1179.)
+ $2\text{H}_2\text{O}$. Slowly decomp. by cold H_2O .
Rapidly decomp. by boiling H_2O . (Hutchins.)
 Hg_2TeO_6 . Insol. in H_2O . Unchanged by boiling with H_2O .
- Sol. in HNO_3 , but more readily sol. in HCl . (Hutchins.)
- Mercuric tellurate.**
Ppt. (Berzelius.)
- Mercurous hydrogen tellurate, $\text{HgHTeO}_4 + 3\text{H}_2\text{O}$.**
Stable in the air if protected from the light
Insol. in H_2O . Decomp. by boiling H_2O or by an excess of cold conc. HgNO_3 + Aq.
Sol. in dil. HNO_3 or dil. acetic acid. (Hutchins, J. Am. Chem. Soc. 1905, 27. 1177.)
- Nickel tellurate.**
Ppt.
- Potassium tellurate, $\text{K}_2\text{TeO}_4 + 5\text{H}_2\text{O}$.**
Deliquesces. Sol. in H_2O . Very sl. sol. in H_2O containing KOH .
100 g. H_2O dissolve at:

0°	20°	30°
8.82	27.53	50.42 g. K_2TeO_4

(Rosenheim and Weinheber, Z. anorg. 1911, 69. 264.)
- Insol. in alcohol. (Berzelius.)
 K_2O , 2TeO_3 . Insol. in H_2O , acids. or alkalies. (B.)
 $\text{KHTeO}_4 + \frac{1}{2}\text{H}_2\text{O}$. Sl. sol. in cold, more sol. in hot H_2O . (Berzelius.)
 K_2O , $3\text{TeO}_3 + 5\text{H}_2\text{O}$. Much more sol. in hot than in cold H_2O . (Hutchins, J. Am. Chem. Soc. 1905, 27. 1174.)
 K_2O , 4TeO_3 . Insol. in H_2O , HCl , or HNO_3 + Aq. Sol. by long heating with conc. HNO_3 + Aq.
 KHTeO_4 , $\text{H}_2\text{TeO}_4 + \frac{1}{2}\text{H}_2\text{O}$. Sl. sol. in H_2O .
- Rubidium tellurate, $\text{Rb}_2\text{TeO}_4 + 3\text{H}_2\text{O}$.**
Sol. in about 10 pts. H_2O . (Norris, Am. Ch. J. 1901, 26. 322.)
- Rubidium hydrogen tellurate, $\text{RbHTeO}_4 + \frac{1}{2}\text{H}_2\text{O}$.**
Sol. in about 20 pts. cold H_2O . Sl. more sol. in hot H_2O . (Norris, Am. Ch. J. 1901, 26. 320.)
- Silver tellurate, $3\text{Ag}_2\text{O}$, TeO_3 .**
Sol. in NH_4OH + Aq.
 $3\text{Ag}_2\text{O}$, 2TeO_3 . Insol. in boiling H_2O .
+ $3\text{H}_2\text{O}$. Ppt. Unchanged by cold H_2O .
Gradually decomp. by boiling H_2O . (Hutchins, J. Am. Chem. Soc. 1905, 27. 1169.)
 Ag_2TeO_4 . Decomp. by H_2O into $3\text{Ag}_2\text{O}$, TeO_3 . Sol. in NH_4OH + Aq.
+ $2\text{H}_2\text{O}$. Insol. in hot and cold H_2O . Sol. in NH_4OH , KCN , $\text{Na}_2\text{S}_2\text{O}_3$, HNO_3 , H_2SO_4 , and $\text{HC}_2\text{H}_3\text{O}_2$ + Aq. Decomp. by conc. HNO_3 , H_2SO_4 or acetic acid. (Hutchins, J. Am. Chem. Soc. 1905, 27. 1165.)

Ag_2TeO_7 . Ppt.
 Ag_2O , 4TeO_3 . Ppt.
 Could not be obtained. (Hutchins, J. Am. Chem. Soc. 1905, **27**, 1168.)

Sodium tellurate, $\text{Na}_2\text{TeO}_4 + 2\text{H}_2\text{O}$.

Very sl. sol. in hot or cold H_2O . When heated to drive off $2\text{H}_2\text{O}$ becomes insol. in H_2O , but sol. in dil. $\text{HNO}_3 + \text{Aq.}$ (Berzelius.)
 1 pt. is sol. in about 130 pts. H_2O at 18° ; 50 pts. H_2O at 100° .

+ $4\text{H}_2\text{O}$. 1 pt. is sol. in about 70 pts. H_2O at 18° ; 40 pts. H_2O at 50° . (Mylius, B. 1901, **34**, 2209.)

$\text{Na}_2\text{Te}_2\text{O}_7 + 4\text{H}_2\text{O} = \text{Na}_2\text{HTeO}_4 + 1\frac{1}{2}\text{H}_2\text{O}$.
 Slowly but completely sol. in H_2O . Sl. sol. in $\text{NaC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ Insol. in alcohol. (Berzelius.)

Na_2O , 4TeO_3 . Insol. in H_2O , acids, or alkalies, except by long boiling with $\text{HNO}_3 + \text{Aq.}$

+ $x\text{H}_2\text{O}$. (α) Slowly sol. in H_2O . (β) Insol. even in boiling H_2O .

$\text{Na}_4\text{TeO}_6 + 8\text{H}_2\text{O}$. Very sol. in H_2O but with decomp. (Mylius.)

Strontium tellurates.

Resemble Ca salts.

Thallous tellurate, Tl_2TeO_4 .

Sl. sol. in H_2O (Dennis, J. Am. Chem. Soc. 1898, **18**, 975.)

Thorium tellurate.

Ppt. Insol. in excess of thorium salts + Aq.

Uranium tellurate, $\text{U}_2(\text{Te}_2\text{O}_7)_2$ (?)

Ppt. Insol. in H_2O or $\text{UO}_2(\text{NO}_3)_2 + \text{Aq.}$

Yttrium tellurate.

Ppt. Insol. in H_2O or Yt salts + Aq.

Zinc tellurate, Zn_3TeO_8 .

Insol. in H_2O

Sol. in HNO_3 , HCl , H_2SO_4 and acetic acid. (Hutchins, J. Am. Chem. Soc. 1905, **27**, 1181.)

Zirconium tellurate.

Ppt. (Berzelius.)

Tellurium, Te.

Insol. in H_2O or $\text{HCl} + \text{Aq.}$ Sl. sol. in hot conc. H_2SO_4 , but separates out on cooling. Sol. in boiling conc. H_2SO_4 . Easily oxidised by HNO_3 or aqua regia. Sol. in boiling very conc. $\text{KOH} + \text{Aq.}$, separating out again on cooling.

Not attacked by boiling conc. $\text{HNO}_3 + \text{Aq.}$ according to Hartung-Schwartzkoff (Ann. Min. (4) **19**, 345).

Sol. in warm conc. $\text{KCN} + \text{Aq.}$

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**, 830.)

100 pts. methylene iodide dissolve 0.1 pt. Te at 12° . (Retgers, Z. anorg. **3**, 343.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0014 g. Te in 6 days. (Gates, J. phys. Ch. 1911, **15**, 143.)

A colloidal solution of Te in H_2O can be obtained. It exists in two modifications, a brown and a blue-gray. Both can be diluted with H_2O or concentrated by boiling without decomp. They are, however, decomp. by electrolytes, especially NH_4Cl . (Gutbier, Z. anorg. 1902, **32**, 53.)

Tellurium dibromide, TeBr_2 .

Decomp. on air or by H_2O . (Rose, Pogg. **21**, 443.)

Conc. tartaric acid dissolves partly without decomp. (Brauner, M. 1891, **12**, 34.)

Tellurium tetrabromide, TeBr_4 .

Sol. in a little, but decomp. by much H_2O .

Completely sol. in tartaric acid + Aq (1:1). (Brauner, M. 1891, **12**, 34.)

Tellurium hydrogen bromide, $\text{TeBr}_4 \cdot \text{HBr} + 5\text{H}_2\text{O}$.

Fumes in the air. Deliquescent. Stable in an atmos. of HBr . (Metsner, C. R. 1897, **124**, 1951.)

Tellurium dichloride, TeCl_2 .

Decomp. on air, or by H_2O or $\text{HCl} + \text{Aq.}$ (Rose, Pogg. **21**, 443.)

Tellurium tetrachloride, TeCl_4 .

Extremely deliquescent. Decomp. by cold H_2O , with separation of oxychloride and tellurous acid. Sol. in hot H_2O with decomp. Sol. in dil. $\text{HCl} + \text{Aq}$ without decomp. (Rose, Pogg. **21**, 443.)

Insol. in sulphur chloride and in CS_2 . (Lenher, J. Am. Chem. Soc. 1902, **24**, 188.)

Tellurium hydrogen chloride, $\text{TeCl}_4 \cdot \text{HCl} + 5\text{H}_2\text{O}$.

Easily decomp. (Metsner, C. R. 1897, **125**, 24.)

Tellurium chloride with MCl .

See Chlorotellurate, M.

Tellurium tetrachloride ammonia,

$\text{TeCl}_4 \cdot 3\text{NH}_3$.

Decomp. by H_2O . (Metsner, C. R. 1897, **124**, 33.)

$\text{TeCl}_4 \cdot 4\text{NH}_3$. Not deliquescent. Decomp. by H_2O . (Espenschied, J. pr. **80**, 480.)

Tellurium tetrachloride sulphur trioxide,
 $\text{TeCl}_4 \cdot \text{SO}_3$.

Ppt. (Prandtl, Z. anorg. 1909, **62**, 247.)

$\text{TeCl}_4 \cdot 2\text{SO}_3$. Decomp. by moisture. On heating at 120° , it gives $\text{TeCl}_4 \cdot \text{SO}_3$. (Prandtl.)

Tellurium tetrafluoride, TeF_4 .(Metzner, C. R. 1897, **125**. 25.) $+\text{H}_2\text{O}$. (Högbom, Bull. Soc. (2) **35**. 60.)**Tellurium hexafluoride, TeF_6 .**Decomp. by H_2O slowly but completely. (Prideaux, Chem. Soc. 1906, **39**. 322.)**Tellurium zirconium fluoride,**

See Fluozirconate, tellurium.

Tellurium diiodide, TeI_2 .Insol. in H_2O . (Rose, Pogg. **21**. 443.)**Tellurium tetraiodide, TeI_4 .**Insol. in cold, decomp. by hot H_2O or alcohol. Sol. in HI , but only sol. in $\text{MI} + \text{Aq}$. (Berzelius.)Data on solubility of TeI_4 in $\text{HI} + \text{I} + \text{Aq}$ are given by Menke (Z. anorg. 1912, **77**. 283.)**Tellurium hydrogen iodide, TeI_4 , $\text{HI} + 8\text{H}_2\text{O}$, and $+9\text{H}_2\text{O}$.**Deliquescent. (Metzner, A. ch. 1898, (7) **15**. 203.)**Tellurium nitride,**

Two forms.

a. Stable at ord. temp.

b. Unstable at ord. temp.

(Franz Fischer, B. 1910, **43**. 1472.) TeN . Not attacked by H_2O or dil. acetic acid.Insol. in liquid NH_3 . Decomp. by $\text{KOH} + \text{Aq}$. (Metzner, A. ch. 1898, (7) **15**. 203.)**Tellurium monoxide, TeO .**Sl. sol. in cold dil. HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$. Easily oxidised by $\text{HNO}_3 + \text{Aq}$ or aqua regia. Decomp. immediately by boiling conc. $\text{HCl} + \text{Aq}$. Slowly decomp. by $\text{KOH} + \text{Aq}$. (Divers and Shimosé, Chem. Soc. **35**. 563.)**Tellurium dioxide, TeO_2 .**Very sl. sol. in H_2O . Sl. attacked by acids. Sl. sol. in NH_4OH or alkali carbonates $+ \text{Aq}$. Easily sol. in NaOH or $\text{KOH} + \text{Aq}$. Not sol. in less than 150,000 pts. H_2O . Easily sol. in warm dil. $\text{HNO}_3 + \text{Aq}$. Sol. in warm $\text{H}_2\text{SO}_4 + \text{Aq}$. (Klein and Morel, Bull. Soc. (2) **43**. 203.)20% $\text{H}_2\text{SO}_4 + \text{Aq}$ dissolves on warming about 0.7%; 30% $\text{H}_2\text{SO}_4 + \text{Aq}$, about 0.85%; 50% $\text{H}_2\text{SO}_4 + \text{Aq}$, about 4.4%.These solutions are supersat. and TeO_2 separates from the more dil. acids on standing. (Brauner, M. 1891, **12**. 34.)

Min. Tellurite.

Tellurium dioxide hydrobromic acid, TeO_2 , 3HBr .(Ditte, C. R. **83**. 336.)**Tellurium dioxide hydrochloric acid, TeO_2 , 2HCl .**(Ditte, C. R. **83**. 336.) TeO_2 , 3HCl . (Ditte.)**Tellurium trioxide, TeO_3 .**Insol. in cold or hot H_2O , cold $\text{HCl} + \text{Aq}$, or cold or hot $\text{HNO}_3 + \text{Aq}$. Insol. in moderately conc. $\text{KOH} + \text{Aq}$, but, when the $\text{KOH} + \text{Aq}$ is very conc., is sol. if boiling.**Tellurium oxide, 2TeO_2 , TeO_3 .**

"Tellurium tellurate."

(Metzner, A. ch. 1898, (7) **15**. 203.)**Tellurium oxybromide.**Insol. in H_2O . (Ditte, A. ch. (5) **10**. 82.)**Tellurium oxybromide sulphur trioxide, TeOBr_2 , 2SO_3 .**Deliquescent. (Prandtl, Z. anorg. 1909, **62**. 247.)**Tellurium oxychloride, TeOCl_2 .**Insol. in H_2O . (Ditte.)**Tellurium oxyfluoride, TeF_4 , $\text{TeO}_2 + 2\text{H}_2\text{O}$.**Sol. in H_2O containing HNO_3 . Decomp. by H_2O . 2TeF_4 , $3\text{TeO}_2 + 6\text{H}_2\text{O}$. Decomp. by H_2O . (Metzner, C. R. 1897, **125**. 25.)**Tellurium sulphide, TeS .**Insol. in CS_2 ; very unstable. (Snelling, J. Am. Chem. Soc. 1912, **34**. 802.)**Tellurium disulphide, TeS_2 .**Insol. in H_2O or dil. acids. Sol. in alkali hydrates or sulphides $+ \text{Aq}$. CS_2 dissolves out S , so that the substance is probably a mixture. (Becker, A. **180**. 257.)**Tellurium trisulphide, TeS_3 .**Insol. in H_2O . Sol. in $\text{K}_2\text{S} + \text{Aq}$.**Tellurium sulphoxide, TeSO_3 .**Decomp. by H_2O . Sol. in H_2SO_4 . (Weber, J. pr. (2) **25**. 218.)Is tellurium sulphite. (Divers, Chem. Soc. **49**. 583.)**Tellurous acid, H_2TeO_3 .**Appreciably sol. in H_2O and acids. Sol. in alkali hydrates or carbonates $+ \text{Aq}$.**Tellurites.**The neutral and acid tellurites of the alkali metals are sol. in H_2O . Ba, Sr, Ca, and Mg tellurites are sl. sol., and the other salts insol. in H_2O . Most tellurites are sol. in $\text{HCl} + \text{Aq}$.

Aluminum tellurite.

Ppt. Insol. in Al salts + Aq. (Berzelius.)

Ammonium tellurite, $(\text{NH}_4)\text{HTeO}_3$, $\text{H}_2\text{TeO}_3 + 3\frac{1}{2}\text{H}_2\text{O}$.

Sol. in H_2O , from which it is precipitated by $\text{NH}_4\text{Cl} + \text{Aq}$ or alcohol. (Berzelius.)

Barium tellurite, BaTeO_3 .

Sl. sol. in H_2O when prepared in the moist way. (Berzelius.)

BaO , 4TeO_2 .

Cadmium tellurite.

Ppt. Sol. in HNO_3 , and $\text{HCl} + \text{Aq}$. (Oppenheim.)

Calcium tellurite, CaTeO_3 .

Sl. sol. in cold, more sol. in hot H_2O . (Berzelius.)

CaO , 4TeO_2 .

Chromium tellurite.

Ppt. Sol. in excess of chromic salts + Aq.

Cobaltous tellurite.

Ppt.

Cupric tellurite.

Insol. in H_2O . (Berzelius.)

Glucinum tellurite.

Insol. in H_2O .

Indium tellurite, $\text{In}_2(\text{TeO}_3)_2$, $2\text{In}(\text{OH})_3$.

Ppt. (Renz, Dissert. 1902.)

Ferrous tellurite.

Ppt.

Ferric tellurite.

Ppt.

Lead tellurite, PbTeO_3 .

Ppt. Easily sol. in acids. (Berzelius.)

Lithium tellurite, Li_2TeO_3 .

Sol. in H_2O . (Berzelius.)

Li_2O , 2TeO_2 . Decomp. by cold H_2O into Li_2TeO_3 and Li_2O , 4TeO_2 . (B.)

Li_2O , 4TeO_2 . Sol. in hot, much less in cold H_2O . (B.)

Magnesium tellurite, MgTeO_3 .

Precipitate. Much more sol. in H_2O than the Ba, Sr, or Ca salt. (Berzelius.)

Manganous tellurite.

Ppt.

Mercurous tellurite.

Ppt.

Mercuric tellurite.

Ppt.

Nickel tellurite.

Ppt.

Potassium tellurite, K_2TeO_3 .

Not deliquescent. Slowly sol. in cold, more quickly in boiling H_2O . (Berzelius.)

K_2O , 2TeO_2 . Completely sol. in boiling H_2O , from which K_2O , 4TeO_2 crystallises. (B.)

K_2O , $4\text{TeO}_2 + 4\text{H}_2\text{O}$. Decomp. by cold H_2O into K_2O , TeO_2 , and K_2O , 2TeO_2 , which dissolve, and H_2TeO_3 , which is insol. (B.)

Potassium hexatellurite, K_2O , $6\text{TeO}_3 + 2\text{H}_2\text{O}$.

Not decomp. by, but sl. sol. in H_2O . (Klein and Morel, C. R. 100. 1140.)

Silver tellurite, Ag_2TeO_3 .

Ppt. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Berzelius.)

The freshly pptd. salt is insol. in H_2O ; sol. in HNO_3 , H_2SO_4 , acetic and tartaric acid; decomp. by HCl . (Lenher, J. Am. Chem. Soc. 1913, 35. 727.)

AgHTeO_3 . Insol. in H_2O . Sol. in $\text{HNO}_3 + \text{Aq}$. (Rose, Pogg. 18. 60.)

Sodium tellurite, Na_2TeO_3 .

Slowly sol. in cold, more quickly in hot H_2O . Precipitated from aqueous solution by alcohol. (Berzelius.)

Na_2O , 2TeO_2 . Decomp. by H_2O as K salt. (B.)

Na_2O , $4\text{TeO}_2 + 5\text{H}_2\text{O}$. As above. (B.)

Strontium tellurite, SrTeO_3 .

Resembles Ba salt.

$\text{SrH}_2\text{Te}_4\text{O}_{10}$. Very sl. sol. in H_2O , more easily in $\text{HNO}_3 + \text{Aq}$.

Thorium tellurite.

Precipitate. Insol. in H_2O or Th salts + Aq.

Stannous tellurite.

Pptd. in presence of 60,000 pts. H_2O . (Fischer.)

Uranium tellurite, $\text{U}_2(\text{TeO}_3)_2$.

Ppt. Insol. in U salts + Aq.

Yttrium tellurite.

Precipitate.

Zinc tellurite, ZnTeO_3 .

Ppt.

Zirconium tellurite.

Ppt.

Terbium, Tb.

Metal has not been isolated.

Has been decomp. into two or more elements by Krüss (Z. anorg. 4. 27).

Terbium chloride, $\text{TbCl}_3 \cdot 6\text{H}_2\text{O}$.

Sol. in H_2O ; very hygroscopic; sol. in alcohol. (Urbain, C. R. 1908, 146. 128.)

Terbium hydroxide.

Sol. in dilute acids. Decomposes NH_4 salts + Aq.

Terbium oxide, T_2O_3 .

Sol. in dil. acids, even after ignition.

Terbium peroxide, Tb_2O_7 .

Sol. in HNO_3 and in hot HCl . (Urbain, C. R. 1907, 146. 127.)

Tetramine chromium compounds.

See—

Bromotetramine chromium compounds.

Chlorotetramine chromium compounds.

Iodotetramine chromium compounds.

Tetramine cobaltic compounds,

$\text{Co}(\text{NH}_3)_4\text{X}_3$.

See—

Bromotetramine cobaltic compounds.

Carbonatotetramine cobaltic compounds.

Chlorotetramine cobaltic compounds.

Croceocobaltic compounds.

Fuscocobaltic compounds.

Flavocobaltic compounds.

Iodotetramine cobaltic compounds.

Nitratotetramine cobaltic compounds.

Praseocobaltic compounds.

Roseotetramine cobaltic compounds.

Sulphatotetramine cobaltic compounds.

See also under octamine cobaltic salts for many tetramine salts as yet unclassified.

Tetramine cobaltic nitrite with MNO_2 ,

$\text{Co}_2(\text{NH}_3)_4(\text{NO}_2)_6, 2\text{MNO}_2$.

See Diamine cobaltic nitrite.

Tetrathionic acid, $\text{H}_2\text{S}_4\text{O}_6$.

Known only in aqueous solution.

Dil. solution can be boiled without decomp. Conc. solution decomp. by boiling.

Addition of H_2SO_4 or HCl makes solution more stable. (Fordos and Gélis, C. R. 15. 920.)

Tetrathionates.

Tetrathionates are all easily sol. in H_2O , but insol. in alcohol

Barium tetrathionate, $\text{BaS}_4\text{O}_6 + 2\text{H}_2\text{O}$.

Very sol. in H_2O , but precipitated by addition of alcohol

Cadmium tetrathionate.

Deliquescent. Solution in H_2O gradually decomposes. (Kessler, Pogg. 74. 249.)

Cæsium tetrathionate, $\text{Cs}_2\text{S}_4\text{O}_6$.

(J. Meyer, B. 1907, 40. 1361.)

Cuprous tetrathionate, $\text{Cu}_2\text{S}_4\text{O}_6$.

Decomp. by H_2O . (Chancel and Diacon, C. R. 1863, 56. 711.)

Cupric tetrathionate, CuS_4O_6 .

Sol. in H_2O .

Decomp. by long boiling. (Curtius and Henkel, J. pr. 1888, (2) 37. 148.)

Lead tetrathionate, $\text{PbS}_4\text{O}_6 + 2\text{H}_2\text{O}$.

Sol. in H_2O .

Manganous hydrogen tetrathionate,

$\text{MnH}_2(\text{S}_4\text{O}_6)_2$.

Deliquescent. Very sol. in H_2O and alcohol. (Curtius and Henkel, J. pr. (2) 37. 148.)

Nickel tetrathionate ammonia, $\text{NiS}_4\text{O}_6, 6\text{NH}_3$.

Ppt. Decomp. by H_2O . Insol. in alcohol. (Ephraim, B. 1913, 46. 3109.)

Potassium tetrathionate, $\text{K}_2\text{S}_4\text{O}_6$.

Soluble in H_2O . Insol. in alcohol.

Difficultly sol. in H_2O . (Kessler, Pogg. 1847, 74. 254.)

Rubidium tetrathionate, $\text{Rb}_2\text{S}_4\text{O}_6$.

Not hygroscopic. (J. Meyer, B. 1907, 40. 1356.)

Sodium tetrathionate, $\text{Na}_2\text{S}_4\text{O}_6$.

Sol. in H_2O . Precipitated therefrom by a great excess of alcohol. (Kessler, J. pr. 36. 13.)

+ $2\text{H}_2\text{O}$. (Berthelot, A. ch. (6) 17. 450.)

Strontium tetrathionate, $\text{SrS}_4\text{O}_6 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Kessler, Pogg. 74. 255.)

More sol. in H_2O than Ba salt.

Zinc tetrathionate.

Sol. in H_2O . (Fordos and Gélis.)

Zinc hydrogen tetrathionate, $\text{ZnH}_2(\text{S}_4\text{O}_6)_2$.

Extremely sol. in H_2O and alcohol. (Curtius and Henkel, J. pr. (2) 37. 147.)

Zinc tetrathionate ammonia, $\text{ZnS}_4\text{O}_6, 3\text{NH}_3$.

Ppt. (Ephraim, B. 1915, 48. 641.)

Thallic acid.**Potassium thallate.**

Known only in aqueous solution. (Carstanjen, J. pr. 101. 55.)

Does not exist. (Lepsius, Chem. Ztg. 1880, 1327.)

Thallium, Tl.

Not attacked by pure H_2O . Easily sol. in dil. H_2SO_4 or HNO_3 + Aq. Difficultly sol. in HCl + Aq. Absolute alcohol dissolves considerable quantity in a short time, also methyl alcohol, and acetic ether. (Böttger.)

Not easily attacked by HF + Aq. (Kuhlmann.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20, 830.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0424 g. Tl in 6 days. (Gates, J. phys. Chem. 1911, 15, 143.)

Thallium arsenide, $TlAs$.

Decomp. by H_2SO_4 . (Carstanjen.)

Thalious azoimide, TlN_3 .

Sl. sol. in H_2O .

0.1712 pt. is sol. in 100 pts. H_2O at 0° ;

0.1965 pt. is sol. in 100 pts. H_2O at 5° ;

0.3 pt. is sol. in 100 pts. H_2O at 16° .

Insol. in abs. alcohol and ether.

(Curtius, J. pr. 1898, (2) 58, 284.)

Thallothallic azoimide, TlN_3 , TlN_5 .

Explosive. Decomp. by hot H_2O and by acids. (Dennis, J. Am. Chem. Soc. 1896, 18, 973.)

Thalious bromide, $TlBr$.

Nearly insol. in cold, sl. sol. in boiling H_2O . (Willm, Bull. Soc. (2) 2, 89.)

1 l. H_2O dissolves 0.00869 g. mol. $TlBr$ at 68.5° . (Noyes, Z. phys. Ch. 6, 248.)

Sl. sol. in H_2O . 0.48×10^{-3} g. is dissolved in a liter of sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46, 603.)

1 l. H_2O dissolves 420 mg. $TlBr$ at 18° . (Kohlrausch, Z. phys. Ch. 1904, 50, 356.)

238 mg. $TlBr$ are contained in 1 l. sat. solution at 0.13° ; 289 mg. at 9.37° ; 423 mg. at 18° ; 579 mg. at 25.68° . (Kohlrausch, Z. phys. Ch. 1908, 64, 168.)

Solubility of $TlBr$ in $Tl(NO_3)$ + Aq at 68.5° .

g. mols. per l.		g. mols. per l.	
$TlNO_3$	$TlBr$	$TlNO_3$	$TlBr$
0	0.00869	0	2.469
0.0163	0.00410	4.336	1.164
0.0294	0.00289	7.820	0.821
0.0955	0.00148	25.400	0.420

(Noyes, Z. phys. Ch. 1890, 6, 248.)

Insol. in acetone (Naumann, B. 1904, 37, 4329); pyridine (Naumann, B. 1904, 37, 4610); acetone (Eidmann, C. C. 1899, II, 1014).

Thallic bromide, $TlBr_3$.

Deliquescent. Easily sol. in H_2O and alcohol. (Willm.)

Insol. in methyl acetate. (Naumann, B. 1909, 42, 3790.)

+ H_2O . Very unstable. Sol. in H_2O ,

alcohol and ether. (Meyer, Z. anorg. 1900, 24, 353.)

+ $4H_2O$. Very sol. in H_2O . (Thomas, C. R. 1902, 134, 546.)

Thallothallic bromide, $TlBr$, $TlBr_3$.

Decomp. by H_2O (Meyer, Z. anorg. 1900, 24, 354.)

$3TlBr$, $TlBr_3$. Decomp. by H_2O into $TlBr$ and $TlBr_3$.

Thallic hydrogen bromide, $TlBr_3$, HBr .

Very sol. in H_2O . (Thomas, C. R. 1902, 134, 546.)

Thallic bromide ammonia, $TlBr_3$, $3NH_3$.

Decomp. by H_2O

Thallium bromochloride, $TlClBr$.

Decomp. by H_2O . (Thomas, C. R. 1901, 132, 1489.)

$TlClBr_2 + 4H_2O$. Ppt.

$TlCl_2Br + 4H_2O$. Ppt. Decomp. by H_2O . (Thomas, C. R. 1902, 134, 546.)

$Tl_2Cl_2Br_4$. Decomp. by H_2O , H_2SO_4 or HNO_3 . (Thomas, C. R. 1900, 131, 894; C. R. 1901, 132, 1489.)

$Tl_2Cl_2Br_2$. Sol. in H_2O . (Thomas, C. R. 1901, 132, 82.)

$TlClBr_3$, $3TlCl$. Cryst. from H_2O containing HNO_3 . (Cushman, Am. Ch. J. 1900, 24, 222.)

$TlCl_3$, $3TlBr$. Sol. in H_2O without decomp. (Cushman.)

$TlBr_3$, $3TlCl$. Decomp. by H_2O . (Cushman.)

$TlBr_3$, $TlCl$. Sol. in H_2O with decomp. (Cushman.)

$TlCl_3$, $2TlBr$, $TlCl$. Sol. in H_2O . (Meyer, Z. anorg. 1900, 24, 355-360.)

$TlBr_3$, $2TlCl$, $TlBr$. Ppt. Decomp. by H_2O (Meyer.)

$(TlCl_3, TlCl)$, $2(TlBr_3, TlBr)$. Ppt. (Meyer.)

$2(TlCl_3, TlCl)$, $(TlBr_3, TlBr)$. Ppt. Decomp. by hot H_2O . (Meyer.)

Thallium bromofluoride, $TlFBr_3$.

Decomp. in moist air.

Sol. in abs. alcohol. (Gewecke, A. 1909, 366, 233.)

Thallium bromofluoride ammonia,

$TlFBr_3$, $4NH_3$.

Decomp. by moisture.

Difficultly sol. in abs. alcohol. (Gewecke, A. 1909, 366, 234.)

Thalious chloride, $TlCl$.

Solubility in pts. H_2O at t° , according to H = Hebbeling; C = Crookes; L = Lamy.

0°	15°	16°	18.5°
504	283.4	377	359
H	C	H	H
100°	100°	100°	100°
about 50	52.5	63	pts. H_2O .
L	C	H	

1 l. H₂O dissolves 0.0161 g. mol. TlCl at 25°. (Noyes, Z. phys. Ch. 6. 249.)
3.26×10² grams are dissolved in 1 liter of sat. solution at 20°. (Böttger, Z. phys. Ch. 1903, 46. 603.)
1 l. H₂O at 25° dissolves 0.01606 g. mol. TlCl. (Geffcken, Z. phys. Ch. 1904, 49. 296.)

Solubility in H₂O at t°.

100 cc. sat. solution contain at:

t°	0°	10°	20°	30°	40°	50°
g. TlCl	0.17	0.24	0.34	0.46	0.60	0.80
t°	60°	70	80°	90°	99.3°	
g. TlCl	1.02	1.29	1.60	1.97	2.41	

(Berkeley, Trans. Roy. Soc. 1904, 203, A, 208.)

1 l. H₂O dissolves 3.040 TlCl at 18°. (Kohlrausch, Z. phys. Ch. 1904, 50. 356.)
2.27 g. are dissolved in 1 l. of sat. solution at 9.54°; 3.05 g. at 17.7°; 3.97 g. at 25.76°. (Kohlrausch, Z. phys. Ch. 1908, 64. 168.)
0.01629 mol. is sol. in 1 l. H₂O at 25°. (Hill, J. Am. Chem. Soc. 1910, 32. 1385.)
0.01607 g. equiv. is sol. in 1 l. H₂O at 25°. (Bray and Winninghoff, J. Am. Chem. Soc. 1911, 33. 1665.)
Much less sol. in H₂O containing HCl or HNO₃.

Solubility in HCl+Aq at 25°. 1 l. dissolves g. mol. TlCl.

g. HCl added	TlCl	g. HCl added	TlCl
0	0.01610	0.1468	0.00316
0.0283	0.00836	1.000	0.00200
0.0560	0.00565

(Noyes, Z. phys. Ch. 6. 249.)

Solubility in HCl+Aq. at 25°.

Concentration of HCl, equivalents per liter	Solubility of TlCl, equivalents per liter
0	0.01612
0.025	0.00869
0.05	0.00585
0.10	0.00384
0.20	0.00254

(Noyes, Z. phys. Ch. 1892, 9. 614.)

Solubility in HNO₃+Aq at 25°.

Normality HNO ₃	Sp. gr. of the solution	g. TlCl dissolved per l.
0.000	0.996	3.952
0.4977	1.0184	5.937
1.0046	1.0359	6.883
2.0452	1.0705	8.143
4.017	1.1362	9.926

(Hill and Simmons, Z. phys. Ch. 1909, 67. 605.)

Nearly insol. in NH₄OH+Aq.
More sol. in K₂CO₃+Aq than in H₂O. 3.86 g. TlCl are sol. in 1 l. H₂O at 25°. 21.84 g. TlCl are sol. in 1 l. 5N-K₂CO₃+Aq at 25°. (Spencer and Le Pla, C. C. 1908, II. 198.)

Solubility in KNO₃+Aq at 25°.

Concentration of KNO ₃ , milliequivalents per l.	Solubility of TlCl milliequivalents per l.
10	16.07
20	17.16
50	18.26
100	19.61
300	23.13
1000	30.72

(Bray and Winninghoff, J. Am. Chem. Soc. 1911, 33. 1670.)

Solubility in K₂SO₄+Aq at 25°.

Concentration of K ₂ SO ₄ , milliequivalents per l.	Solubility of TlCl milliequivalents per l.
10	16.07
20	17.79
50	19.42
100	21.37
300	26.00
1000	34.16

(Bray and Winninghoff, J. Am. Chem. Soc. 1911, 33. 1670.)

Solubility in Tl₂SO₄+Aq at 25°.

Concentration of Tl ₂ SO ₄ , milliequivalents per l.	Solubility of TlCl milliequivalents per l.
10	16.07
20	10.34
50	6.77
100	4.68

(Bray and Winninghoff, J. Am. Chem. Soc. 1911, 33. 1670.)

Solubility of TlCl in salts+Aq at 25°.

Salt	Concentration of salt g. equiv. per l.	TlCl dissolved g. equiv. per l.
NH ₄ Cl	0.025	0.00872
	0.05	0.00593
	0.2	0.00271
CaCl ₂	0.025	0.00899
	0.05	0.00624
	0.10	0.00417
	0.20	0.00284

Solubility of TlCl in salts + Aq at 25° .—
Continued.

Salt	Concentration of salt g. equiv. per l.	TlCl dissolved g. equiv. per l.
CdCl_2	0.025	0.01040
	0.05	0.0078
	0.10	0.00578
	0.20	0.00425
CuCl_2	0.025	0.00905
	0.05	0.00614
	0.10	0.00422
	0.20	0.00291
MgCl_2	0.025	0.00904
	0.05	0.00618
	0.10	0.00413
	0.20	0.00275
MnCl_2	0.025	0.00898
	0.05	0.00617
	0.10	0.00412
	0.20	0.00286
KCl	0.025	0.00872
	0.05	0.00593
	0.1	0.00399
	0.2	0.00265
NaCl	0.025	0.00869
	0.05	0.00592
	0.10	0.00395
	0.20	0.00271
ZnCl_2	0.025	0.00899
	0.05	0.00627
	0.10	0.00412
	0.20	0.00281
TlClO_3	0.025	0.00897
TlNO_3	0.025	0.00883
	0.05	0.00626
	0.10	0.00423

(Noyes, Z. phys. Ch. 1892, 9. 609.)

Solubility of TlCl in salts + Aq at 25° .

Salt	Mols TlCl sol. in 1 liter of				
	0.5-N solution	N solution	2-N solution	3-N solution	4-N solution
NH_4NO_3	0.02587	0.03121	0.03966		
KNO_3	0.02566	0.03077	0.03904		
NaNO_3	0.02564	0.03054	0.03851	0.04544	0.05128
LiNO_3	0.02542	0.03035	0.03785	0.04438	...
KClO_3	0.02370				
NaClO_3	0.02320	0.02687	0.03060	0.03303	0.03850

(Geffcken, Z. phys. Ch. 1904, 49. 295.)

Insol. in alcohol. Easily sol. in hot HgCl_2 + Aq. (Carstanjen.)Solubility of TlCl in $\text{HC}_2\text{H}_3\text{O}_2$ + Aq at 25° .
(g. equiv. per l.)

Acid	TlCl
0.000	0.01629
0.5134	0.01580
1.013	0.01495
2.016	0.0132
4.180	0.0099
8.130	0.0054
11.49	0.0026
14.31	0.0012
16.01	0.0005

(Hill, J. Am. Chem. Soc. 1910, 32. 1189.)

Insol. in pyridine. (Naumann, B. 1904, 37. 4610); acetone. (Naumann, B. 1904, 37. 4329.)

Thallic chloride, TlCl_3 .*Anhydrous.*Easily sol. in H_2O and in most ord. solvents.

In contact with moist air, it rapidly becomes hydrated. (Thomas, C. R. 1902, 135. 1053.)

Difficultly sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Very sol. in acetone. (Renz, B. 1902, 35. 1110.)

Difficultly sol. in acetone. (Naumann, B. 1904, 37. 4328.)

+ H_2O . Deliquescent, and very easily sol. in H_2O . (Werther.)Deliquescent, and very easily sol. in H_2O . (Werther.)+ $4\text{H}_2\text{O}$. 86.2 pts. are sol. in 100 pts. H_2O at 17° . Sp. gr. of sat. aq. solution at 17° = 1.85. (Thomas, C. R. 1902, 135. 1052.)

Very hygroscopic. (Meyer, Z. anorg. 1900, 24. 336.)

Very sol. in alcohol and ether. (Meyer, Z. anorg. 1900, 24. 338.)

+ $7\frac{1}{2}\text{H}_2\text{O}$. Deliquescent. (Werther.)**Thallothallic chloride, 3TlCl , TlCl_3 .**1 pt. dissolves in pts. H_2O at t° , according to C = Crookes; H = Hebbeling; L = Lamy.

15°	17°	100°	100°
380.1	346	52.9	20–25 pts. H_2O .
C	H	C	L

Sl. decomp. by dissolving. (Lamy.)

Thallic hydrogen chloride, TlCl_3 , HCl + $3\text{H}_2\text{O}$.

Very hygroscopic.

Decomp. by H_2O . (Meyer, Z. anorg. 1900, 24. 337.)**Thallium tungsten chloride, $\text{Tl}_3\text{W}_2\text{Cl}_9$.**Nearly insol. in H_2O .Sol. in a hot mixture of equal pts. H_2O and conc. HCl .

Sl. sol. in conc. HCl.
Nearly insol. in most organic solvents.
(Olsson, B. 1913, 46. 575.)

Thallic zinc chloride, $2\text{TlCl}_3, \text{ZnCl}_2 + 6\text{H}_2\text{O}$.

Can be cryst. from H_2O . (Gewecke, A. 1909, 366. 224.)

Thallic chloride ammonia, $\text{TlCl}_3, 3\text{NH}_3$.

Decomp. by H_2O . Sol. in $\text{HCl} + \text{Aq}$ (Willm.)

Thallium chlorofluoride, TlFCl_2 .

Very hygroscopic.

Decomp. by moist air.

Easily sol. in abs. alcohol. (Gewecke, A. 1909, 366. 230.)

+ $3\text{H}_2\text{O}$. Not hygroscopic.

Decomp. by H_2O , alcohol and ether. (Gewecke.)

Thallium chlorofluoride ammonia, $\text{TlFCl}_2, 4\text{NH}_3$.

Decomp. by H_2O .

Difficultly sol. in abs. alcohol and in ether. (Gewecke, A. 1909, 366. 232.)

Thallium chlorofluoride potassium chloride, $2\text{TlFCl}_2, \text{KCl}$.

Ppt. (Gewecke, A. 1909, 366. 231.)

Thallous fluoride, TlF .

Sol. in $1\frac{1}{4}$ pts. H_2O at 15° , and in much less hot H_2O . Difficultly sol. in alcohol. (Buchner, W. A. B. 52. 2. 644.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Deliquescent. (Willm.)

Thallic fluoride, TlF_3 .

Insol. in H_2O and cold $\text{HCl} + \text{Aq}$. (Willm.)

Cannot be obtained in pure state. (Gewecke, A. 1909, 366. 218.)

Thallous hydrogen fluoride, TlF, HF .

Sol. in 1 pt. H_2O . (Buchner.)

Thallous tungstyl fluoride,

See Fluoxytungstate, thallous.

Thallous vanadium fluoride.

See Fluovanadate, thallous.

Thallous vanadyl fluoride.

See Fluoxyvanadate, thallous.

Thallous hydroxide, TlOH .

Sol. in H_2O and alcohol.

Solubility of TlOH in H_2O at t° .

t°	g. equiv. TlOH per l	Sp. gr. 15°
0	1.151	1.231
18.5	1.554	1.317
19.5	1.582	1.322
29	1.803	1.342
23.1	1.861	1.377
33.1	1.967	1.400
36	2.075	1.417
40	2.240	1.446
44.5	2.442	...
54.1	2.940	...
59.4	3.281	...
64.6	3.601	...
78.5	4.673	...
90.0	5.705	...
99.2	6.708	...

(Bahr, Z. anorg. 1911, 71. 87.)

The solubility of Tl_2O in H_2O at temperatures is the same as that of $\text{TlOH} + \text{H}_2\text{O}$. (Willm, Bull. Soc. (2) 5. 354)

Thallic hydroxide, $\text{Tl}_2\text{O}_3, \text{H}_2\text{O} = \text{TlO}(\text{OH})$.

Insol. in H_2O . Sol. in dil. acids and ammonium salts + Aq. Insol. in caustic solutions.

$\text{Tl}(\text{OH})_3$. Easily sol. in dil. HCl or H_2O + Aq. (Carnegie, C. N. 60. 113.)

Thallous iodide, TlI .

Very sl. sol. in H_2O .

1 pt. TlI is sol. in pts. H_2O at t° . (According to Crookes; H = according to Hering; L = according to Lamy; W = according to Werther.)

35°	15°	16°	16-17°	19.4°
20,000	4450	16,000	11,676	14,654 pts.
W	C	L	H	W

20°	23.4°	45°	100°	100°
11,954	10,482	5407	842	804 pts. E
W	W	W	C	H

Sol. in 17,000 pts. H_2O at 20° . (Lo anal. 30. 342.)

Sat. solution at 20.15° contains 63. or 1.92×10^4 g. mol. TlI per l. (Böttg phys. Ch. 1903, 46. 603.)

1 l. H_2O dissolves 56 mg. TlI at 9.90° ; 56 mg. at 18.1° ; 84.7 mg. at 18.1° ; 84.7 mg. at 18.1° . (Kohlrausch, Z. phys. Ch. 1904, 50. 35)

36.2 mg. are dissolved in 1 l. of sat. sol. at 9.90° ; 56 mg. at 18.1° ; 84.7 mg. at 18.1° . (Kohlrausch, Z. phys. Ch. 1908, 64. 16)

Solubility in H_2O at $25^\circ = 1.76 \times 10^4$ per litre. (Spencer, Z. phys. Ch. 191 708.)

Not decomp. by dil. H_2SO_4 , HCl , or HNO_3 + Aq. Decomp. by hot dil. HNO_3 + Aq. cold conc. HNO_3 . Sol. in aqua regia.

Also less sol. in acetic acid than in H_2O . (Carstanjen.)

Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ (Werther.) Not wholly insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ and solubility is increased by presence of $(\text{NH}_4)_2\text{SO}_4$ or $\text{NH}_4\text{Cl.}$ (Baubigny, C. R. 113. 544.)

Sol. in 13,000 pts. $\text{NH}_4\text{OH} + \text{Aq.}$ ($6\frac{1}{2}\%$ or $2\frac{1}{2}\%$ NH_3). Sol. in 17,000 pts. $\text{NH}_4\text{OH} + \text{Aq.}$ ($1\frac{1}{4}\%$ NH_3). (Long.)

Insol. in dil. $\text{KI} + \text{Aq.}$ (1% KI). (Baubigny.)

Much more insol. in $\text{KI} + \text{Aq.}$ than in H_2O ; 1 pt. dissolves in 75,000 pts. dil. $\text{KI} + \text{Aq.}$ (Lamy.)

Nearly insol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq.}$ and absolutely insol. therein in presence of Pb salts. (Werther, C. N. 53. 51.)

Sol. in 56,336 pts. 85% alcohol at 13° . (Werther.) Sol. in 18,934 pts. 98% alcohol at 19° . (Hebberling.)

When TII is shaken with alcohol of 78°B (1 vol. $\text{H}_2\text{O} + 3$ vols. 98% alcohol) at 22° , and let stand with TII for 24 hours, and then evaporated to $\frac{1}{6}$ vol., there is shown no ppt. by $\text{NH}_4\text{SH} + \text{Aq.}$ (Baubigny.)

Sol. in 260,000 pts. 90% alcohol, and 37,000 pts. 50% alcohol at 20° . (Long.)

Insol. in methylene iodide. (Retgers, Z. anorg. 3. 343.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C., 1899, II. 1014.)

Insol. in pyridine. (Naumann, B. 1904, 37. 4601.)

Thallic iodide, TII_3 .

Sol. in alcohol.

Decomp. slowly in the air. (Wells, Z. anorg. 1894, 6. 313.)

Sol. in ether.

Thallothallic iodide, $\text{Tl}_3\text{I}_4 = 5\text{TII}, \text{TII}_3$.

Sol. in H_2O . (Jørgensen, J. pr. (2) 6. 82.)

Thallium nitride.

Very unstable. (Franz Fischer, B. 1901, 43. 1470.)

Thallous oxide, Tl_2O .

Deliquescent. Sol. in H_2O .

See Thallous hydroxide.

Thallic oxide, Tl_2O_3 .

Insol. in H_2O . Not attacked by cold H_2SO_4 . Sol. in hot H_2SO_4 . Sol. in cold $\text{HCl} + \text{Aq.}$

Insol. in alkalis + Aq. (Werther, J. pr. 91. 385.)

Black modification.

Less sol. in dil. acids than the brown modification. Solution is accompanied by a slight reduction to thallous salt.

More sol. in 10% HCl than in 10% H_2SO_4 or HNO_3 .

More sol. in conc. than in dil. acids. (Rabe, Z. anorg. 1906, 48. 431.)

Brown modification.

Easily sol. in dil. mineral acids on warming, with slight reduction to thallous salt.

More sol. in conc. acid. (Rabe.)

Thallium dioxide, TlO_2 .

Insol. in H_2O . (Piccini, Gazz. ch. it. 17. 450.)

Thallic oxide ammonia, $\text{Tl}_2\text{O}_3, 6\text{NH}_3$.

Decomp. by much H_2O . Insol. in alcohol. (Carstanjen.)

Thallic oxyfluoride, TlOF .

Insol. in H_2O . Slowly decomp. by boiling with H_2O .

Sol. in mineral acids. Almost insol. in HF . (Gewecke, A. 1909, 366. 226.)

Thallium phosphide (?).

Ppt. (Crookes.)

Thallous selenide, Tl_2Se .

Insol. in H_2O . Scarcely attacked by cold dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ but dissolves when heated. (Carstanjen.)

Thallothallic selenide.

Not attacked by cold conc. or boiling dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ Conc. H_2SO_4 decomposes. (Carstanjen.)

Thallous sulphide, Tl_2S .

Insol. in H_2O , $(\text{NH}_4)_2\text{S} + \text{Aq.}$, $\text{NH}_4\text{OH} + \text{Aq.}$, $\text{KCN} + \text{Aq.}$ and in alkali carbonates, and hydrates + Aq. Difficultly sol. in a solution of oxalic acid or acetic acid. (Crookes.) Easily sol. in HNO_3 , and $\text{H}_2\text{SO}_4 + \text{Aq.}$ Difficultly sol. in $\text{HCl} + \text{Aq.}$ (Willm.)

Sl. sol. in H_2O .

0.21×10^{-4} g. is dissolved in 1 l. sat. solution at 20° . (Böttger, Z. phys. Ch. 1903, 46. 603.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329.)

Thallic sulphide, Tl_2S_3 .

Insol. in H_2O . Insol. in cold, sol. in warm dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ without separation of S . Sol. in other dilute acids with separation of S . (Carstanjen.)

Thallium pentasulphide, Tl_2S_5 .

Ppt. (Hofmann, B. 1903, 36. 3092.)

Thallothallic sulphide, $5\text{Tl}_2\text{S}, 3\text{Tl}_2\text{S}_3$.

Very slowly decomp. by cold dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$

$\text{Tl}_2\text{S}, \text{Tl}_2\text{S}_3$. (Carstanjen.)

$\text{Tl}_2\text{S}, 2\text{Tl}_2\text{S}_3$. Decomp. by dil. acids. (Schneider, J. pr. (2) 10. 55.)

Thallium telluride, Tl_2Te .

(Fabre, C. R. 106. 673.)

Thio-

For compounds with prefix thio-, see also under sulpho-.

Thioantimonic acid.

See Sulphantimonic acid.

Thioarsenic acid.

See Sulpharsenic acid.

Thiomolybdic acid.

See Sulphomolybdic acid.

Thionamic acid, $\text{NH}_2\text{SO}_2 = \text{NH}_2\text{SO}(\text{OH})$.

Very deliquescent, and sol. in H_2O .

H_2O solution decomp. gradually. (Rose, Pogg. 33. 275; 42. 425.)

Ammonium thionamate, $\text{NH}_2\text{SO}(\text{ONH}_2)$.

Deliquescent. Sol. in H_2O ; easily decomp. when in solution. (Rose.)

Very sol. in alcohol with decomp. Sl. sol. in dry ether. (Divers and Ogawa, C. C. 1900, I. 1259.)

Dithionic acid.

See Dithionic acid.

Trithionic acid.

See Trithionic acid.

Tetrathionic acid.

See Tetrathionic acid.

Pentathionic acid.

See Pentathionic acid.

Thionyl bromide, SOBr_2 .

Unstable.

Decomp. by H_2O . (Besson, C. R. 1896, 122. 322.)

Thionyl bromochloride, SOClBr .

Decomp. slowly in the cold, rapidly at 115° .

Decomp. by H_2O . (Besson C. R. 1896, 122. 321.)

Thionyl chloride, SOCl_2 .

Sol. in CHCl_3 , and C_6H_6 . (Oddo, Gazz. ch. it. 1899, 29. (2) 318.)

Thionyl fluoride, SOF_2 .

Decomp. by H_2O .

Sol. in AsCl_3 , C_6H_6 , ether and oil of turpentine. (Moissan, C. R. 1900, 130. 1439.)

Thiophosphamic acid, $\text{H}_2\text{PNH}_2\text{O}_2\text{S}$ (?).

Known only in its salts. (Gladstone and Holmes, Chem. Soc. (2) 3. 1.)

Cadmium thiophosphamate, $\text{CdPNH}_2\text{O}_2\text{S}$.

Sol. in dil. acids, and $\text{NH}_4\text{OH} + \text{Aq}$. (G. and H.)

Lead —, $\text{PbPNH}_2\text{O}_2\text{S}$.

Ppt. Sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Gladstone and Holmes, Chem. Soc. (2) 3. 1.)

Thiophosphodiamic acid, $\text{H}_2\text{PN}_2\text{H}_4\text{OS}$.

Known only in solution, which soon decomposes. (G. and H.)

Cadmium thiophosphodiamate, $\text{Cd}(\text{PN}_2\text{H}_4\text{OS})_2$.

Insol. in H_2O ; sol. in dil. acids, and $\text{NH}_4\text{OH} + \text{Aq}$. (G. and H.)

Cupric —, $\text{Cu}(\text{PN}_2\text{H}_4\text{OS})_2$.

Insol. in H_2O , dil. HCl , or $\text{NH}_4\text{OH} + \text{Aq}$. Sol. in $\text{KCN} + \text{Aq}$. (Gladstone and Holmes, Chem. Soc. (2) 3. 1.)

Lead —, $\text{Pb}(\text{PN}_2\text{H}_4\text{OS})_2$.

Insol. in H_2O . Sol. in dil. $\text{HNO}_3 + \text{Aq}$.

Nickel —, $\text{Ni}(\text{PN}_2\text{H}_4\text{OS})_2$.

Sol. in dil. acids, and $\text{NH}_4\text{OH} + \text{Aq}$. (Gladstone and Holmes, Chem. Soc. (2) 3. 1.)

Zinc —, $\text{Zn}(\text{PN}_2\text{H}_4\text{OS})_2$.

Ppt. Sol. in dil. acids, and $\text{NH}_4\text{OH} + \text{Aq}$. (Gladstone and Holmes.)

Thiophosphonitrile, PSN .

Not decomp. by cold H_2O . Slowly decomp. by boiling H_2O . Easily decomp. by boiling dil. HCl . (Stock, B. 1906, 39. 1974.)

Thiophosphoric acid, $\text{H}_2\text{PSO}_3 = \text{PS}(\text{OH})_2$.

Known only in its salts.

Ammonium magnesium thiophosphate, $\text{NH}_4\text{MgPSO}_3 + 9\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . (Kubierschky, J. pr. (2) 31. 100.)

Barium —, $\text{Ba}_3(\text{PSO}_3)_2$.

Insol. in H_2O . (Wurts, A. ch. (3) 20. 473.)

Cobalt —.

Insol. in H_2O , but partially decomp. when boiled therewith. (Wurts.)

Cupric —.

Insol. in H_2O ; very easily decomp. (Wurts.)

Ferric —.

Insol. in H_2O . (Wurts.)

Magnesium —, $\text{Mg}_3(\text{PSO}_3)_2 + 20\text{H}_2\text{O}$.

Sl. sol. in cold H_2O . (Kubierschky, J. pr. (2) 31. 99.)

Nickel thiophosphate.

Insol. in H_2O , but decomp. when boiled therewith. (Wurtz.)

Potassium —, K_2PSO_3 .

Very sol. in H_2O . Known only in aqueous solution. (Wurtz.)

Sodium —, $Na_2PSO_3 + 12H_2O$.

Easily sol. in boiling H_2O . Cryst. out on cooling. (Wurtz, A. ch. (3) 20. 472.)

Insol. in alcohol.

Strontium —.

Insol. in H_2O . (Wurtz.)

Dithiometaphosphoric acid.**Ammonium dithiometaphosphate, NH_4PS_2O .**

Decomp. by H_2O . (Stock, B. 1906, 39. 1990.)

Monothioorthophosphoric acid.**Monoammonium monothioorthophosphate, $O:P(SNH_4)(OH)_2$.**

Sol. in H_2O .

Insol. in alcohol. (Stock, B. 1906, 39. 1990.)

Triammonium monothioorthophosphate, $SNH_4.PO.(ONH_4)_2$.

(Stock.)

Dithioorthophosphoric acid.**Ammonium dithiophosphate, $(NH_4)_2PS_2O_3 + 2H_2O$.**

Sl. efflorescent. Sol. in H_2O . (Kubierschky, J. pr. (2) 31. 93.)

Ammonium magnesium —, $NH_4MgPS_2O_3 + 6H_2O$.

Sl. sol. in cold, H_2O . (Kubierschky.)

Barium —, $Ba_2(PS_2O_3)_2 + 8H_2O$.

Precipitate. (Kubierschky, J. pr. (2) 31. 103.)

+18 H_2O . As the *trithio* compound. (Ephraim, B. 1910, 43. 287.)

Calcium —.

Very easily decomposed. (Kubierschky.)

Sodium —, $Na_2PS_2O_3 + 11H_2O$.

Very sol. in H_2O . (Kubierschky, J. pr. (2) 31. 93.)

Trithioorthophosphoric acid.**Ammonium trithioorthophosphate, $(NH_4)_3PS_3O + H_2O$.**

(Stock, B. 1906, 39. 1985.)

Barium trithioorthophosphate, $Ba_2(PS_3O)_2 + 20H_2O$.

Decomp. by H_2O and dil. acids.

Sol. in conc. HNO_3 with oxidation of the sulphur to H_2SO_4 . (Ephraim, B. 1910, 43. 286.)

Magnesium —, $Mg_2(PS_3O)_2 + 20H_2O$.

Decomp. by H_2O and dil. acids. (Ephraim.)

Thiophosphorous acid.**Ammonium thiophosphite (?), $(NH_4)_4P_2S_2O_3 + 3H_2O$.**

Sol. in H_2O . (Lemoine, C. R. 98. 45.) +6 H_2O .

Sodium thiophosphite (?), $Na_4P_2S_2O_3 + 5H_2O = P_2O_3, 2Na_2S + 5H_2O$.

Sol. in H_2O . (Lemoine, C. R. 98. 45.)

$Na_4P_2S_2O_3 + 4H_2O = P_2O_3, 3Na_2O, 2H_2S + 2H_2O$. Sol. in H_2O . (Lemoine, *l. c.*)

Thiophosphoryl triamide, $PS(NH_2)_3$.

Rapidly decomp. by H_2O . Scarcely sol. in alcohol, ether, or CS_2 . (Chevrier, C. R. 66. 748.)

Metathiophosphoryl bromide, PS_2Br .

Decomp. by H_2O . Insol. in ether. (Michaelis, A. 164. 9.)

Orthothiophosphoryl bromide, $PSBr_3$.

Slowly decomp. by cold, rapidly by hot H_2O but volatile with only partial decomp. with steam. Easily sol. in ether, CS_2 , PCl_3 , PBr_3 . Decomp. by cold alcohol. Forms hydrate $PSBr_3 + H_2O$. (Michaelis, A. 164. 9.)

Pyrothiophosphoryl bromide, $P_2S_2Br_4$.

Decomp. by H_2O and alcohol. Sol. in CS_2 and ether. (Michaelis.)

Thiophosphoryl phosphorus bromide, $PSBr_3, PBr_3$.

Decomp. by H_2O into $PSBr_3$. (Michaelis.)

Thiophosphoryl dibromochloride, $PSClBr_2$.

Decomp. by H_2O and alkalis.

Fumes in the air. (Besson, C. R. 1896, 122. 1059.)

Thiophosphoryl bromodichloride, $PSCl_2Br$.

Decomp. by H_2O and alkalis. Reacts violently with HNO_3 . (Besson, C. R. 1896, 120. 1058.)

Thiophosphoryl chloride, $PSCl_3$.

Very slowly decomp. by H_2O , and may be distilled with steam without much decomp. Decomp. by alcohol. Miscible with CS_2 . (Baudrimont, J. pr. 87. 301.)

Sol. in CCl_4 and C_6H_6 . (Oddo, Gazz. ch. it. 1899, 29. (2) 318.)

Thiophosphoryl pentachloride, PS_2Cl_5 (?).

Decomp. by H_2O . Sol. in alkalis with residue of S. Attacked violently by HNO_3 , alcohol, ether, oil of turpentine. Miscible with CS_2 . (Gladstone, Chem. Soc. 3. 5.)

Thiophosphoryl fluoride, PSF_3 .

Slowly sol. in H_2O with decomp. Sl. sol. in ether.

Insol. in H_2SO_4 , CS_2 , or benzene. (Thorpe and Rodger, Chem. Soc. 55. 306.)

More sol. in KOH or $\text{NaOH} + \text{Aq}$ than in H_2O .

Thiophosphoryl iodide, P_2SI_2 .

Very sol. in CS_2 . Unstable when heated. Fumes in the air. (Besson, C. R. 1896, 122. 1201.)

Thiosulphuric (formerly Hyposulphurous) acid, $\text{H}_2\text{S}_2\text{O}_3$.

Known only in aqueous solution, which is extremely unstable, and decomposes very quickly after its formation. The time before decomposition is exactly proportional to the ratio of the weight of H_2O to the weight of $\text{H}_2\text{S}_2\text{O}_3$ present; i. e., if one solution contains twice as much H_2O for a given amt. of $\text{H}_2\text{S}_2\text{O}_3$ as a second solution, the first solution will decompose in twice the length of time. The length of time is about 20 secs. at 10° , and 2 secs. at 50° for conc. solutions, to 120 secs. at 10° and 12 secs. at 50° for very dilute solutions. (See Landolt (B. 16. 2958) for further figures; also Winkelmann (B. 18. 406).)

Thiosulphates.

The thiosulphates of the alkalis and of Ca and Sr are easily sol. in H_2O ; Ba and Sr salts are sl. sol. and the other salts insol. The salts of the metals dissolve in alkali thiosulphates + Aq. All are insol. in alcohol.

Double Salts of Thiosulphuric acid. It is impossible to determine whether substances of this class are true chem. individuals. Many described by Svensson and others are doubtless isomorphic mixtures, whose comp. depends on the temp. and conc. of the solution in which pptd. (Rosenheim, Z. anorg. 1900, 25. 72.)

Ammonium thiosulphate, $(\text{NH}_4)_2\text{S}_2\text{O}_3$.

Very deliquescent. Very sol. in H_2O .

Not deliquescent. (Fock and Klüss, B. 1889, 22. 3099.)

Crystallises with $\frac{1}{2}\text{H}_2\text{O}$. (Rammelsberg, Pogg. 56. 298.) Anhydrous. (Arppe, A. 96. 113.)

Insol. in alcohol. (Arppe.)

Sol. in acetone. (Eidmann, C. C. 1899, II. 1014.)

Difficultly sol. in acetone. (Naumann, B. 1904, 37. 4328.)

Ammonium cadmium thiosulphate, $3(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{CdS}_2\text{O}_3 + 3\text{H}_2\text{O}$.

Can be recryst. from warm H_2O . (Fock and Klüss, B. 23. 1758.)

+ H_2O . (F. and K.)

$(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{CdS}_2\text{O}_3$. (F. and K.)

Ammonium cuprous thiosulphate, $(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Less sol. in H_2O than $2(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 1\frac{1}{2}\text{H}_2\text{O}$. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 91.)

$2(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 1\frac{1}{2}\text{H}_2\text{O}$. Very sol. in H_2O . Insol. in alcohol. (Rosenheim and Steinhäuser.)

Ammonium cuprous thiosulphate cuprous iodide, $7(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3, 8\text{CuI} + 4\text{H}_2\text{O}$.

Insol. in H_2O . (Brun, C. R. 1892, 114. 668.)

Could not be obtained. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 107.)

Ammonium cuprous sodium thiosulphate ammonia, $3\text{Cu}_2\text{S}_2\text{O}_3, 4\text{Na}_2\text{S}_2\text{O}_3, (\text{NH}_4)_2\text{S}_2\text{O}_3, 6\text{NH}_3$.

Ppt. When dry is fairly stable in the air. Partially decomp. by H_2O .

Sol. in dil. H_2SO_4 or acetic acid. (Shinn, J. Am. Chem. Soc. 1904, 26. 948.)

Ammonium lead thiosulphate, $2(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{PbS}_2\text{O}_3 + 3\text{H}_2\text{O}$.

Easily and completely sol. in cold H_2O , but deposits PbS_2O_3 by standing or warming. (Rammelsberg, Pogg. 56. 312.)

Ammonium magnesium thiosulphate, $(\text{NH}_4)_2\text{Mg}(\text{S}_2\text{O}_3)_2 + 6\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O . (Kessler, Pogg. 74. 283.)

Not deliquescent. (Fock and Klüss, B. 23. 540.)

Ammonium mercuric thiosulphate, $4(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{HgS}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Sol. in H_2O , from which it is precipitated by alcohol. Extremely easily decomp. (Rammelsberg, Pogg. 56. 318.)

Ammonium potassium thiosulphate, $\text{NH}_4\text{KS}_2\text{O}_3$.

Sol. in H_2O . (Fock and Klüss, B. 23. 536.)

Ammonium silver thiosulphate, $2(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + x\text{H}_2\text{O}$.

Easily sol. in H_2O . Somewhat sol. in alcohol. (Herschel, Edinb. Phil. J. 1. 398.)

$(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + x\text{H}_2\text{O}$. Nearly insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$, from which it is reprecipitated by an acid. (Herschel.)

Ammonium zinc thiosulphate,
 $(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{ZnS}_2\text{O}_3 + \text{H}_2\text{O}.$

Very sol. in H_2O . (Rosenheim and Davidsohn, Z. anorg. 1904, **41**. 238.)

Ammonium thiosulphate ammonium cuprous bromide, $\text{CuBr}, \text{NH}_4\text{Br}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O . (Rosenheim, Z. anorg. 1900, **25**. 107.)

The double salts of ammonium thiosulphate with silver and copper haloids are true chemical compounds and may be recryst. from H_2O without decomp. (Rosenheim, Z. anorg. 1900, **25**. 100.)

Ammonium thiosulphate ammonium silver bromide, $\text{AgBr}, \text{NH}_4\text{Br}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O . (Rosenheim.)

Ammonium thiosulphate ammonium cuprous chloride, $\text{CuCl}, \text{NH}_4\text{Cl}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O and in $\text{NH}_4\text{OH} + \text{Aq}$. (Rosenheim.)

Ammonium thiosulphate ammonium silver chloride, $\text{AgCl}, \text{NH}_4\text{Cl}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in cold H_2O and $\text{NH}_4\text{OH} + \text{Aq}$.

Decomp. by boiling with H_2O and by dil. acids. (Rosenheim.)

Ammonium thiosulphate ammonium cuprous cyanide.

Composition not constant. (Rosenheim.)

Ammonium thiosulphate ammonium silver cyanide.

Composition not constant. (Rosenheim.)

Ammonium thiosulphate ammonium cuprous iodide, $\text{CuI}, \text{NH}_4\text{I}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O . (Rosenheim.)

Ammonium thiosulphate ammonium cuprous iodide, $4(\text{NH}_4)_2\text{S}_2\text{O}_3, \text{NH}_4\text{I}, \text{CuI}.$

Very sol. in H_2O . Decomp. by boiling. (Brun, C. R. 1892, **114**. 668.)

Ammonium thiosulphate ammonium silver iodide, $\text{AgI}, \text{NH}_4\text{I}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O . (Rosenheim.)

Ammonium thiosulphate ammonium cuprous sulphocyanide, $\text{CuSCN}, \text{NH}_4\text{SCN}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O . (Rosenheim.)

Ammonium thiosulphate ammonium silver sulphocyanide, $\text{AgSCN}, \text{NH}_4\text{SCN}, 4(\text{NH}_4)_2\text{S}_2\text{O}_3.$

Sol. in H_2O . (Rosenheim.)

Ammonium thiosulphate cuprous iodide,
 $(\text{NH}_4)_2\text{S}_2\text{O}_3, 2\text{CuI} + \text{H}_2\text{O}.$

Insol. in H_2O . (Brun, C. R. 1892, **114**. 668.)

Could not be obtained. (Rosenheim and Steinhäuser.)

Barium thiosulphate, $\text{BaS}_2\text{O}_3 + \text{H}_2\text{O}.$

Sl. sol. in H_2O . (Rose, Pogg. **21**. 437.)

Insol. in alcohol.

1 pt. cannot be dissolved in 2000 pts. H_2O .

Sol. in dil. $\text{HCl} + \text{Aq}$ without decomposition. (Herschel, **1819**.)

Pptd. from $\text{BaS}_2\text{O}_3 + \text{Aq}$ by dil. alcohol. (Sobrero and Selmi, A. ch. (3) **28**. 211.)

Insol. in acetone. (Naumann, B. 1904, **37**. 4329.)

Barium bismuth thiosulphate, $\text{Ba}_3[\text{Bi}(\text{S}_2\text{O}_3)_3]_2.$

Sol. in H_2O with decomp. (Hauser, Z. anorg. 1903, **35**. 9.)

Barium cadmium thiosulphate, $2\text{BaS}_2\text{O}_3, \text{CdS}_2\text{O}_3 + 8\text{H}_2\text{O}.$

Sl. sol. in H_2O . (Fock and Klüss, B. **23**. 1761.)

$3\text{BaS}_2\text{O}_3, \text{CdS}_2\text{O}_3 + 8\text{H}_2\text{O}$. Sl. sol. in H_2O .

Barium cuprous thiosulphate.

Easily sol. in hot, difficultly sol. in cold H_2O . (Cohen, Chem. Soc. **51**. 38.)

$2\text{BaS}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 7\text{H}_2\text{O}$. Nearly insol. in H_2O . (Vortmann, M. **9**. 165.)

Barium gold thiosulphate.

Sl. sol. in H_2O . Insol. in alcohol. (Fordos and Gélis.)

Barium lead thiosulphate.

Difficultly sol. in H_2O . (Rammelsberg, Pogg. **56**. 313.)

Barium thiosulphate chloride, $\text{BaS}_2\text{O}_3, \text{BaCl}_2 + 2\text{H}_2\text{O}.$

Sol. in H_2O . (Fock and Klüss, B. **23**. 3001.)

Bismuth caesium thiosulphate, $\text{Cs}_3\text{Bi}(\text{S}_2\text{O}_3)_3.$

Sol. in H_2O .

Insol. in alcohol. (Hauser, Z. anorg. 1903, **35**. 8.)

Bismuth potassium thiosulphate, $\text{K}_3\text{Bi}(\text{S}_2\text{O}_3)_3 + \frac{1}{2}\text{H}_2\text{O}.$

Solubility in H_2O .

100 cc. of the sat. solution contain 3.5 g. at 2° ; 7.0 g. at 18° . At 18° the solution decomposes.

More sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ than in pure H_2O .

Insol. in alcohol. (Hauser, Z. anorg. 1903, **35**. 5.)

$+ \text{H}_2\text{O}$. Sol. in H_2O . Insol. in alcohol. (Carnot, C. R. **83**. 390.)

Bismuth rubidium thiosulphate,
 $\text{Rb}_2\text{Bi}(\text{S}_2\text{O}_3)_3 + \frac{1}{2}\text{H}_2\text{O}.$ Sol. in H_2O .Insol. in acid alcohol. (Hauser, Z. anorg. 1903, **35**. 7.)+ H_2O . Sol. in H_2O . (Hauser, Z. anorg. 1903, **35**. 8.)**Bismuth sodium thiosulphate,**Very sol. in H_2O , and also in alcohol. (Carnot, C. R. **83**. 338.) $\text{Na}_2\text{Bi}(\text{S}_2\text{O}_3)_3$. Decomp. by H_2O . Easily sol. in 50% alcohol. (Hauser, Z. anorg. 1903, **35**. 3.)**Cadmium thiosulphate, $\text{CdS}_2\text{O}_3 + 2\text{H}_2\text{O}$.**Sol. in H_2O . Insol. in alcohol. (Vortmann and Padberg, B. **22**. 2638.)**Cadmium potassium thiosulphate, $3\text{CdS}_2\text{O}_3, 5\text{K}_2\text{S}_2\text{O}_3$.**Cannot be recryst. without decomp. (Fock and Klüss, B. **23**. 1753.) $\text{CdS}_2\text{O}_3, 3\text{K}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Can be crystallised from H_2O without decomp. (F. and K.)**Cadmium sodium thiosulphate, $\text{CdS}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 16\text{H}_2\text{O}$.**Not deliquescent. Sol. in H_2O . (Jochum, C. C. **1885**, 642.)+ $9\text{H}_2\text{O}$. (Vortmann and Padberg, B. **22**. 2639.)+ $3\text{H}_2\text{O}$. Deliquescent. (Fock and Klüss, B. **23**. 1157.) $2\text{CdS}_2\text{O}_3, \text{Na}_2\text{S}_2\text{O}_3 + 7\text{H}_2\text{O}$. (V. and P.) $3\text{CdS}_2\text{O}_3, \text{Na}_2\text{S}_2\text{O}_3 + 9\text{H}_2\text{O}$. (V. and P.)**Cadmium strontium thiosulphate, $\text{CdS}_2\text{O}_3, 3\text{SrS}_2\text{O}_3 + 10\text{H}_2\text{O}$.**(Fock and Klüss, B. **23**. 1763.)**Cæsium thiosulphate, $\text{Cs}_2\text{S}_2\text{O}_3$.**Easily sol. in H_2O . (Chabrié, C. R. 1901, **133**. 297.)+ $2\text{H}_2\text{O}$. Very hygroscopic. (J. Meyer, B. 1907, **40**. 1360.)**Cæsium cuprous thiosulphate, $\text{Cs}_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.**Easily sol. in H_2O with decomp. (J. Meyer, B. 1907, **40**. 1361.)**Cæsium lead thiosulphate, $\text{Cs}_2\text{S}_2\text{O}_3, \text{PbS}_2\text{O}_3 + 2\text{H}_2\text{O}$.**

Not hygroscopic.

 $2\text{Cs}_2\text{S}_2\text{O}_3, \text{PbS}_2\text{O}_3 + 3\text{H}_2\text{O}$. Hygroscopic. (J. Meyer.)**Cæsium magnesium thiosulphate, $\text{Cs}_2\text{S}_2\text{O}_3, \text{MgS}_2\text{O}_3 + 6\text{H}_2\text{O}$.**Easily sol. in H_2O . (Meyer.)**Cæsium silver thiosulphate, $2\text{Cs}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + 3\text{H}_2\text{O}$.**

Not hygroscopic.

Decomp. by hot H_2O . (J. Meyer.)**Calcium thiosulphate, $\text{CaS}_2\text{O}_3 + 6\text{H}_2\text{O}$.**Sol. in 1 pt. H_2O at 3° .Aqueous solution saturated at 10° has sp. gr. 1.300. Solution with sp. gr. 1.11437 at 15.5° contains 0.2081 of its weight in CaS_2O_3 .Decomp. on heating. Insol. in alcohol (sp. gr. 0.8234). (Herschel, A. ch. **14**. 355.)100 g. sat. solution contains 29.4 g. CaS_2O_3 at 9° , and 34.7 g. CaS_2O_3 at 25° . (Kremann and Rodemund, M. 1914, **35**. 1065.)**Solubility of $\text{CaS}_2\text{O}_3 + \text{Na}_2\text{S}_2\text{O}_3$ in H_2O .**

t°	% $\text{Na}_2\text{S}_2\text{O}_3$	% CaS_2O_3	Solid phase
9	0	29.4	$\text{CaS}_2\text{O}_3, 6\text{H}_2\text{O}$
	11.04	22.64	"
	25.21	15.84	" + $\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$
	31.01	7.70	$\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$
25	0	34.7	$\text{CaS}_2\text{O}_3, 6\text{H}_2\text{O}$
	9.24	29.69	"
	15.67	21.41	"
	18.34	25.18	"
	28.24	21.14	"
	30.19	20.33	" + $\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$
	31.24	18.43	$\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$
	35.04	11.61	"

(Kremann and Rodemund, M. 1914, **35**. 1065.)**Calcium lead thiosulphate, $2\text{CaS}_2\text{O}_3, \text{PbS}_2\text{O}_3 + 4\text{H}_2\text{O}$.**Decomp. by H_2O . (Rammelsberg.)**Calcium potassium thiosulphate, $\text{CaS}_2\text{O}_3, 3\text{K}_2\text{S}_2\text{O}_3 + 5\text{H}_2\text{O}$.**Sol. in H_2O . (Fock and Klüss, B. **24**. 3016.)**Calcium silver thiosulphate, $2\text{CaS}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + x\text{H}_2\text{O}$.**Easily sol. in H_2O ; less sol. in alcohol. $\text{CaS}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + x\text{H}_2\text{O}$. Sl. sol. in H_2O . abundantly in $\text{NH}_4\text{OH} + \text{Aq}$. (Herschel, 1819.)**Calcium sodium thiosulphate nitrate, $\text{CaNa}_2(\text{S}_2\text{O}_3)_2\text{NO}_3 + 11\text{H}_2\text{O}$.**(Kremann and Rodemund, M. 1914, **35**. 1065.)**Cobaltous thiosulphate, $\text{CoS}_2\text{O}_3 + 6\text{H}_2\text{O}$.**Sol. in H_2O . (Rammelsberg.)**Cobaltous sodium thiosulphate, $2\text{CoS}_2\text{O}_3, 5\text{Na}_2\text{S}_2\text{O}_3 + 25\text{H}_2\text{O}$.**Efflorescent. Sol. in H_2O . (Jochum.)

Could not be obtained by Vortmann and Padberg.

$\text{CoS}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 15\text{H}_2\text{O}$. Sol. in H_2O . (Vortmann and Padberg, B. 22. 2641.)

Cuprous thiosulphate, $\text{Cu}_2\text{O}, 3\text{S}_2\text{O}_3 + 2\text{H}_2\text{O} = \text{Cu}_2\text{H}_4(\text{S}_2\text{O}_3)_2$.

Sl. sol. in H_2O . Abundantly sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$, $\text{NH}_4\text{Cl} + \text{Aq}$, $\text{NH}_4\text{OH} + \text{Aq}$, or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Sol. in HCl or $\text{HNO}_3 + \text{Aq}$. (v. Hauer, W. A. B. 13. 443.)

Cuprous hydrazine thiosulphate,
 $\text{Cu}_2\text{S}_2\text{O}_3, (\text{N}_2\text{H}_4)_2\text{H}_2\text{S}_2\text{O}_3 + \frac{1}{2}\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in $\text{NH}_4\text{OH} + \text{Aq}$ and in dil. acids. (Ferratini, C. C. 1912, I. 1281.)

Cupric lead thiosulphate, $\text{Pb}(\text{CuS}_2\text{O}_3)_2 + 3\text{H}_2\text{O}$. (?)

Very sol. in H_2O and quickly decomp. (Girard, C. C. 1904, I. 253.)

Cuprous mercurous thiosulphate, $5\text{Cu}_2\text{S}_2\text{O}_3, 3\text{Hg}_2\text{S}_2\text{O}_3$.

Insol. or sl. sol. in cold, decomp. by boiling H_2O . $\text{HNO}_3 + \text{Aq}$ dissolves out Cu . (Rammelsberg, Pogg. 56. 319.)

Cuprous potassium thiosulphate, $\text{Cu}_2\text{S}_2\text{O}_3, \text{K}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Sl. sol. in H_2O ; decomp. on heating with pptn. of CuS . Easily sol. in $\text{K}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Rammelsberg, Pogg. 56. 321.)

$\text{Cu}_2\text{S}_2\text{O}_3, 2\text{K}_2\text{S}_2\text{O}_3$. Very sol. in cold H_2O ; insol. in $\text{K}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Cohen, Chem. Soc. 51. 39.)

$+3\text{H}_2\text{O}$. Scarcely sol. in cold, sol. with sl. decomp. in hot H_2O . Sol. in $\text{HCl} + \text{Aq}$ with evolution of SO_2 .

$\text{Cu}_2\text{S}_2\text{O}_3, 3\text{K}_2\text{S}_2\text{O}_3 + 3\text{H}_2\text{O}$. More sol. in H_2O than $\text{Cu}_2\text{S}_2\text{O}_3, \text{K}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Solution is not decomp. by boiling. Sol. in excess of $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg.)

Cuprous rubidium thiosulphate, $\text{Rb}_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Ppt. (J. Meyer, B. 1907, 40. 1357.)

$2\text{Rb}_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Ppt. (J. Meyer.)

$3\text{Rb}_2\text{S}_2\text{O}_3, \text{Cu}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Ppt. (J. Meyer.)

Cuprous silver sodium thiosulphate ammonia, $\text{Cu}_2\text{S}_2\text{O}_3, 2\text{Ag}_2\text{S}_2\text{O}_3, 5\text{Na}_2\text{S}_2\text{O}_3, 6\text{NH}_3$.

Ppt. Becomes dark when exposed to light. Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Shinn, J. Am. Chem. Soc. 1904, 26. 949.)

Cuprous sodium thiosulphate, $2\text{Cu}_2\text{S}_2\text{O}_3, 7\text{Na}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Ppt. from aqueous solution by alcohol. (Jochum, C. C. 1885. 642.)

$+12\text{H}_2\text{O}$. Sol. in very dil. $\text{HCl} + \text{Aq}$. (Jochum.)

$\text{Cu}_2\text{S}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Sol. in H_2O ; insol. in alcohol. (Rammelsberg.)

$+6\text{H}_2\text{O}$. (Jochum.)

$3\text{Cu}_2\text{S}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3 + 8\text{H}_2\text{O}$. Decomp. by H_2C . (Vortmann.)

$+5\text{H}_2\text{O}$. (Lenz, A. 40. 99.) Formula according to Jochum is—

$5\text{Cu}_2\text{S}_2\text{O}_3, 4\text{Na}_2\text{S}_2\text{O}_3 + 8\text{H}_2\text{O}$. Insol. in H_2O or alcohol. Sol. in $\text{HCl} + \text{Aq}$ without evolution of SO_2 , also in dil. H_2SO_4 or $\text{HNO}_3 + \text{Aq}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Jochum.)

$+6\text{H}_2\text{O}$. As above. (Jochum.)

$\text{Cu}_2\text{S}_2\text{O}_3, \text{Na}_2\text{S}_2\text{O}_3 + \text{H}_2\text{O}$. Insol. in H_2O ; sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Russell, Ch. 7tg. 9. 233.)

$+2\text{H}_2\text{O}$. Insol. in H_2O and alcohol.

Decomp. by dil. acids. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 84.)

$+2\frac{1}{2}\text{H}_2\text{O}$. Sol. in H_2O . Pptd. by alcohol. (Bhaduri, Z. anorg. 1898, 17. 1.)

$+3\text{H}_2\text{O}$. Decomp. by H_2O . (Vortmann, M. 1888, 9. 168.)

$3\text{Cu}_2\text{S}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3$. (Gerard, C. C. 1904, I. 253.)

$+5\text{H}_2\text{O}$. Sol. in 352 pts. H_2O . (Russell, Ch. Z. 1885, 9. 223.)

100 cc. 5% $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ dissolve 12.28 g.

" " 7.5% " " 17.46 g.

" " 10% " " 22.54 g.

(Russell, Ch. Z. 1885, 9. 223.)

$4\text{Cu}_2\text{S}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 6\text{H}_2\text{O}$. Insol. in H_2O and alcohol.

Decomp. by dil. acids. (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 84.)

$+9\text{H}_2\text{O}$. Sol. in H_2O . Pptd. by alcohol. (Bhaduri, Z. anorg. 1898, 17. 1.)

$7\text{Cu}_2\text{S}_2\text{O}_3, 5\text{Na}_2\text{S}_2\text{O}_3 + 16\text{H}_2\text{O}$. Decomp. by H_2O and by alcohol. (Bhaduri, Z. anorg. 1898, 17. 1.)

It is impossible to determine whether any of these compds. are complex or double salts. As a class they are not easily sol. and decomp. in solution. They may therefore be mixtures whose comp. depended upon the temp. and conc. of the solution in which they were pptd. (Rosenheim, Z. anorg. 1900, 25. 81.)

Cuprocupric sodium thiosulphate ammonia,
 $\text{Cu}_2\text{S}_2\text{O}_3, \text{CuS}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3, 4\text{NH}_3$.

Insol. in, but decomp. by hot H_2O . Sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ or $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Schütte, C. R. 42. 1267.)

Cuprous sodium thiosulphate cuprous chloride, $\text{Cu}_2\text{S}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3, 2\text{CuCl}$.

(Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 86.)

Cupric sodium thiosulphate cupric sulphide,
 $\text{Cu}_2\text{S}_2\text{O}_3, \text{Na}_2\text{S}_2\text{O}_3, \text{CuS} + 4\text{H}_2\text{O}$.

Sl. sol. in H_2O ; easily sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$, and $\text{NH}_4\text{OH} + \text{Aq}$; insol. in alcohol. (Lenz, A. 40. 99.)

$\text{Cu}_2\text{S}_2\text{O}_3, \text{Na}_2\text{S}_2\text{O}_3, 2\text{CuS}$. Sol. in H_2O or dil. $\text{HCl} + \text{Aq}$. (Kessel, B. 11. 1585.)

Cuprous sodium thiosulphate sodium chloride, $3\text{Cu}_2\text{S}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3, 4\text{NaCl} + 8\text{H}_2\text{O}$.

Sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Siewert, Zeit. ges. Naturwiss. **26**. 486.)

Ppt. (Bhaduri, Z. anorg. 1898, **17**. 3.)

Cupric thiosulphate ammonia, $\text{CuS}_2\text{O}_3, 4\text{NH}_3$.

100 pts. H_2O dissolve 21.79 pts. at 25° . (Pudschies, Dissert.)

Cuprocupric thiosulphate ammonium chloride, $\text{Cu}_2\text{O}, \text{CuO}, 3\text{S}_2\text{O}_3, 2\text{NH}_4\text{Cl}$.

Sol. in $\text{HNO}_3 + \text{Aq}$ with separation of S. (v. Hauer, W. A. B. **13**. 447.)

Glucinum thiosulphate, $\text{GlS}_2\text{O}_3 + 11\text{H}_2\text{O}$.

(Faktor, C. C. **1901**, II. 878.)

Gold (aurous) hydrogen thiosulphate, $\text{Au}_2\text{S}_2\text{O}_3, 3\text{H}_2\text{S}_2\text{O}_3$.

Known only in solution. (Fordos and Gélis, A. ch. (3) **13**. 394.)

Gold (aurous) sodium thiosulphate, $\text{Au}_2\text{S}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O ; solution decomp. on heating. Insol. in absolute, sl. sol. in dil. alcohol. (Fordos and Gélis.)

$\text{Au}_2\text{S}_2\text{O}_3, 6\text{Na}_2\text{S}_2\text{O}_3 + 10\text{H}_2\text{O}$. Very sol. in H_2O , but decomp. by boiling. (Jochum, C. C. **1885**. 642.)

Iron (ferrous) thiosulphate, $\text{FeS}_2\text{O}_3 + 5\text{H}_2\text{O}$.

Deliquescent. Very sol. in H_2O or alcohol. (Koene, Pogg. **63**. 241.)

Iron (ferrous) sodium thiosulphate, $\text{FeS}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 8\text{H}_2\text{O}$.

Very sol. in H_2O , and easily decomp. (Vortmann and Padberg, B. **22**. 2641.)

Lead thiosulphate, PbS_2O_3 .

Sol. in 3266 pts. H_2O . Sol. in alkali thiosulphates + Aq. (Rammelsberg, Pogg. **56**. 308.)

Lead lithium thiosulphate, $\text{PbS}_2\text{O}_3, \text{Li}_2\text{S}_2\text{O}_3$.

Very hygroscopic.

Decomp. by H_2O with separation of PbS . (J. Meyer, B. 1907, **40**. 1355.)

Lead potassium thiosulphate, $\text{PbS}_2\text{O}_3, 3\text{K}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Sol. in H_2O with partial separation of PbS_2O_3 . Sol. in $\text{K}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Rammelsberg, Pogg. **56**. 310.)

Lead rubidium thiosulphate, $2\text{Rb}_2\text{S}_2\text{O}_3, \text{PbS}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Difficultly sol. in cold H_2O .

Decomp. by H_2O . (J. Meyer, B. 1907, **40**. 1358.)

Lead sodium thiosulphate, $\text{PbS}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3$.

Sl. sol. in H_2O . Very easily sol. in $\text{Na}_2\text{C}_2\text{H}_3\text{O}_2$ and $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Lenz, A. **40**. 98.)

Insol. in alcohol.

$2\text{PbS}_2\text{O}_3, 5\text{Na}_2\text{S}_2\text{O}_3 + 60\text{H}_2\text{O}$. Easily decomp. (Jochum, C. C. **1885**. 642.)

$\text{PbS}_2\text{O}_3, 3\text{Na}_2\text{S}_2\text{O}_3 + 12\text{H}_2\text{O}$. Decomp. in boiling aqueous solution. (Vortmann and Padberg, B. **22**. 2637.)

Lead strontium thiosulphate.

Sol. in H_2O . Precipitated as a syrup by alcohol. (Rammelsberg.)

Lithium thiosulphate, $\text{Li}_2\text{S}_2\text{O}_3 + 3\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O and absolute alcohol. (Fock and Klüss, B. **22**. 3099.)

Lithium silver thiosulphate, $\text{Li}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + \text{H}_2\text{O}$.

Hygroscopic.

Decomp. by boiling H_2O and by acids. (Meyer and Eggeling, B. 1907, **40**. 1355.)

Magnesium thiosulphate, $\text{MgS}_2\text{O}_3 + 6\text{H}_2\text{O}$.

Very easily sol. in H_2O . Precipitated from conc. solution by alcohol. (Rammelsberg, Pogg. **56**. 303.)

Magnesium potassium thiosulphate, $\text{MgK}_2(\text{S}_2\text{O}_3)_2 + 6\text{H}_2\text{O}$.

Deliquescent, and sol. in H_2O . Less sol. than $\text{K}_2\text{S}_2\text{O}_3$. (Rammelsberg, Pogg. **56**. 304.)

Not deliquescent. (Fock and Klüss, B. **23**. 539.)

Magnesium rubidium thiosulphate, $\text{MgS}_2\text{O}_3, \text{Rb}_2\text{S}_2\text{O}_3 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . (Meyer, B. 1907, **40**. 1358.)

Manganous thiosulphate, MnS_2O_3 .

Sol. in H_2O , from which it is pptd. by alcohol. (Rammelsberg, Pogg. **56**. 305.)

+ $5\text{H}_2\text{O}$. Decomp. very easily. (Vortmann and Padberg, B. **22**. 2641.)

Manganous sodium thiosulphate, $\text{MnS}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3 + 16\text{H}_2\text{O}$.

Sol. in H_2O . Insol. or but sl. sol. in alcohol (Jochum, C. C. **1885**. 642.)

Mercuric potassium thiosulphate, $3\text{HgS}_2\text{O}_3, 5\text{K}_2\text{S}_2\text{O}_3$.

Sol. in 10 pts. H_2O at 15° , and $\frac{1}{2}$ pt. at 100° . Aqueous solution decomp. on standing or heating.

Insol. in alcohol. (Kirchhoff, Scher. J. **2**. 30.)

$\text{HgS}_2\text{O}_3, 3\text{K}_2\text{S}_2\text{O}_3 + 3\text{H}_2\text{O}$. (Fock and Klüss, B. **24**. 1353.)

$\text{HgS}_2\text{O}_3, 5\text{K}_2\text{S}_2\text{O}_3 + \text{H}_2\text{O}$. (F. and K.)

Nickel thiosulphate, $\text{NiS}_2\text{O}_3 + 6\text{H}_2\text{O}$.

Permanent. Sol. in H_2O . (Rammelsberg, Pogg. 56. 306.)

Nickel sodium thiosulphate, $2\text{NiS}_2\text{O}_3, 5\text{Na}_2\text{S}_2\text{O}_3 + 25\text{H}_2\text{O}$.

Efflorescent. Sol. in H_2O . (Jochum.)

Nickel thiosulphate ammonia, $\text{NiS}_2\text{O}_3, 4\text{NH}_3 + 6\text{H}_2\text{O}$.

Decomp. on air. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg, Pogg. 56. 306.)

$\text{NiS}_2\text{O}_3, 5\text{NH}_3 + \text{H}_2\text{O}$. (Ephraim, B. 1913, 46. 3108.)

$\text{NiS}_2\text{O}_3, 6\text{NH}_3 + 3\text{H}_2\text{O}$. (Vortmann and Padberg, B. 22. 2641.)

Platinous sodium thiosulphate.

See Platothiosulphate, sodium.

Potassium thiosulphate, $\text{K}_2\text{S}_2\text{O}_3 + 1/3, 1, 1 1/2, \text{H}_2\text{O}$, and $2\text{H}_2\text{O}$.

Very deliquescent. Very sol. in H_2O with absorption of heat. Solution is stable on the air. Insol. in alcohol.

100 pts. H_2O dissolve pts. $\text{K}_2\text{S}_2\text{O}_3$ at t° .

t°	Pts. $\text{K}_2\text{S}_2\text{O}_3$	Solid phase
0	96.1	$\text{K}_2\text{S}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$
17	150.5	$3\text{K}_2\text{S}_2\text{O}_3 \cdot 5\text{H}_2\text{O}$
20	155.4	"
25	165.0	"
30	175.7	"
35	202.4	$3\text{K}_2\text{S}_2\text{O}_3 \cdot 5\text{H}_2\text{O} + \text{K}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}$
40	204.7	$\text{K}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}$
45	208.7	"
50	215.2	"
55	227.7	"
60	238.3	"
56.1	234.5	$\text{K}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O} + 3\text{K}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}$
65	245.8	$3\text{K}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}$
70	255.2	"
75	268.0	"
78.3	292.0	$3\text{K}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O} + \text{K}_2\text{S}_2\text{O}_3$
80	293.1	$\text{K}_2\text{S}_2\text{O}_3$
85	298.5	"
90	312.0	"

(Inohiko Io, Mem. Coll. Sc. Kyoto, 1911, 3. 212.)

Sol. in dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$ without decomp. (Mathieu-Plessy, C. R. 101. 59.)

Insol. in ethyl acetate. (Casaseca, C. R. 30. 821.)

Potassium silver thiosulphate, $2\text{K}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3$.

Sol. in H_2O . (Cohen.)

$\text{K}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3$. Sl. sol. in H_2O . (Herschel.)

$3\text{K}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Rather sl. sol. in H_2O . (Rosenheim and Steinhäuser, Z. anorg. 1900, 25. 76.)

Sl. sol. in H_2O . (Rosenheim.)

Potassium silver thiosulphate ammonia, $\text{KAgS}_2\text{O}_3, 2\text{NH}_3$.

Very sl. sol. in H_2O . Easily sol. in hot $\text{NH}_4\text{OH} + \text{Aq}$. (Schwicker, B. 22. 1735.)

$5\text{K}_2\text{S}_2\text{O}_3, 3\text{Ag}_2\text{S}_2\text{O}_3, \text{NH}_3$. Difficultly sol. in H_2O with decomp.

Sol. in hot $\text{NH}_4\text{OH} + \text{Aq}$ with partial decomp. (J. Meyer, B. 1907, 40. 1359.)

Potassium sodium thiosulphate.

(a) $\text{KNaS}_2\text{O}_3 + 2\text{H}_2\text{O}$. Very sol. in H_2O . 100 pts. H_2O dissolve 213.7 pts. salt at 15° . (Schwicker, B. 22. 1733.)

(b) $\text{NaKS}_2\text{O}_3 + 2\text{H}_2\text{O}$. 100 pts. H_2O dissolve 205.3 pts. salt at 15° . (Schwicker.)

Potassium strontium thiosulphate, $\text{K}_2\text{S}_2\text{O}_3, \text{SrS}_2\text{O}_3 + 5\text{H}_2\text{O}$.

Sol. in H_2O . (Fock and Klüss, B. 24. 3017.)

Potassium zinc thiosulphate, $\text{K}_2\text{S}_2\text{O}_3, \text{ZnS}_2\text{O}_3 + \text{H}_2\text{O}$.

Very sol. in H_2O . (Rosenheim and Davidsohn, Z. anorg. 1904, 41. 238.)

Potassium thiosulphate sodium chloride, $\text{K}_2\text{S}_2\text{O}_3, \text{NaCl}$.

Sol. in H_2O . (Pape, Pogg. 139. 238.)

Rubidium thiosulphate, $\text{Rb}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Very hygroscopic; easily sol. in H_2O . (J. Meyer, B. 1907, 40. 1356.)

Rubidium silver thiosulphate, $2\text{Rb}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$.

Ppt. Not hygroscopic. Difficultly sol. in cold H_2O . Decomp. by warm H_2O . (J. Meyer.)

Rubidium silver thiosulphate ammonia, $\text{Rb}_2\text{S}_2\text{O}_3, \text{Ag}_2\text{S}_2\text{O}_3, \text{NH}_3$.

Ppt. (J. Meyer.)

$3\text{Rb}_2\text{S}_2\text{O}_3, 4\text{Ag}_2\text{S}_2\text{O}_3, \text{NH}_3$. Ppt. Decomp. in the air. (J. Meyer.)

Samarium thiosulphate.

(Cleve.)

Scandium thiosulphate, basic, $\text{Sc}(\text{OH})\text{S}_2\text{O}_3$.

Ppt. (R. J. Meyer, Z. anorg. 1914, 86. 282.)

Silver thiosulphate, $\text{Ag}_2\text{S}_2\text{O}_3$.

Sl. sol. in H_2O . Sol. in NH_4OH or alkali thiosulphates + Aq . (Herschel, Edinb. Phil. J. 1. 26.)

Silver sodium thiosulphate, $\text{Ag}_2\text{S}_2\text{O}_3, \text{Na}_2\text{S}_2\text{O}_3 + \text{H}_2\text{O}$.

Sl. sol. in H_2O . Easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$, also in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$ to form—

$\text{Ag}_2\text{S}_2\text{O}_3, 2\text{Na}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$. Easily sol. in H_2O or $\text{NH}_4\text{OH} + \text{Aq}$; somewhat sol. in al-

cohol, especially if warm or dilute. (Lenz, A. 40. 94.)

$\text{Ag}_2\text{S}_2\text{O}_3, 6\text{Na}_2\text{S}_2\text{O}_3 + 21\text{H}_2\text{O}$. Sol. in H_2O . (Jochum, C. C. 1885. 642.)

Silver sodium thiosulphate acetylde acetylene, $2\text{Na}_2\text{S}_2\text{O}_3, 7\text{Ag}_2\text{S}_2\text{O}_3, 18\text{Ag}_2\text{C}_2, 32\text{C}_2\text{H}_2$.

Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in alkalies. (Bhaduri, Z. anorg. 1913, 79. 356.)

$7\text{Na}_2\text{S}_2\text{O}_3, 5\text{Ag}_2\text{S}_2\text{O}_3, 86\text{Ag}_2\text{C}_2, 13\text{C}_2\text{H}_2$. (Bhaduri.)

Silver sodium thiosulphate ammonia, $\text{NaAgS}_2\text{O}_3, \text{NH}_3$.

Very unstable. (Schwicker, B. 22. 1736.)

Silver strontium thiosulphate, $\text{Ag}_2\text{S}_2\text{O}_3, \text{SrS}_2\text{O}_3$.

Nearly insol. in H_2O . Very sl. sol. in $\text{SrS}_2\text{O}_3 + \text{Aq}$; easily sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Herschel.)

Sodium thiosulphate, $\text{Na}_2\text{S}_2\text{O}_3$ and +1, 2 and $5\text{H}_2\text{O}$.

100 pts. H_2O dissolve:
At 16° , 65 pts. $\text{Na}_2\text{S}_2\text{O}_3$.
" 20° , 69 " "
" 25° , 75 " "
" 30° , 82 " "
" 35° , 89 " "
" 40° , 98 " "
" 45° , 109 " "
" 47° , 114 " "
(Mulder.)

100 pts. H_2O dissolve at 0° , 47.6 pts. $\text{Na}_2\text{S}_2\text{O}_3$; at 20° , 69.5 pts. ; at 40° , 104 pts. ; at 60° , 192.3 pts. (Kremers, Pogg. 99. 50)

100 pts. H_2O dissolve 171 pts. cryst. (=108.9 pts. anhydrous)salt at 19.5° to form a solution of 1.3875 sp. gr. (Schiff, A. 113. 350.)

By supersaturation 100 pts. H_2O may dissolve 217.4 pts. $\text{Na}_2\text{S}_2\text{O}_3$ at 0° . (Kremers.)

Solubility in H_2O .

t°	% $\text{Na}_2\text{S}_2\text{O}_3$	Solid phase
0	34.43	$\text{Na}_2\text{S}_2\text{O}_3 + 5\text{H}_2\text{O}$
10	37.89	"
20	41.17	"
30	45.86	"
40	50.65	"
45	54.49	"
50	62.92	$\text{Na}_2\text{S}_2\text{O}_3 + 2\text{H}_2\text{O}$
60	67.39	"
72	70.39	"
80.5	71.33	"
90.5	71.76	"
100	72.68	"

(Taylor, Proc. Edinburgh Soc. 1898, 22. 49.)

Solubility of anhydrous $\text{Na}_2\text{S}_2\text{O}_3$ in H_2O at t° .

t°	g. $\text{Na}_2\text{S}_2\text{O}_3$ per 100 g.	
	Solution	H_2O
40	67.40	206.70
45	67.60	208.60
50	67.76	210.20
55	68.15	214.00
60	68.48	217.30
65	68.80	220.50
70	69.05	223.10
75	69.35	226.30
80	69.80	231.80

(Young and Burke, J. Am. Chem. Soc. 1906, 28. 327.)

The solubility of the hydrates of $\text{Na}_2\text{S}_2\text{O}_3$ are exceedingly complicated. There are five groups of hydrates, (I) primary, (II) secondary, (III) tertiary, (IV) quaternary, and (V), quintary, and transitions occur between members of the same and different groups. $\text{Na}_2\text{S}_2\text{O}_3 + 5\text{H}_2\text{O}$ (I) is the commercial thiosulphate. The relations and solubility of the various hydrates is shown in the following table.

Solubility of $\text{Na}_2\text{S}_2\text{O}_3$ in H_2O at t° .

I. Primary hydrates.

t°	g. $\text{Na}_2\text{S}_2\text{O}_3$ per 100 g.		Solid phase
	Solution	H_2O	
0	33.40	50.15	$\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$ (I)
5	35.33	54.64	"
10	37.37	59.69	"
15	39.11	64.22	"
20	41.20	70.07	"
25	43.15	75.90	"
30	45.19	82.45	"
35	47.71	91.24	"
40	50.83	105.37	"
45	55.33	123.87	"
48.17	" + $\text{Na}_2\text{S}_2\text{O}_3, 2\text{H}_2\text{O}$ (I)
0	52.73	111.60	$\text{Na}_2\text{S}_2\text{O}_3, 2\text{H}_2\text{O}$ (I)
5	53.45	114.90	"
10	53.94	117.10	"
20	55.15	122.68	"
25	56.03	127.43	"
30	57.13	133.27	"
40	59.38	146.20	"
45	60.73	154.70	"
50	62.28	165.11	"
55	63.85	176.60	"
60	65.68	191.30	"
65	68.04	212.90	"
66.5	" + $\text{Na}_2\text{S}_2\text{O}_3$

Solubility of $\text{Na}_2\text{S}_2\text{O}_3$ in H_2O at t° .—
Continued.

II. Secondary hydrates.

t°	g. $\text{Na}_2\text{S}_2\text{O}_3$ per 100 g.		Solid phase
	Solu- tion	H_2O	
	41.96	72.30	$\text{Na}_2\text{S}_2\text{O}_3 \cdot 5\text{H}_2\text{O}(\text{II})$
	43.56	77.17	"
	45.25	82.05	"
	47.27	89.36	"
	49.38	97.55	"
	52.15	108.98	"
	56.57	130.26	"
22	" $\text{Na}_2\text{S}_2\text{O}_3 \cdot 4\text{H}_2\text{O}(\text{II})$
5	58.59	141.48	$\text{Na}_2\text{S}_2\text{O}_3 \cdot 4\text{H}_2\text{O}(\text{II})$
2	60.51	153.23	"
6	62.80	168.52	"
65	" $+\text{Na}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}(\text{II})$

	60.47	153.00	$\text{Na}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}(\text{II})$
	60.74	154.70	"
	61.04	156.70	"
	61.57	161.20	"
	62.11	163.90	"
	62.73	168.30	"
	63.56	174.40	"
	64.32	180.20	"
	65.22	187.60	"
	66.02	194.30	"
	66.82	201.40	"
	67.90	211.50	"
5	" $+\text{Na}_2\text{S}_2\text{O}_3$

III. Tertiary hydrates.

	46.14	85.67	$\text{Na}_2\text{S}_2\text{O}_3 \cdot 6\text{H}_2\text{O}$ (III and IV)
	48.44	93.95	"
	51.66	106.80	"
	54.96	122.00	"
.35	" $+\text{Na}_2\text{S}_2\text{O}_3 \cdot \frac{1}{2}\text{H}_2\text{O}(\text{IV})$
.3	" $+\text{Na}_2\text{S}_2\text{O}_3 \cdot \frac{1}{2}\text{H}_2\text{O}(\text{III})$

	57.42	134.80	$\text{Na}_2\text{S}_2\text{O}_3 \cdot \frac{1}{2}\text{H}_2\text{O}(\text{III})$
	57.84	137.20	"
	58.28	139.70	"
	58.80	142.70	"
	59.28	145.60	"
	60.18	151.10	"
	60.78	155.00	"
	61.57	160.20	"
	62.60	167.40	"
	63.97	177.50	"
5	64.68	183.00	"
.5	" $+\text{Na}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}(\text{III})$

5	64.78	183.90	$\text{Na}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}(\text{III})$
	65.30	188.20	"
.5	65.89	193.20	"
	66.45	198.10	"
	68.07	213.10	"
	" $+\text{Na}_2\text{S}_2\text{O}_3$

Solubility of $\text{Na}_2\text{S}_2\text{O}_3$ in H_2O at t° .—
Continued.

IV. Quaternary hydrate.

t°	g. $\text{Na}_2\text{S}_2\text{O}_3$ per 100 g.		Solid phase
	Solu- tion	H_2O	
0	57.63	136.00	$\text{Na}_2\text{S}_2\text{O}_3 \cdot \frac{1}{2}\text{H}_2\text{O}(\text{IV})$
5	58.08	138.60	"
10	58.49	140.90	"
15	59.00	143.90	"
20	59.57	147.30	"
25	60.35	152.30	"
30	61.03	156.60	"
35	61.94	162.80	"
40	62.95	169.90	"
45	64.22	179.50	"
50	65.45	189.50	"
55	67.07	203.70	"
58	" $+\text{Na}_2\text{S}_2\text{O}_3$

V. Quintary hydrates.

0	57.63	136.00	$\text{Na}_2\text{S}_2\text{O}_3 \cdot 2\text{H}_2\text{O}(\text{V})$
5	58.23	139.40	"
10	59.05	144.20	"
15	60.02	150.10	"
20	61.02	156.50	"
25	62.30	165.30	"
30	63.56	174.40	"
35	65.27	188.00	"
27.5	" $+\text{Na}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}(\text{V})$

30	63.34	172.80	$\text{Na}_2\text{S}_2\text{O}_3 \cdot \text{H}_2\text{O}(\text{V})$
35	64.07	178.40	"
40	64.75	183.70	"
45	65.58	190.50	"
50	66.58	199.20	"
55	67.59	209.50	"
43	" $+\text{Na}_2\text{S}_2\text{O}_3 \cdot \frac{1}{2}\text{H}_2\text{O}(\text{V})$

25	64.21	179.40	$\text{Na}_2\text{S}_2\text{O}_3 \cdot \frac{1}{2}\text{H}_2\text{O}(\text{V})$
35	64.60	182.50	"
40	64.99	185.60	"
45	65.61	190.02	"
50	66.02	194.30	"
55	66.57	199.10	"
60	67.40	206.70	"
65	68.24	214.90	"
70	69.06	...	"
70	" $\text{Na}_2\text{S}_2\text{O}_3$

(Young and Burke, J. Am. Chem. Soc. 1906, 28, 321.)

Heat is absorbed by dissolving in H_2O .
110 pts. $\text{Na}_2\text{S}_2\text{O}_3 + 5\text{H}_2\text{O} + 100$ pts. H_2O
lower temp. from 10.7° to 8° . (Rüdorff, B. 2, 68.) $+\frac{1}{2}\text{H}_2\text{O}$. (Young and Burke, J. Am. Chem. Soc. 1906, 28, 321.) $+\text{H}_2\text{O}$. $+\frac{1}{2}\text{H}_2\text{O}$. (Young and Burke.) $+\frac{1}{2}\text{H}_2\text{O}$. (Young and Burke.)

+2H₂O.
+4H₂O. M.-pt. 41.60°. (Young and Burke.)
+5H₂O. M.-pt. 45° (Kopp); 48° (Kremers), 50° (Mulder); 48.5° (Tilden, Chem. Soc. 45. 409); 47.9° (Taylor, Proc. Edinburgh, Soc. 1898, 22. 249); 48.09° (Richards and Churchill, Z. phys. Ch. 1899, 28. 314); 48.45° (Young and Burke, J. Am. Chem. Soc. 1906, 28. 324.)

Labile modification melts at 32°. (Parmentier and Amat, C. R. 96. 735.)

Sp. gr. of Na₂S₂O₃+Aq at 19°.
% = %Na₂S₂O₃+5H₂O.

%	Sp. gr.	%	Sp. gr.	%	Sp. gr.
1	1.0052	18	1.0975	35	1.1986
2	1.0105	19	1.1031	36	1.2048
3	1.0158	20	1.1087	37	1.2110
4	1.0211	21	1.1145	38	1.2172
5	1.0264	22	1.1204	39	1.2234
6	1.0317	23	1.1263	40	1.2297
7	1.0370	24	1.1322	41	1.2362
8	1.0423	25	1.1381	42	1.2427
9	1.0476	26	1.1440	43	1.2492
10	1.0529	27	1.1499	44	1.2558
11	1.0584	28	1.1558	45	1.2624
12	1.0639	29	1.1617	46	1.2690
13	1.0695	30	1.1676	47	1.2756
14	1.0751	31	1.1738	48	1.2822
15	1.0807	32	1.1800	49	1.2888
16	1.0863	33	1.1862	50	1.2954
17	1.0919	34	1.1924

(Schiff, A. 113. 118.)

B.-pt. of Na₂S₂O₃+Aq. P=pts. Na₂S₂O₃ to 100 pts. H₂O.

B.-pt.	P	B.-pt.	P	B.-pt.	P
101°	14	110°	104	119°	201
102	27	111	113	120	214.5
103	39	112	122	121	229
104	49.5	113	131.5	122	244
105	59	114	141.5	123	262
106	68	115	152	124	283
107	77	116	164	125	311
108	86	117	175.75	126	348
109	95	118	188

(Gerlach, Z. anal. 26. 436.)

Moderately sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, 20. 829.)

100 pts. absolute ethyl alcohol dissolve at room temperature 2.5 mg. Na₂S₂O₃. (Bödtker, Z. phys. Ch. 1897, 22. 410.)

100 pts. absolute ethyl alcohol dissolve at room temperature 3.4 mg. Na₂S₂O₃+5H₂O. (Bödtker, Z. phys. Ch. 1897, 22. 510.)

Sol. in oil of turpentine (Edison, Am. Chemist. 7. 127). Insol. therein (Techn. J. B. 27. 1003).

Insol. in ethyl acetate. (Casaseca, C. R. 30. 821.)

Sodium thallos thiosulphate, 3Na₂S₂O₃, 2Tl₂S₂O₃+10H₂O.

Sol. in H₂O. (Werther.)

+8H₂O. (Jochum.)

2Na₂S₂O₃, Tl₂S₂O₃+8H₂O. (Vortmann and Padberg, B. 22. 2638.)

Sodium zinc thiosulphate, Na₂S₂O₃, 2ZnS₂O₃+23H₂O.

Sol. in H₂O. (Jochum, C. C. 1885. 642.)

3Na₂S₂O₃, 2ZnS₂O₃+10H₂O. Deliquescent. (Vortmann and Padberg, B. 22. 2640.)

Sodium thiosulphate mercuric iodide, 2Na₂S₂O₃, HgI₂.

Decomp. by H₂O and by alcohol. (Eder and Ulm, M. 1882, 3. 197.)

Strontium thiosulphate, SrS₂O₃+5H₂O.

Permanent. Sol. in 6 pts. cold H₂O (Gay-Lussac); in 4 pts. H₂O at 13°, and 1.75 pts. boiling H₂O (Herschel, 1819).

1 g. is sol. in 3.7 cc. H₂O at room temp. (Antenrieth, Z. anal. 1898, 37. 293.)

Gradually efflorescent. Insol. in alcohol (Herschel.)

Thallos thiosulphate.

Ppt. Sl. sol. in cold, easily sol. in hot H₂O. (Crookes.)

Easily sol. in Na₂S₂O₃+Aq. (Jochum.)

Tin thiosulphate (?).

Sol. in H₂O.

Uranyl thiosulphate, (UO₂)S₂O₃.

Ppt. (Faktor, C. C. 1901, II. 878.)

Zinc thiosulphate, ZnS₂O₃+xH₂O.

Very deliquescent, and very sol. in H₂O and alcohol. (Rammelsberg.)

Zinc thiosulphate ammonia, ZnS₂O₃, 2NH₃.

Decomp. by H₂O. Sol. in NH₄OH+Aq. from which it is pptd. by alcohol. (Rammelsberg, Pogg. 56. 62.)

Thio(trithiazyl bromide, N₃S₃Br.

Decomp. H₂O and alkalis.

Stable in the air. (Muthmann, B. 1897, 30. 630.)

Thiodithiazyl dichloride, S₂N₂Cl₂.

See Nitrogen sulphochloride.

Thiotrithiazyl chloride, S₃N₃Cl.

See Nitrogen sulphochloride.

Thiotrithiazyl iodide, N₃S₃I.

Decomp. spontaneously in the air. Very unstable. (Muthmann, B. 1897, 30. 631.)

Thiothiazyl nitrate, $S_4N_2NO_3$.

Sol. in H_2O with decomp. Sol. in HNO_3 + Aq. (Demargay, C. R. 91. 1006.)

Sol. in cold H_2O with decomp.

Completely insol. in organic solvents, as CS_2 , $CHCl_3$, acetone, and acetic ether. (Muthmann and Seitter, B. 1897, 30. 629.)

Thiothiazyl sulphate $(S_4N_2)HSO_4$.

Stable on air. Sol. in H_2O with decomp. (Demargay, C. R. 91. 854. 1066.)

Thiothiazyl sulphocyanide, N_4S_4CNS .

Stable in the air.

Sol. in benzene and chloroform (Muthmann and Seitter, B. 1897, 30. 631.)

Dithiothiazyl dichloride, $S_4N_2Cl_2$.

See Nitrogen sulphochloride.

Thorium, Th.

Not oxidised by boiling H_2O .

Quickly sol. (Chydenius, Pogg. 119. 43), very slowly sol. by long boiling - (Berzelius, Pogg. 16. 385) in HNO_3 + Aq. Insol. in cold, easily sol. in warm dil. H_2SO_4 + Aq. Slowly sol. in cold, rapidly in hot HCl + Aq. Easily oxidised by aqua regia. Insol. in KOH + Aq or HF + Aq.

Sl. sol. in dil. H_2SO_4 + Aq; decomp. by conc. H_2SO_4 . Very al. sol. in dil., and less in conc. HNO_3 + Aq. Easily sol. in conc. HCl + Aq, and aqua regia. (Nilson, B. 15. 2521.)

Thorium B.

Somewhat sol. in hot H_2O , which dissolves 70% in an hour. More rapidly sol. in dil. acids or hot KI + Aq. Somewhat sol. in organic solvents. Insol. in CS_2 and in methylene iodide. (Hogley, Phil. Mag. 1913, (6) 35. 331.)

Thorium C.

Somewhat sol. in hot H_2O . More rapidly sol. in hot KI + Aq. and in dil. acids. More sol. in organic solvents than thorium B. 20% sol. in CS_2 or methylene iodide in 10 minutes. (Hogley, Phil. Mag. 1913, (6) 35. 331.)

Thorium tetraboride, ThB_4 .

Sol. in cold conc. HNO_3 and HCl and in hot conc. H_2SO_4 . (Jassonneix, C. R. 1905, 141. 192.)

Thorium hexaboride, ThB_6 .

Sol. in hot dil. or conc. HNO_3 ; insol. in H_2SO_4 , HCl , HF and aq. alkalis. (Jassonneix, C. R. 1905, 141. 193.)

Thorium dibromide, $ThBr_2$.

Sol. in H_2O with partial decomp. (Troost and Ouvrard, A. ch. (6) 17. 227.)

Thorium tetrabromide, $ThBr_4$.

Sol. in H_2O . (Berzelius.)

Very hygroscopic, and sol. in H_2O with partial decomp. (Troost and Ouvrard, A. ch. (6) 17. 229.)

Sl. sol. in organic solvents. (Matthews, J. Am. Chem. Soc. 1898, 20. 840.)

+7 H_2O . Sol. in alcohol. (Rosenheim, B. 1900, 33. 979.)

+8 H_2O . Very sol. in alcohol and H_2O . Insol. in chloroform and light petroleum. (Lesinsky, Z. anorg. 1897, 15. 82.)

Thorium tetrabromide ammonia, $ThBr_4 \cdot 3NH_3$.

Insol. in organic solvents. (Matthews, J. Am. Chem. Soc. 1898, 20. 840.)

Thorium carbide, ThC_2 .

Almost insol. in conc. acids; decomp. by H_2O and by dil. acids. (Moissan, C. R. 1896, 123. 577.)

Thorium tetrachloride, $ThCl_4$.

Anhydrous. Extremely deliquescent, and sol. in H_2O with evolution of heat. Sol. in alcohol.

Sol. in ether. (Matthews, J. Am. Chem. Soc. 1898, 20. 824.)

+7 H_2O . Deliquesces in the air. Very sol. in H_2O and alcohol. Insol. in ether. (Krüss, Z. anorg. 1897, 14. 368.)

+8 H_2O . Hygroscopic. Easily sol. in H_2O and absolute alcohol. Not pptd. from solution in alcohol by ether.

+9 H_2O . As above. (Rosenheim, Z. anorg. 1903, 35. 426.)

Thorium tetrachloride ammonia, $ThCl_4 \cdot 6NH_3$.

Insol. in ether. Decomp. by H_2O .

$ThCl_4 \cdot 8NH_3$. Fumes in moist air. Decomp. by H_2O . (Matthews, J. Am. Chem. Soc. 1898, 20. 824.)

The compds. of $ThCl_4$ with NH_3 are of the type $ThCl_4 \cdot nNH_3$ and may be classified into three groups.

(1) $n=4, 6, 7, 12, 18$. Compds. are decomp. by H_2O .

(2) $n=4, 6, 7$. Compds. are not decomp. by H_2O .

(3) $n=6, 7, 12, 18$. The first two compds. of this series are identical with the last two of group (2).

$ThCl_4 \cdot 4NH_3$ of group (2) is the only one in the whole series stable above 120° . (Chauvenet, C. R. 1910, 151. 387.)

Thorium fluoride, $ThF_4 \cdot 4H_2O$.

Insol. in H_2O or HF + Aq.

Thorium hydride, ThH_2 .

Decomp. by dil. HCl + Aq. (Winkler, B. 34. 873.)

Thorium hydroxide, $\text{Th}(\text{OH})_4$.

Insol. in H_2O .

Sol. in acids, except oxalic, molybdic, and hydrofluoric acids.

Insol. in alkali hydroxides, but easily sol. in alkali carbonates + Aq. More sol. in $\text{NH}_4\text{OH} + (\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ than in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$ alone. (Berzelius.) Not pptd. in presence of tartaric and citric acids. (Chydenius, Pogg. 119. 43.)

$4\text{ThO}_2, \text{H}_2\text{O}$. Insol. in water and acids at boiling temp.

Thorium hydroxybromide, $\text{Th}(\text{OH})_2\text{Br}_2 + 11\text{H}_2\text{O}$.

Sol. in alcohol. (Rosenheim, B. 1900, 33. 979.)

Thorium hydroxychloride, $(\text{OH})\text{ThCl}_3 + 11\text{H}_2\text{O}$.

Sol. in alcohol. (Rosenheim, B. 1900, 33. 978.)

$\text{Th}(\text{OH})_2\text{Cl}_2 + 5\text{H}_2\text{O}$. Slowly takes up H_2O from the air.

Sol. in H_2O without decomp.

Sol. in alcohol. Pptd. from solution in alcohol by ether. (Rosenheim, Z. anorg. 1903, 35. 425.)

+ $8\text{H}_2\text{O}$. Hydrosopic; sol. in alcohol. (Rosenheim, B. 1900, 33. 978.)

Thorium hydroxyiodide, $\text{Th}(\text{OH})\text{I}_3 + 10\text{H}_2\text{O}$.

Evolves iodine in the light. (Rosenheim, Z. anorg. 1903, 35. 430.)

Thorium iodide.

Sol. in H_2O .

Thorium nitride, Th_3N_4 .

Decomp. by H_2O . (Matignon, C. R. 1901, 132. 37.)

Thorium oxide, ThO_2 .

When ignited is insol. in HCl , and $\text{HNO}_3 + \text{Aq}$. Sol. in H_2SO_4 by heating to boiling and subsequent addition of H_2O . Insol. in alkali hydrates or carbonates + Aq.

Thorium metoxide, $\text{Th}_2\text{O}_3 + \text{H}_2\text{O}$.

Sl. sol. in HCl . (Locke, Z. anorg. 1894, 7. 348.)

+ $2\text{H}_2\text{O}$. Loses 1 H_2O at 100° . Sol. in H_2O ; insol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Locke.)

Metathorium oxide.

Sol. in H_2O after having been treated with conc. HNO_3 or $\text{HCl} + \text{Aq}$, even if previously ignited.

$\text{ThO}_2, x\text{Th}(\text{OH})_4$. Compare Th_2O_3 . (Locke) (Stevens, Z. anorg. 1901, 27. 42.)

Thorium peroxide, Th_2O_7 .

Precipitate. (Cleve, C. R. 100. 605.)

Thorium oxychloride.

Decomp. by H_2O into ThCl_4 and ThO_2 . ThOCl_2 . Sol. in H_2O .

Insol. in abs. alcohol. (Matignon, A. ch. 1907, (8) 10. 133.)

+ $3\text{H}_2\text{O}$.

+ $5\text{H}_2\text{O}$. (Matignon, A. ch. 1907, (8) 10. 135.)

Metathorium oxychloride, $\text{ThO}_2, x\text{ThCl}_4$.

Hydrosopic; sol. in H_2O ; insol. in abs. alcohol. (Stevens, Z. anorg. 1901, 27. 47.)

Thorium oxyfluoride, ThOF_2 .

Insol. in H_2O .

Sol. in H_2SO_4 with decomp. (Chauvenet, C. R. 1908, 146. 974.)

Thorium oxysulphide, $\text{ThS}_2, 2\text{ThO}_2$.

(Chydenius.)

Thorium phosphide.

Insol. in H_2O . (Berzelius.)

Thorium silicide, ThSi_2 .

Sol. in aq. min. acids; insol. in aq. alkalis. Decomp. by fusion with NaOH or KOH . (Hönigschmid, C. R. 1906, 142. 158.)

Thorium sulphide, ThS_2 .

Insol. in warm H_2SO_4 . Very slightly attacked by HNO_3 or $\text{HCl} + \text{Aq}$. Sol. in hot aqua regia. (Berzelius.)

Thoromolybdic acid.**Ammonium thoromolybdate,**

$(\text{NH}_4)_2\text{Th}(\text{Mo}_2\text{O}_7)_6 + 8\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in dil. acids. (Barbieri, C. A. 1913. 372.)

$(\text{NH}_4)_4\text{H}_2\text{Th}(\text{Mo}_2\text{O}_7)_6 + 11\text{H}_2\text{O}$. Insol. in H_2O ; sol. in dil. acids. (Barbieri.)

Silver thoromolybdate, $\text{Ag}_2\text{Th}(\text{Mo}_2\text{O}_7)_4$.

Insol. in H_2O ; sl. sol. in $\text{HNO}_3 + \text{Aq}$, but nearly insol. in presence of AgNO_3 . (Barbieri.)

Sodium thoromolybdate, $\text{Na}_2\text{Th}(\text{Mo}_2\text{O}_7)_4 + 15\text{H}_2\text{O}$.

Insol. in H_2O ; sol. in dil. acids. (Barbieri.)

$\text{Na}_4\text{H}_2\text{Th}(\text{Mo}_2\text{O}_7)_6 + 17\text{H}_2\text{O}$. Insol. in H_2O ; sol. in dil. acids. (Barbieri.)

Thulium, Tm .**Thulium chloride, $\text{Tm}_2\text{Cl}_6 + 14\text{H}_2\text{O}$.**

Very sol. in H_2O and in alcohol. (James, J. Am. Chem. Soc. 1911, 33. 1342.)

Thulium hydroxide.

Easily sol. in dil. acids. (James, J. Am. Chem. Soc. 1911, 33. 1342.)

Thallium oxide, Tm_2O_3 .

Slowly sol. in hot conc. acids. (James, J. Am. Chem. Soc. 1911, **33**, 1342.)

Tin, Sn.

Insol. in H_2O . Tin is not attacked by distilled H_2O when air is passed through it for a week.

Slowly sol. in dil. cold $HCl + Aq$, but rapidly sol. if hot and conc. Slowly sol. in hot dil. $H_2SO_4 + Aq$, but decomp. by hot conc. H_2SO_4 .

Readily sol. in cold aqua regia. Attacked violently by conc. $HNO_3 + Aq$ with pptn. of SnO_2 . Completely sol. in dil. cold $HNO_3 + Aq$ (1 pt. HNO_3 : 1 pt. H_2O) at 22° . (Hay, C. N. **22**, 298.) Not attacked by pure conc. $HNO_3 + Aq$ of 1.512-1.419 sp. gr., but violently attacked by less conc. acid. Also attacked by most conc. acid if it contains NO_2 . (Millon, A. ch. (3) **6**, 95.)

If Sn is placed in dil. $HNO_3 + Aq$ of 1.15 sp. gr. it is sl. dissolved, but soon pptd. again as SnO_2 . If a small amt. of NH_4Cl is added, the Sn remains permanently in solution; $HCl + Aq$ has a similar action. (Ordway, Am. J. Sci. (2) **23**, 220.) Easily sol. in the cold in mixture of 1 vol H_2SO_4 , 2 vols. HNO_3 , and 3 vols. H_2O . (Basset, C. N. **53**, 172.)

$HNO_3 + Aq$ containing less than 12% HNO_3 attacks Sn and forms a stannous salt, which decomposes, giving a turbid solution. $HNO_3 + Aq$ (12-45% HNO_3) completely dissolves Sn, but solution becomes turbid on standing. $HNO_3 + Aq$ containing more than 45% HNO_3 does not dissolve Sn, but forms a white substance, which is sol. in H_2O if over 70% acid is used; this solution soon becomes turbid. (Montemartini, Gazz. ch. it. **22**, 384.)

Sn dissolves in $HNO_3 + Aq$ at low temps. ($0-21^\circ$). When very dil. $HNO_3 + Aq$ (14% HNO_3) is used, the amount of stannous salt formed decreases only slightly with increase of temp. while with 30-40% acid it falls to zero at 21° . (Walker, J. Soc. Chem. Ind. **1893**, 845.)

In presence of Fe, Cr or Al, $HNO_3 + Aq$ acts on Sn to form soluble products, from which conc. HNO_3 ppt. all Sn as meta-stannic acid. (van Leent, C. C. **1899**, I, 101.)

Much more sol. in acids when small quantities of metallic salts have been added. This is most noticeable when $PtCl_4$ or tartar emetic is added to $HCl + Aq$. $HCl + Aq$ with tartar emetic exerts 11 times, and with $PtCl_4$ 13 times the action exhibited by pure acid. (Millon, C. R. **21**, 47.)

Sol. in 2N $HClO_4 + Aq$. (Hendrixson, J. Am. Chem. Soc. 1904, **26**, 755.)

Pyrosulphuric acid dissolves Sn on warming. (Divers, Chem. Soc. 1885, **47**, 639.)

Hot telluric acid attacks Sn. (Hutchins, J. Am. Chem. Soc. 1905, **27**, 1183.)

Sn is attacked by 17% $HN_3 + Aq$. (Curtius and Rissom, J. pr. 1898, (2) **58**, 299.)

Sol. in a solution of Na in liquid NH_3 .

(Kraus, J. Am. Chem. Soc. 1907, **29**, 1582.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**, 830.)

Sol. in boiling alum + Aq (1 pt. alum to 4 pta. H_2O).

Sol. in $KHSO_4$, NH_4Cl (1:4), and $K_2C_2H_3O_4 + Aq$. Sl. sol. in $KC_2H_3O_4 + Aq$, but not attacked by $MgSO_4$, K_2SO_4 , KNO_3 , or $Na_2SO_4 + Aq$. (Cludius, J. pr. **9**, 161.)

Sol. in alkalis + Aq.

Attacked easily by conc. $NaCl$, KCl , or $NH_4NO_3 + Aq$, not attacked by $NH_4Cl + Aq$. (Hallock, Am. Ch. J. **6**, 52.)

Sol. in $Fe(NO_3)_3 + Aq$ in presence of $HNO_3 + Aq$ in proportion of 1 atom Sn to 1 atom Fe. (Lepes and Storch, W. A. B. **98**, 2b, 268.)

Solubility in dil. saline solutions.

100 ccm. H_2O containing 0.5 g. $NaCl$ or KCl dissolve 6 mg. Sn from 11.8 sq. cm in one week when air without CO_2 is passed through the solution, but none at all when the air contains CO_2 .

100 ccm. H_2O containing 1 g. NH_4Cl dissolve 5 mg. Sn under above conditions without CO_2 , and none with CO_2 .

With 1 g. $MgCl_2$, 1 mg. Sn was dissolved without CO_2 , and none with CO_2 .

With 1 g. K_2SO_4 , 2 mg. Sn were dissolved without CO_2 , and none with CO_2 .

With 1 g. KNO_3 , 3 mg. Sn were dissolved without CO_2 , and 1 mg. with CO_2 .

With 1 g. Na_2CO_3 , 7 mg. Sn were dissolved without CO_2 .

With 1 g. $NaOH$, 220 mg. Sn were dissolved without CO_2 .

$CaO, H_2 + Aq$ did not dissolve. (Wagner, Dingl. **231**, 260.)

Not attacked by sugar + Aq. (Klein, C. R. **102**, 1170.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0134 g. Sn in 6 days. (Gates, J. phys. Chem. 1911, **15**, 143.)

Tin antimonide, $SnSb$.

(Stead, J. Soc. Chem. Ind. 1897, **16**, 205.)

Tin arsenide, Sn_3As_4 .

(Stead, J. Soc. Chem. Ind. 1897, **16**, 206.)

Tin (stannous) bromide, $SnBr_2$.

Sol. in H_2O .

Sol. in pyridine. (Naumann, B. 1904, **37**, 1609.)

Mol. weight determined in pyridine and ethyl sulphide (Werner, Z. anorg. 1897, **15**, 23.)

Tin (stannic) bromide, basic, $SnBr_2 \cdot OH + 3H_2O$.

Sol. in H_2O . Decomp. in aq. solution when warmed.

Sol. in ether, methyl alcohol, ethyl alcohol, acetone, acetic acid and esters of organic acids. Nearly insol. in benzene, ligroin and $CHCl_3$. (Pfeiffer, Z. anorg. 1914, **37**, 242.)

Tin (stannic) bromide, SnBr_4 .

Deliquescent. Sol. in H_2O without evolution of heat. (Balard.)

Decomp. by H_2O much more quickly than SnCl_4 . (Lorenz, Z. anorg. 1895, **9**, 378.)

Easily sol. in AsBr_3 . (Walden, Z. anorg. 1902, **29**, 374.); PCl_3 , PBr_3 , and S_2Cl_2 . (Walden, Z. anorg. 1900, **35**, 217.)

Sol. in acetone. (Naumann, B. 1904, **37**, 4328.)

+ $4\text{H}_2\text{O}$. (Preis and Raymann, C. C. 1882, 773.)

Stannic hydrogen bromide, $\text{SnBr}_4 \cdot 2\text{HBr}$.

See Bromostannic acid.

Stannic bromide with MBr .

See Bromostannate, M .

Tin (stannic) bromochloride, SnClBr_3 .

Fumes in moist air; decomp. by H_2O . (Besson, C. R. 1897, **124**, 685.)

SnCl_2Br_2 . Fumes in moist air. Decomp. by H_2O . (Besson.)

SnCl_3Br . Fumes in moist air. Decomp. by H_2O . (Besson.)

Tin (stannic) bromiodide, SnBr_3I .

Sol. in cold H_2O . Decomp. in aq. solution at 80° .

SnBr_2I .

SnBr_2I_2 . (Lenormand, C. C. 1899, II, 521; J. Pharm. 1899, **10**, 114.)

Tin (stannous) chloride, SnCl_2 , and + $2\text{H}_2\text{O}$.

Not deliquescent. 100 pts. H_2O dissolve 83.9 pts. SnCl_2 at 0° . (Engel, A. ch. (6) **17**, 347.) 100 pts. H_2O dissolve 269.8 pts. SnCl_2 at 15° , and sat. solution has sp. gr. 1.827. (Michel and Krafft, A. ch. (3) **41**, 478.) Sol. in a certain amount of H_2O without decomp., but more H_2O causes pptn. of SnO , SnCl_2 .

$\text{SnCl}_2 + \text{Aq}$ absorbs O from air.

Melts in crystal H_2O at 46° . (Ordway)

Sat. solution boils at 121.7° .

Sp. gr. of $\text{SnCl}_2 + \text{Aq}$ at 15° containing:

5	10	15	20 % $\text{SnCl}_2 + 2\text{H}_2\text{O}$
1.0331	1.0684	1.1050	1.1442

25	30	35	40 % $\text{SnCl}_2 + 2\text{H}_2\text{O}$
1.1855	1.2300	1.2779	1.3298

45	50	55	60 % $\text{SnCl}_2 + 2\text{H}_2\text{O}$
1.3850	1.4451	1.5106	1.5823

65	70	75 % $\text{SnCl}_2 + 2\text{H}_2\text{O}$
1.6598	1.7452	1.8399

(Gerlach, Dingl. 186, 131.)

Solubility of SnCl_2 in $\text{HCl} + \text{Aq}$. $\frac{\text{Sn}}{\text{HCl}}$

$\frac{1}{2}$ molecules SnCl_2 in milligrams 10 ccm. solution; HCl = molecules 1 milligrammes in ditto; H_2O = amt present in grammes.

$\frac{\text{SnCl}_2}{2}$	HCl	Sum of equiv.	sp. gr. of solution
74	■	74	1.532
66.7	6.6	73.3	1.489
63.75	13.54	77.29	1.472
68.4	24.8	93.2	1.524
81.2	34.9	116.1	1.625
94.2	40.0	134.2	1.724
117.6	44	161.6	1.883
147.6	49.4	197.0	2.114
156.4	■	222.4	2.190
157	78	235	2.199

(Engel, A. ch. (6) **17**, 347.)

Solubility is thus diminished by HCl while there are less than 8-10 mols. $\frac{1}{2}$ 1 mol. SnCl_2 . When that limit is passed solubility rapidly increases. (Engel.)

Sol. in very dil. HCl or tartaric acid Sol. in $\text{KOH} + \text{Aq}$. Sol. in conc. SnOCl_2 (Gerlach.) Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$.

Anhydrous SnCl_2 is partially sol. in NH_3 . (Gore, Am. Ch. J. 1898, **20**, 830) Sol. in absolute alcohol. Insol. in turpentine.

11.41 pts. SnCl_2 are sol. in 100 pts. $\text{C}_2\text{H}_5\text{OH}$ at 0° .

11.38 pts. SnCl_2 are sol. in 100 pts. $\text{C}_2\text{H}_5\text{OH}$ at 16° .

11.38 pts. SnCl_2 are sol. in 100 pts. $\text{C}_2\text{H}_5\text{OH}$ at 35.5° . (Lassacynski, B. 1894, **27**, 2286)

Anhydrous SnCl_2 is sol. in ether Jong, Z. anal 1902, **41**, 596.)

1 g. anhydrous SnCl_2 is sol. in 1.8 g. $\text{C}_2\text{H}_5\text{OH}$ at 18° . Sp. gr. of sat. solution $18^\circ/4$ (Naumann, B. 1904, **37**, 4336.)

Sol. in acetone and in methylal. mann, C. C. 1899, II, 1014.)

Anhydrous SnCl_2 is sol. in methylal to the extent of 15.7%. (Schröder Steiner, J. pr. 1909, (2) **79**, 63.)

31.20 pts. $\text{SnCl}_2 + 2\text{H}_2\text{O}$ are sol. in 1 ethyl acetate at -2° .

35.53 pts. $\text{SnCl}_2 + 2\text{H}_2\text{O}$ are sol. in 1 ethyl acetate at $+22^\circ$.

73.44 pts. $\text{SnCl}_2 + 2\text{H}_2\text{O}$ are sol. in 1 ethyl acetate at 82° . (Lassacynski, B. **27**, 2286.)

1 pt. anhydrous SnCl_2 is sol. in 22. ethyl acetate at 18° . $D_{18^\circ/4} = 1.48$ (Naumann, B. 1910, **43**, 319.)

Insol. in ethyl amine (Shann, J. Ch. 1907, **11**, 538); pyridine (Nau B. 1904, **37**, 4609); benzonitrile (Nau B. 1914, **47**, 1369.)

Insol. in CS_2 . (Aretowski, Z. anorg. 1894, 6, 257.)

Sol. in urethane. (Castoro, Z. anorg. 1899, 20, 61.)

Mol. weight determined in pyridine and ethyl sulphide. (Werner, Z. anorg. 1897, 15, 22.)

Tin (stannic) chloride, basic, $\text{SnCl}_4\text{OH} + 3\text{H}_2\text{O}$.

Hygroscopic.

Sol. in H_2O .

Sol. in ether, alcohol, acetone, acetic acid. Nearly insol. in ligroin and benzene. (Pfeiffer Z. anorg. 1914, 37, 241.)

Tin (stannic) chloride, SnCl_4 .

(a) Ordinary modification.—Deliquescent. Sol. in H_2O . On diluting $\text{SnCl}_4 + \text{Aq}$ and boiling, SnO_2 separates out. $\text{SnCl}_4 + \text{Aq}$ is not pptd. by HNO_3 , HCl , or $\text{H}_2\text{SO}_4 + \text{Aq}$; $\text{H}_3\text{PO}_4 + \text{Aq}$ ppts. in a few days, and $\text{H}_2\text{AsO}_4 + \text{Aq}$ in a short time. No ppt is formed by K_2SO_4 , Na_2SO_4 , KCl , NaCl , NH_4Cl , KNO_3 , etc. + Aq .

Sp. gr. of $\text{SnCl}_4 + \text{Aq}$ at 15° .

% SnCl_4 + $5\text{H}_2\text{O}$	Sp. gr.	% SnCl_4 + $5\text{H}_2\text{O}$	Sp. gr.	% SnCl_4 + $5\text{H}_2\text{O}$	Sp. gr.
2	1.012	34	1.226	66	1.538
4	1.024	36	1.242	68	1.563
6	1.036	38	1.259	70	1.587
8	1.048	40	1.276	72	1.614
10	1.059	42	1.293	74	1.641
12	1.072	44	1.310	76	1.669
14	1.084	46	1.329	78	1.698
16	1.097	48	1.347	80	1.727
18	1.110	50	1.366	82	1.759
20	1.124	52	1.386	84	1.791
22	1.137	54	1.406	86	1.824
24	1.151	56	1.426	88	1.859
26	1.165	58	1.447	90	1.894
28	1.180	60	1.468	92	1.932
30	1.195	62	1.491	94	1.969
32	1.210	64	1.514	95	1.988

(Gerlach, Dingl. 178, 49.)

Sp. gr. of $\text{SnCl}_4 + \text{Aq}$.

Deg. Baumé	% Sn	Deg. Baumé	% Sn	Deg. Baumé	% Sn
65.7	29.45	55	24.47	34	14.90
65	29.12	54	24.02	33	14.45
64	28.64	53	23.56	32	14.00
63	28.17	52	23.11	31	13.56
62	27.70	51	22.65	30	13.11
61	27.24	50	22.20	29	12.67
60	26.77	49	21.74	28	12.23
59	26.30	48	21.29	27	11.79
58	25.84	47	20.83	26	11.35
57	25.38	46	20.38	25	10.91
56	24.93				

(Heermann, Ch. Z. 1907, 31, 680.)

Sol. in S_2Cl_2 . (Walden, Z. anorg. 1900, 25, 217.)

Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, 25, 211.)

Very sol. in liquid NH_3 . (Gore, Am. Ch. J. 1899, 20, 830.)

Very sol. in absolute alcohol, from which it is pptd. by H_2O . Easily sol. in ether; decomp. by oil of turpentine. Miscible with CS_2 and Br_2 .

Sol. in acetone. (Naumann, B. 1904, 37, 4328.)

Sol. in acetone and in methylal. (Eidmann, C. C. 1899, II, 1014.)

Sol. in ethyl acetone. (Naumann, B. 1904, 37, 3601.)

Distribution of SnCl_4 between H_2O and xylene.

n = pta. by wt. of Cl in 100 pta. of H_2O layer.

m = pta. by wt. of Cl in 100 pta. of xylene layer.

k = partition coefficient.

50 cc. xylene + 60 g. $\text{SnCl}_4 \cdot 5\text{H}_2\text{O}$.

t°	n	m	k
66°	40.35	0.08	504.4
80°	39.95	0.175	228.5
97.5°	40.24	0.33	122.1
111°	40.27	0.68	59.3

(Smirnov, Z. phys. Ch. 1907, 58, 377.)

50 cc. xylene + 60 g. $\text{SnCl}_4 \cdot 4\text{H}_2\text{O}$.

t°	n	m	k
66°	41.905	0.925	45.3
80°	41.915	1.555	27.0
100°	41.845	2.515	16.7
111°	41.68	3.235	12.9

(Smirnov.)

50 cc. xylene + 60 g. $\text{SnCl}_4 \cdot 3\text{H}_2\text{O}$.

t°	n	m	k
80°	43.205	9.95	4.4
94°	42.545	9.325	4.6
100°	42.645	10.56	5.1
111°	42.31	10.03	4.2

(Smirnov.)

+ $2\text{H}_2\text{O}$. Sol. in H_2O .

+ $3\text{H}_2\text{O}$. Tr. pt. 83° . (Meyerhoffer, Bull. Soc. 1891 (3) 6, 85.)

+ $4\text{H}_2\text{O}$. Tr. pt. 63° . (Meyerhoffer.)

+ $5\text{H}_2\text{O}$. Very deliquescent, and sol. in H_2O . Decomp. by alcohol. Sol. in $\text{HCl} + \text{Aq}$. Tr. pt. 56° . (Meyerhoffer.)

+ $8\text{H}_2\text{O}$. More deliquescent than the $5\text{H}_2\text{O}$ salt. Tr. pt. 19° . (Meyerhoffer.)

+ $9\text{H}_2\text{O}$. (Nöllner, Z. Ch. 1896, 45.)

(b) *Metastannic chloride*.—Sol. in cold H_2O ; solution coagulates on boiling. Conc. HCl + Aq ppts. from $SnCl_4$ + Aq. When solution does not contain HCl , the addition of HCl + Aq causes a ppt., which dissolves in H_2O . HNO_3 , and H_2SO_4 + Aq also ppt. K_2SO_4 , Na_2SO_4 , and $NaCl$ + Aq produce ppts., insol. in H_2O , but sol. in HCl + Aq. NH_4Cl or KCl + Aq do not ppt. KNO_3 + Aq ppts. slowly. (Rose.)

Tin (stannous) hydrogen chloride, $SnCl_2$, $HCl + 3H_2O$.

Decomp. by H_2O .

Melts at -25° . (Engel, C. R. 106. 1398.)

Tin (stannic) hydrogen chloride.

See Chlorostannic acid.

Tin (stannous) hydrazine chloride, $SnCl_2$, $2N_2H_4$, HCl .

Very hygroscopic.

Sol. in H_2O and abs. alcohol. (Curtius, J. pr. 1894, (2) 50. 341.)

Tin (stannic) chloride with MCl .

See Chlorostannate, M.

Tin (stannous) chloride ammonia, $SnCl_2$, NH_3 .

(Berzelius.)

$SnCl_2$, $4NH_3$. Ppt. (Naumann, B. 1904, 37. 4336.)

Tin (stannic) chloride ammonia, $SnCl_4$, $2NH_3$.

Sol. in cold H_2O without decomp., but decomposes by heating.

Tin (stannous) chloride arsenate.

See Arsenate chloride, stannous.

Tin (stannic) chloride cyanhydric acid, $SnCl_4$, $2HCN$.

Decomp. on moist air or with H_2O . (Klein, A. 74. 85.)

Tin (stannous) chloride hydrazine, $SnCl_2$, $2N_2H_4$.

Decomp. by H_2O .

Insol. in NH_4OH + Aq. (Franzen, Z. anorg. 1908, 60. 286.)

Tin (stannic) chloride nitrogen sulphide, $SnCl_4$, $2N_2S_4$.

Insol. in most solvents.

Decomp. by warm NH_4OH + Aq.

Decomposes in the air. (Wölbling, Z. anorg. 1908, 57. 284.)

Decomp. by H_2O . (Davis, Chem. Soc. 1908, 222.)

Tin (stannic) chloride phosphine, $3SnCl_4$, $2PH_3$.

Decomp. by H_2O . (Rose, Pogg. 24.

Tin (stannous) chloride potassium st sulphate.

See Sulphate, potassium stannous st chloride.

Tin (stannic) chloride sulphur tetrachloride, $SnCl_4$, $2SCl_4$.

Very hygroscopic.

Sol. in $CHCl_3$, ligroin, petroleum CS_2 , $POCl_3$; very sol. in complete absolute ether, in benzene, acetaceti and in CCl_4 . (Ruff, B. 1904, 37. 4517.)

Tin (stannic) chloride sulphide, $2SnCl_4$, S .

See Stannic sulphochloride.

Tin (stannic) chlorobromide, $SnClBr_2$.

Decomp. by H_2O . (Ladenburg, A. 8. 60.)

$SnCl_2Br_2$. Decomp. by H_2O . (Ladenburg.)

Tin (stannous) chloriodide, $SnClI_2$.

Decomp. immediately by H_2O . (Phil. Trans. 1845. 363.)

Tin (stannic) chloriodide, $SnCl_4$, I_2 .

Fumes in the air.

Decomp. by H_2O . (Lenormand, J. 1898. 8.)

$SnClI_2$. (Lenormand, J. Pharm. 18 114.)

Tin (stannous) fluoride, SnF_2 .

Easily sol. in H_2O . (Berzelius, Pogg.

Tin (stannic) fluoride, SnF_4 .

Very hygroscopic.

Sol. in H_2O . Slowly decomp. in action with separation of SnO_2 . (Ruff, E 37. 681.)

Tin (stannic) fluoride with MF.

See Fluostannate, M.

Tin (stannous) hydroxide, $2SnO$, H_2O .

Decomp. to SnO when boiled with H_2O . More easily sol. in acids than SnO . Sol. in $NaOH$, and KOH + Aq, even dil. Insol. or very sl. sol. in NH_4OH , $(NH_4)_2CO_3$, and K_2CO_3 + Aq; sol. in CaO , H_2O , and BaO , H_2O , with decomp. on boiling. (Freymy, A. ch. (3) 12. 460.) sl. sol. in NH_4Cl + Aq hot or cold. (Sl. sol. in $NaCH_3O_2$ + Aq. (Mercer.)

Solubility in NaOH + Aq.

G. Na in 20 ccm.	G. Sn in 20 ccm.
0.2480	0.1904
0.3680	0.2614
0.6394	0.4304
0.8326	0.5560
0.9661	0.7849
2.1234	1.8934

(Rubenbauer, Z. anorg. 1902, 30. 335.)

Not pptd. in presence of Na citrate. (Spiller.)

Sol. in water-glass + Aq. (Ordway.)

$\text{SnO}_2 \cdot \text{H}_2\text{O}$. Solubility in 1 l. H_2O = 0.0003135 g. mol. at 25°. (Golde Schmidt, Z. phys. Ch. 1906, 56. 389.)

Tin hydroxide, $\text{SnO} \cdot 6\text{SnO}_2 + 5\text{H}_2\text{O}$.
+ 9 H_2O . (Schiff, A. 120. 153.)

Tin sesquihydroxide, $\text{Sn}_2\text{O}_3 \cdot x\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in NH_4OH + Aq. (Fuchs, J. pr. 5. 318.)

Tin (stannic) hydroxide.

" α " modification.

Obtained by pptn. by alkali in stannic chloride solution.

Freshly pptd. substance when air dried contains 73.5% H_2O ; when dried over H_2SO_4 or in a vacuum for 1 month 12.6% H_2O . Heated to glowing loses all H_2O and passes into the anhydride. The " α " form is capable of existing in all degrees of hydration. (Lorenz, Z. anorg. 1895, 9. 372-375.)

" α " stannic hydrate is a white amorphous substance which is very sol. in HNO_3 when moist; sol. in H_2SO_4 even dil.; sol. in HCl and not pptd. by an excess. Very sol. in NaOH + Aq. and is not pptd. by an excess.

A solution of α stannic acid in HCl is identical with a solution of freshly prepared aqueous stannic chloride and gives no ppt. with dil. HCl , H_2SO_4 , HNO_3 or arsenic acid even on long standing.

" β " modification.

Obtained by oxidizing and dissolving Sn in HNO_3 , and from solution of sodium stannate by pptn. Freshly pptd. from HNO_3 when air dried contains 21.3% H_2O , and when dried over H_2SO_4 or in a vacuum 11.3%,—corresponding to $\text{Sn}(\text{OH})_4$ and $\text{SnO}(\text{OH})_2$ respectively.

Freshly pptd. from sodium stannate solution and air dried contains 22.5% H_2O and when dried over H_2SO_4 or in a vacuum contains 12.1%,—corresponding to $\text{Sn}(\text{OH})_4$ and $\text{SnO}(\text{OH})_2$. Passes into the anhydride when heated to glowing.

The " β " form is capable of existing in all degrees of hydration. It is a white amorphous substance which is insol. in HNO_3 ; in-

sol in H_2SO_4 even when conc.; insol. in HCl but changed by contact with the acid in that when the acid has been removed the ppt. is readily sol. in H_2O , though pptd. again from solution by addition of HCl . When freshly prepared the " β " form is sol. in NaOH + Aq. but is pptd. by an excess of NaOH .

A solution of " β " stannic acid in HCl behaves quite differently from an aq. solution of stannic chloride in that it ppts. metastannic sulphate when treated with H_2SO_4 .

This ppt. dissolves when heated with dilute HNO_3 or HCl , but the solution on standing spontaneously forms another ppt. A solution of " β " stannic acid in HCl gives a ppt. when treated with arsenic acid. (Lorenz, Z. anorg. 1895, 9. 372.)

See also Stannic acid.

Tin hydroxyl chloride, $\text{SnO}(\text{OH})\text{Cl}$.

See Chlorostannic acid.

Tin (stannous) iodide, SnI_2 , and +2 H_2O .

Sl. sol. in cold, more abundantly in hot H_2O , without decomp.

Solubility in H_2O .

t°	Pts. SnI_2 in 100 pts. solution	t°	Pts. SnI_2 in 100 pts. solution
98.5	3.43	97.3	3.70
84.9	3.05	87.4	3.24
73.9	2.56	77.6	2.75
60.1	2.09	67.5	2.34
51.5	1.79	59.7	2.03
41.0	1.50	49.5	1.72
30.5	1.21	39.4	1.38
20.8	1.03	29.6	1.11
	...	19.8	0.96

(Young, J. Am. Chem. Soc. 1897, 19. 846.)

Solubility of SnI_2 in HI + Aq at t°.

Pts. SnI_2 per 100 pts. solvent.

t°	5.83% HI	9.60% HI	15.20% HI	20.44% HI	24.86% HI	30.40% HI	36.83% HI
20	0.98	0.20	0.60	1.81	4.20	10.86	25.31
30	1.16	0.23	0.64	1.81	4.06	10.28	23.46
40	1.40	0.33	0.71	1.90	4.12	10.06	23.15
50	1.69	0.46	0.82	2.12	4.34	10.35	23.76
60	2.07	0.66	1.11	2.51	4.78	11.03	24.64
70	2.48	0.91	1.37	2.92	5.43	11.97	25.72
80	2.95	1.23	1.83	3.70	6.38	13.30	27.23
90	3.46	1.65	2.40	4.58	7.82	15.52	29.84
100	4.03	2.23	3.63	5.82	9.60	...	34.05

(Young, J. Am. Chem. Soc. 1897, 19. 851.)

Solubility of SnI_2 at low temp. in 29.95% $\text{HI} + \text{Aq.}$

Temp.	Pts. in 100 pts. solution	Pts. in 100 pts. solvent
1.5	12.96	14.89
1.5	13.15	15.14
6.0	12.35	14.09
10.5	11.01	12.36
15.2	10.48	11.70
24.8	9.36	10.33
30.7	8.78	9.62
34.8	8.70	9.50
40.3	9.51	10.50

(Young, J. Am. Chem. Soc. 1897, **19**. 854.)Solubility of SnI_2 at low temp. in 39.6% $\text{HI} + \text{Aq.}$

Temp.	Pts. in 100 pts. of solution		Pts. in 100 pts. of solvent
	I	II	
0°	13.52	13.56	15.66
5.7°	16.44	16.37	19.71
10.5°	19.47	19.60	24.27
15.7°	23.56	23.68	30.92
20.3°	25.50	25.60	34.30

(Young, J. Am. Chem. Soc. 1897, **19**. 852-853.)

Sol. in $\text{SnCl}_2 + \text{Aq.}$ Sol. in warm alkali chlorides or iodides + Aq. ; also in dil. $\text{HCl} + \text{Aq.}$ Very sl. sol. in CHCl_3 , CS_2 , or C_6H_6 . (Personne, C. R. **54**. 216.)

Sol. in $\text{KOH} + \text{Aq.}$ (Rose.)Sol. in acetone. (Naumann, B. 1904, **37**. 4328.)**Tin (stannic) iodide, SnI_4 .**Decomp. by H_2O into SnO_2 and HI .Very sol. in PCl_3 . (Beckmann, Z. anorg. 1906, **51**. 110.)Sol. in POCl_3 . (Walden, Z. anorg. 1900, **25**. 212.)Easily sol. in PCl_3 and PBr_3 . (Walden, Z. anorg. 1900, **25**. 211.)Sol. in liquid AsBr_3 forming a solution with sp. gr. = 3.731 at 15°. (Retgers, Z. phys. Ch. 1893, **11**. 342.)Sol. in SOCl_2 , S_2Cl_2 and SO_2Cl_2 . (Walden, Z. anorg. 1900, **25**. 215.)Sol. in SnCl_4 . (Walden.)Sol. in anhydrous alcohol, ether, and benzene. 1 pt. CS_2 dissolves 1.45 pts. SnI_4 at ordinary temp. (Schneider, Pogg. **127**. 624.)100 pts. methylene iodide, CH_2I_2 , dissolve 22.9 pts. SnI_4 at 10°. Sp. gr. of solution = 3.481. (Z. anorg. **8**. 343.)

Solubility in organic solvents at t°.

Solvent	t°	G. SnI_4 in 100 g. of the sat. solution	Sp. gr. of the sat. solution
CCl_4	22.4	5.25	1.59
CCl_4	50.0	12.50	1.63
CHCl_3	28.0	8.21	1.50
C_6H_6	20.2	12.65	0.95

(McDermott, J. Am. Chem. Soc. 1911, **33**. 1964.)Sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)Sol. in acetone. (Eidman, C. C. **1890**, II. 1014.)Solubility in CS_2 .

100 g. of the sat. solution contain at:

−58°	−84°	−89°	−94°	−114.5°
16.27	10.22	9.68	10.65	9.41 g. SnI_4

(Arctowski, Z. anorg. 1896, **11**. 274.)Sol. in allyl mustard oil. (Mathews, J. phys. Ch. 1905, **9**. 647.)**Tin (stannous) hydrogen iodide, SnI_2, HI .**Not obtained in pure state. (Young, J. Am. Chem. Soc. 1897, **19**. 856.)**Tin (stannous) iodide ammonia, $\text{SnI}_2, 2\text{NH}_3$.**(Ephraim and Schmidt, B. 1909, **42**. 3857.) $\text{SnI}_4, 8\text{NH}_3$. (Ephraim and Schmidt.)**Tin (stannic) iodide ammonia, $\text{SnI}_4, 3\text{NH}_3$.**(Personne, C. R. **54**. 218.) $\text{SnI}_4, 4\text{NH}_3$. (Personne.) $\text{SnI}_4, 8\text{NH}_3$. (Rammelsberg, Pogg. **48**. 169.)**Tin iodosulphide,**

See Tin sulphoidide.

Tin monoxide (Stannous oxide), SnO .Insol. in H_2O . Sol. in acids. Very sl. sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq.}$ (Rose.) Insol. in NaOH or $\text{KCH} + \text{Aq.}$ Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 830.)Insol. in acetone. (Naumann, B. 1904, **37**. 4329.)**Tin dioxide (Stannic oxide), SnO_2 .**Insol. in H_2O or conc. acids except conc H_2SO_4 . Insol. in conc. alkalis or $\text{NH}_4\text{OH} + \text{Aq.}$ Not absolutely insol. in dil. $\text{HNO}_3 + \text{Aq.}$ (Mulder.)Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 830.)

Min. Cassiterite (Tin stone). Not attacked by acids.

1 sesquioxide, Sn_2O_3 .

While moist, easily sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ sol. in dil., more easily in conc. $\text{HCl} + \text{Aq.}$ (erzelius.)

n (stannic) oxybromide, $\text{Sn}_2\text{Br}_2\text{O} + 12\text{H}_2\text{O}$.

Decomp. by H_2O into SnBr_2 and H_2SnO_3 . $\text{Sn}_2\text{Br}_2\text{O}_2$. As above. (Preis and Rayann, C. C. 1882. 773.)

n (stannic) oxybromide nitrogen pentoxide, $\text{SnO}_2, 3\text{Br}_2, \text{N}_2\text{O}_5$.

Decomp. by H_2O . (Thomas, C. R. 1896, 2. 33.)

in (stannous) oxychloride, $\text{SnO}, \text{SnCl}_2 + 3\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in HCl , $\text{HC}_2\text{H}_3\text{O}_2$, and l. HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq.}$ (J. Davy, Schw. 10. 325.)

$\text{Sn}_2\text{Cl}_{14}\text{O}_8 + 10\text{H}_2\text{O}$. Easily sol. in H_2O orcohol.

Can be recrystallized from alcohol but not from H_2O . (Tschermak, W. A. B. 44. 736.)

$3\text{SnO}_2, 2\text{SnCl}_2 + 6\text{H}_2\text{O}$. Very sl. sol. in H_2O . Sol. in dil. acids. (Ditte, A. ch. 1882, 27. 146.)

$4\text{SnO}, \text{SnCl}_2 + 6\text{H}_2\text{O}$. (Ditte.)

in (stannic) oxychloride, $\text{SnO}_2, \text{SnCl}_4$.

Sol. in H_2O . (Scheurer-Kestner, A. ch. 47. 6.)

in (metastannic) oxychloride, $3\text{SnO}_2, \text{SnCl}_4 + 3\text{H}_2\text{O}$.

Sol. in little, decomp. by much H_2O . (Weber, Pogg. 122. 368.)

$4\text{SnO}_2, \text{SnCl}_4 + 7\text{H}_2\text{O}$. (Weber.)

"*Metastannyl chloride β* ," $\text{Sn}_2\text{O}_3\text{Cl}_2$. Dequescent. Sol. without decomp. in a small amount of H_2O or in a large amount of H_2O containing a few drops HCl .

Sol. in abs. alcohol. (Engel, C. R. 1897, 24. 767.)

$+4\text{H}_2\text{O}$ and $+9\text{H}_2\text{O}$. Sol. in H_2O acidified with one drop of HCl . Pptd. by excess HCl . (Engel, C. R. 1897, 124. 768.)

"*Parastannyl chloride*," $\text{Sn}_2\text{O}_3\text{Cl}_2 + 2\text{H}_2\text{O}$. Decomp. by excess H_2O .

Sol. in H_2O ; pptd. by HCl . (Engel, C. R. 1897, 125. 465.)

in (stannic) oxychloride nitrogen pentoxide, $\text{SnOCl}_2, 3\text{SnCl}_4, \text{N}_2\text{O}_5$.

Hydroscopic; sol. in H_2O .

Decomp. by heat. (Thomas, C. R. 1896, 22. 32.)

in (stannous) oxyiodide, $\text{SnO}, 3\text{SnI}_2, 2\text{SnO}, 3\text{SnI}_2, \text{SnO}, \text{SnI}_2$; and $2\text{SnO}, \text{SnI}_2$.

Decomp. by much H_2O . (Personne, C. R. 4. 216.)

Tin oxysulphide, $\text{Sn}_2\text{S}_2\text{O} + 11\text{H}_2\text{O}$.

Very sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$; slowly sol. in H_2O . (Schmidt, B. 1894, 27. 2739.)

Tin phosphide, Sn_2P .

(Ragg, C. C. 1898, II. 170.)

SnP . Sol. in $\text{HCl} + \text{Aq.}$ Insol. in $\text{HNO}_3 + \text{Aq.}$

SnP_2 . Not attacked by HCl . Easily attacked by aqua regia. (Emmerling, B. 1879, 12. 155.)

SnP_3 . Insol. in HCl . Slowly attacked by dil. HNO_3 at 50° . Oxidized by fuming HNO_3 with ignition. (Jolibois, C. R. 1909, 148. 638.)

Sn_2P_2 . Insol. in mercury.

Decomp. by HCl . (Stead, J. Soc. Chem. Ind. 1897, 16. 206.)

Sn_4P_2 . Attacked by HCl , HNO_3 , and alkalies. (Jolibois, C. R. 1909, 148. 637.)

The only true compounds are Sn_4P_2 and SnP_3 . (Jolibois, C. R. 1909, 148. 637.)

Tin phosphochloride, $\text{Sn}_2\text{P}_2\text{Cl}_4$.

(Mahn, Jena. Zeit. 5. 1660.)

Tin (stannous) selenide, SnSe .

Decomp. by boiling $\text{HCl} + \text{Aq.}$ Slowly oxidised by boiling $\text{HNO}_3 + \text{Aq.}$ and easily dissolved in aqua regia (Schneider, Pogg. 127. 624.) Easily sol. in alkalies + Aq. (Uelsmann, A. 116. 122), or scarcely even on boiling (Schneider), according to method of preparation. Sol. in alkali sulphides or selenides + Aq.

Tin (stannic) selenide, SnSe_2 .

Not attacked by H_2O or dil. acids; scarcely attacked by boiling conc. $\text{HCl} + \text{Aq.}$; gradually decomp. by hot $\text{HNO}_3 + \text{Aq.}$; easily dissolved by warm aqua regia, and hot conc. H_2SO_4 .

Sol. in cold, more easily in warm KOH , NaOH , or $\text{NH}_4\text{OH} + \text{Aq.}$ (Uelsmann, A. 116. 122.)

Tin (stannous) sulphide, SnS .

1 l. H_2O dissolves 0.14×10^{-6} mols. SnS at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

Insol. in dil., sol. in conc. $\text{HCl} + \text{Aq.}$ Sl. sol. in hot conc. $\text{HNO}_3 + \text{Aq.}$ Insol. in $\text{KOH} + \text{Aq.}$

$+ \text{H}_2\text{O}$. Insol. in H_2O , $\text{H}_2\text{S} + \text{Aq.}$ or dil. acids; sol. with decomp. in conc. acids; easily sol. in hot conc. $\text{HCl} + \text{Aq.}$ Insol. in $\text{H}_2\text{SO}_4 + \text{Aq.}$ Insol. in $\text{NH}_4\text{OH} + \text{Aq.}$ Insol. in NH_4Cl , or $\text{NH}_4\text{NO}_3 + \text{Aq.}$ Scarcely sol. in $(\text{NH}_4)_2\text{S} + \text{Aq.}$ but easily sol. in the same on addition of S . (Rose.)

10% $\text{NaOH} + \text{Aq.}$ dissolves SnS by violent boiling.

Insol. in cold, sl. sol. in hot $\text{Na}_2\text{SO}_3 + \text{Aq.}$ (Materne, C. C. 1906, II. 557.)

Sol. in alkali polysulphides + Aq.

Insol. in acetone. (Eidmann, C. C. 1899, 18. 100.)

II. 1014); (Naumann, B. 1904, 37. 4329.); ethyl acetate. (Naumann, B. 1910, 43. 314.)

Tin (stannic) sulphide, SnS_2 .

Anhydrous. (*Mosaic gold.*) Insol. in HCl or $\text{HNO}_3 + \text{Aq.}$ but decomp. by aqua regia. Sol. in hot $\text{KOH} + \text{Aq.}$ or $\text{K}_2\text{CO}_3 + \text{Aq.}$; also in hot K_2S , $\text{Na}_2\text{S} + \text{Aq.}$, and $(\text{NH}_4)_2\text{S} + \text{Aq.}$

1 l. H_2O dissolves 1.13×10^{-6} mols. SnS_2 at 18° . (Weigel, Z. phys. Ch. 1907, 58. 294.)

+ $x\text{H}_2\text{O}$ Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq.}$ but readily in KOH , K_2S , or $\text{Na}_2\text{S} + \text{Aq.}$; also in hot conc. $\text{HCl} + \text{Aq.}$ Decomp. by hot $\text{HNO}_3 + \text{Aq.}$ Insol. in $\text{KHSO}_4 + \text{Aq.}$ Sol. in $\text{K}_2\text{CO}_3 + \text{Aq.}$ Insol. in NH_4Cl , and $\text{NH}_4\text{NO}_3 + \text{Aq.}$ (Brett.)

Pptd. SnS_2 is insol. in cold, sol. in hot $\text{Na}_2\text{B}_4\text{O}_7 + \text{Aq.}$ Sol. in $\text{Na}_2\text{CO}_3 + \text{Aq.}$ Very sol. in $\text{NaOH} + \text{Aq.}$ (Materna, C. C. 1906, II. 557.)

Sol. in boiling conc. $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq.}$ (Clarke, C. N. 21. 124.)

Insol. in methyl acetate (Naumann, B. 1909, 42. 3790); ethyl acetate (Naumann, B. 1910, 43. 314.); acetone (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Tin sesquisulphide, Sn_2S_3 .

Sol. in moderately conc. HCl. (Antony and Niccoli, Gazz. ch. it. 1892, 22. (2) 408.)

Tin sulphochloride, $\text{SnS}_2 \cdot 2\text{SnCl}_4$.

H_2O_2 dissolves out SnCl_4 . (Dumas, Schw. J. 66. 409.)

$\text{SnS}_2 \cdot \text{Cl}_{11} = \text{SnCl}_4 \cdot 2\text{SnCl}_4$. Sol. in H_2O with separation of S.

Gradually sol. in dil. $\text{HNO}_3 + \text{Aq.}$

Sol. in POCl_3 . (Casselmann, A. 83. 267.)

Tin sulpholodide, SnS_2I_2 .

Decomp. by H_2O into SnO_2 , S, and HI; by cold conc. $\text{HCl} + \text{Aq.}$ with separation of S, also by aqua regia, and $\text{HNO}_3 + \text{Aq.}$

Cold $\text{KOH} + \text{Aq.}$ separates S and SnO_2 .

Completely sol. in hot $\text{KOH} + \text{Aq.}$

Sol. in cold, more easily in hot CS_2 or CHCl_3 .

Decomp. by alcohol. (Schneider, Pogg. 111. 249.)

Tin sulphophosphide, $\text{Sn}_2\text{P}_2\text{S}_5$.

Insol. in HCl, HNO_3 , and aqua regia.

Sol. in aq. alkali hydroxides, containing Cl_2 or Br_2 in solution. (Granger, C. R. 1896, 122. 322.)

Tin (stannous) telluride, SnTe .

Not attacked by conc. $\text{HCl} + \text{Aq.}$ (Ditte, C. R. 97. 42.)

Titanic acid, $\text{TiO}_2 \cdot x\text{H}_2\text{O}$.

α -Titanic acid.—Insol. in H_2O or alcohol. When dried in the cold, is completely sol. in

acids, especially HCl, or dil. $\text{H}_2\text{SO}_4 + \text{Aq.}$ but when the solution in acids is boiled, it is converted into β -titanic acid. Very sl. sol. even when moist in $\text{H}_2\text{SO}_4 + \text{Aq.}$ (Berthier, sl. sol. in alkali carbonates + Aq. A complete solution in an alkali carbonate + Aq. can only be obtained by adding a Ti salt drop by drop to the alkaline solution, and allowing the ppt. to dissolve entirely before adding more Ti salt. On boiling the solution in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ (or in K_2CO_3 or $\text{Na}_2\text{CO}_3 + \text{Aq.}$ with NH_4Cl) the titanic acid is pptd.

Relatively easily sol. in mineral acids, decreasing in the following order HCl, $\text{HN}(\text{O})_2$, H_2SO_4 . Insol. in perchloric acid. (Landecker, Z. anorg. 1909, 64. 67.)

Sol. in dil. H_2SO_4 . 40 g. $\text{H}_2\text{O} + 70$ g. H_2SO_4 (sp. gr. 1.145) dissolves 0.33 g. TiO_2 in 15 min. (Hall and Smith, Proc Am Phil. Soc. 1905, 44. 193.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.)

β -Titanic acid, Metatitanic acid.—Insol. in H_2O , acids except HF, or alkali hydrates or carbonates + Aq. When digested with conc H_2SO_4 until acid is evaporated, the residue is sol. in H_2O . (Berzelius.)

γ -Titanic acid.—Sol. in pure H_2O , but β -acid is pptd. by boiling. (Knop, A. 123 351.)

Colloidal $\text{TiO}_2 \cdot x\text{H}_2\text{O} + \text{Aq.}$ has been prepared by Graham (Chem. Soc. 17. 325.)

Barium titanate, $2\text{BaO} \cdot 3\text{TiO}_2$.

(Bourgeois, C. R. 103. 141.)

Barium pertitanate peroxide.

See Pertitanate, barium peroxide.

Calcium titanate, CaTiO_3 .

(Ebelmen, C. R. 22. 711.)

Min. *Perovskite*. Scarcely attacked by $\text{HCl} + \text{Aq.}$ or other acids, except hot H_2SO_4 , which decomposes it.

$\text{CaO} \cdot 2\text{TiO}_2$. Min. *Titanomorphite*. Partially decomp. by $\text{HCl} + \text{Aq.}$ completely by H_2SO_4 .

Cobaltous titanate, CoTiO_3 .

(Bourgeois, C. C. 1893, I. 226.)

Ferrous orthotitanate, Fe_2TiO_4 .

(Hautefeuille, C. R. 69. 733.)

Ferroferric titanate, $\text{FeTiO}_3 \cdot x\text{Fe}_2\text{O}_3$.

Min. *Menaccasite*. Very sl. sol. in HCl or aqua regia with separation of TiO_2 .

Ferric titanate.

Not attacked by boiling H_2SO_4 or conc $\text{HCl} + \text{Aq.}$ (Wöhler and Liebig, Pogg. 21. 578.)

Magnesium titanate, $MgTiO_3$.

Insol. in H_2O and acids. (Hautefeuille, A. ch. (4) 4. 169.)

Min. *Geikielite*.

When finely powdered, is easily sol. in hot HCl , or in cold HF in a few hours. (Dick, Miner. Mag. 1894, 10. 146.)

Mg_2TiO_4 . Slowly decomp. by boiling with $HNO_3 + Aq$. (Hautefeuille, A. ch. (4) 4. 169.)

Potassium titanate, K_2TiO_3 .

Anhydrous. Decomp. with H_2O .

+ $4H_2O$. Deliquescent. Very sol. in H_2O . Precipitated from aqueous solution by alcohol. (Demoly, Compt. chim. 1849. 325.)

Potassium titanate, acid, $K_2O, 3TiO_2 + 2H_2O$

Insol. in H_2O . (Demoly.)

$K_2O, 6TiO_2 + 2H_2O$. (Demoly.)

$K_2O, 3TiO_2 + 3H_2O$. Insol. in H_2O . Completely sol. in $HCl + Aq$ if only cold H_2O is used for washing. When heated to 100° , no longer completely sol. in $HCl + Aq$. (Rose, Pogg. 74. 563.)

$K_2O, 12TiO_2$. (Rose, Gilb. Ann. 73. 78.)

Sodium titanate, Na_2TiO_3 .

Anhydrous. Decomp. by H_2O into $NaOH$, and an acid titanate, insol. in H_2O .

+ $4H_2O$. Deliquescent. Very sol. in H_2O . Precipitated from aqueous solution by alcohol. (Demoly.)

Sodium titanate, acid, $2Na_2O, 9TiO_2 + 5H_2O$.

If not heated to 100° , is sol. in cold $HCl + Aq$. (Rose, Gilb. Ann. 73. 78.)

$2Na_2O, 3TiO_2$. Insol. in H_2O ; slowly sol. in cold, easily in hot $HCl + Aq$. (Cormimbœuf, C. R. 115. 823.)

$Na_2O, 2TiO_2$. As above. (C.)

$Na_2O, 3TiO_2$. Insol. in H_2O , and nearly so in boiling $HCl + Aq$. (C.)

Strontium titanate, $2SrO, 3TiO_2$.

(Bourgeois, C. R. 103. 141.)

Zinc titanate, $ZnO, TiO_2(?)$.

(Lévy, A. ch. (6) 24. 456.)

$2ZnO, TiO_2(?)$. (Lévy.)

$3ZnO, 2TiO_2$. Slowly attacked by warm H_2SO_4 or $HNO_3 + Aq$, and by $H_2SO_4 + HF$. Wholly sol. in cold $HCl + Aq$. (Lévy.)

$4ZnO, 5TiO_2$. Not attacked by cold conc. acids, but sol. by boiling except in $HCl + Aq$. (Lévy.)

$ZnO, 3TiO_2$. Insol. in H_2O , alcohol, or ether. Dil. HNO_3 , H_2SO_4 , or $HCl + Aq$ do not attack even on boiling; boiling H_2SO_4 dissolves with difficulty; not attacked by conc. boiling alkalies + Aq . (Lévy, A. ch. (6) 25. 471.)

Pertitanic acid.

See Pertitanic acid.

Titanium, Ti .

Decomp. H_2O even under 100° (Wöhler); not attacked by H_2O under 500° . (Kern, C. N. 33. 57.)

Does not decomp. H_2O at 100° . (Schneider, Z. anorg. 1894, 8. 85.)

Sol. in $HCl + Aq$ if warmed. Rapidly sol. in $HF + Aq$. Sol. in cold dil. $H_2SO_4 + Aq$, $HNO_3 + Aq$, or $HC_2H_3O_2 + Aq$. Dissolves almost instantaneously in $HF + Aq$. (Merz.)

Sol. in molten lead and iron; sol. in HCl , HNO_3 , and aqua regia. (Moissan, C. R. 1895, 120. 293.)

Amorphous. Loses its spontaneous inflammability when left for a time in contact with H_2O . (Schneider, Z. anorg. 1895, 8. 85.)

Titanium amide, $Ti(NH_2)_4$.

Violently attacked by H_2O . (Stähler, B. 1905, 38. 2629.)

Titanium tribromide, $TiBr_3 + 6H_2O$.

Very hygroscopic. (Stähler, B. 1904, 37. 4409.)

Titanium tetrabromide, $TiBr_4$.

Deliquescent. Decomp. by H_2O . (Duppa, C. R. 42. 352.)

Sol. in absolute alcohol and in dry ether. (Rosenheim and Schütte, Z. anorg. 1900, 24. 238.)

Titanium bromonitride, $TiNBr$.

Decomp. by a small amount of H_2O . On addition of more H_2O , a part dissolves forming a solution which decomp. on warming with separation of titanous acid. It behaves similarly toward dil. HNO_3 , dil. HCl and dil. H_2SO_4 . Completely sol. in warm dil. H_2SO_4 . (Ruff, B. 1908, 41. 2262.)

Titanium carbide, TiC .

Sol. in $HNO_3 + Aq$. (Shimer, C. N. 55. 71.)

Insol. in HCl . Slowly sol. in aqua regia. (Moissan, C. R. 1895, 120. 295.)

Titanium carbide nitride, $Ti_{10}C_2N_8 = Ti(CN)_2, 3Ti_2N_2$.

Insol. in, and not attacked by boiling HNO_3 or H_2SO_4 (Wollaston), but sol. in $HNO_3 + HF$ (Berzelius).

Titanium dichloride, $TiCl_2$.

Very deliquescent. Decomposes H_2O with violence. Insol. in ether, CS_2 , or $CHCl_3$. Decomp. by 99.5% alcohol.

Titanium trichloride, $TiCl_3$.

Deliquescent. Sol. in H_2O with evolution of heat.

+ $4H_2O$. (Glatzel, B. 9. 1829.)

+ $6H_2O$. Very sol. in H_2O . (Polidori, Z. anorg. 1898, 19. 307.)

Titanium tetrachloride, TiCl_4 .

Anhydrous. Sol. in H_2O with evolution of much heat.

+5 H_2O . Deliquescent.

Titanium sulphuryl chloride, $\text{TiCl}_4\text{SO}_3 = \text{TiCl}_3\text{OSO}_2\text{Cl}$.

Deliquesces gradually in moist air. (Clausnitzer, B. 11. 2011.)

Titanium chloride ammonia, $\text{TiCl}_4 \cdot 4\text{NH}_3$.

Deliquescent. Solution in H_2O is not quite clear. (Rose.)

According to Persoz (A. ch. 46. 315), is $\text{TiCl}_4 \cdot 6\text{NH}_3$.

$\text{TiCl}_4 \cdot 6\text{NH}_3$ and $\text{TiCl}_4 \cdot 4\text{NH}_3$.

Both compds. are unstable in moist air; insol. in ether. (Rosenheim, Z. anorg. 1901, 26. 245.)

$\text{TiCl}_4 \cdot 8\text{NH}_3$. Violently decomp. by H_2O . (Stähler, B. 1905, 38. 2627.)

Titanium tetrachloride cyanobromide, $\text{TiCl}_4 \cdot \text{NCClBr}$.

(Schneider, Z. anorg. 1894, 8. 92.)

Titanium chloride cyanhydric acid, $\text{TiCl}_4 \cdot 2\text{HCN}$.

Deliquescent. Sol. in H_2O with evolution of heat. (Wöhler, A. 73. 226.)

Titanium trichloride nitrogen sulphide, $2\text{TiCl}_3 \cdot \text{N}_4\text{S}_4$.

Decomp. rapidly in air. (Davis, Chem. Soc. 1906, 89. (2) 1576.)

Titanium tetrachloride nitrogen sulphide, $\text{TiCl}_4 \cdot \text{N}_4\text{S}_4$.

Hydrosopic.

Decomp. by H_2O , HNO_3 , HCl , KOH and alcohol. (Wölbling, Z. anorg. 1908, 57. 282.)

Titanium chloride phosphine.

Decomp. by H_2O , $\text{HCl} + \text{Aq}$, $\text{KOH} + \text{Aq}$, $\text{K}_2\text{CO}_3 + \text{Aq}$, or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Rose.)

Titanium tetrachloride phosphoryl chloride, $\text{TiCl}_4 \cdot 2\text{POCl}_3$.

(Ruff, B. 1903, 36. 1783.)

Titanium chloronitride, TiNCl .

Decomp. by small amount cold H_2O . On the addition of more H_2O it is only partially decomp. For complete solution, the addition of dil. HCl or a mixture of warm dil. H_2SO_4 and HF is necessary. Easily sol. in conc. HNO_3 and in conc. H_2SO_4 . (Ruff, B. 1908, 41. 2259.)

Titanium difluoride.

(Hautefeuille, C. R. 57. 151.)

Probably sesquifluoride.

Titanium sesquifluoride, Ti_2F_7 .

Appears to be two modifications, one sol. in H_2O , and the other insol. in H_2O . (Hautefeuille, C. R. 59. 189.)

Insol. in H_2O . (Weber, Pogg. 120. 292.)

Titanium tetrafluoride, TiF_4 .

Decomp. by H_2O . (Unverdorben.)

Sol. in H_2O , but solution decomp. upon evaporation. (Marignac, Ann. Min. (5) 15. 258.)

Sol. in H_2O . (Emich, M. 1904, 25. 910.) Very hygroscopic.

Sol. in H_2O . Sl. sol. in conc. $\text{HF} + \text{Aq}$.

Sol. in cold POCl_3 without decomp. Decomp. in warm POCl_3 .

Sol. in alcohol and dry pyridine.

Insol. in ether, CS_2 , CCl_4 , SiCl_4 , SiBr_4 , SO_2Cl_2 , SOCl_2 , SCl_2 , AsCl_3 , SO_3 , CrO_3 , PCl_5 . (Ruff, B. 1903, 36. 1780.)

+2 H_2O . Sol. in H_2O . (Ruff, B. 1903, 36. 1780.)

Titanium hydrogen fluoride, 2HF , $\text{TiF}_4 = \text{H}_2\text{TiF}_6$.

Sol. in H_2O with decomposition and separation of a basic salt. Corresponds to fluosilicic acid, and may be considered as fluotitanic acid H_2TiF_6 .

Titanium fluoride with MF.

See Fluotitanate, M.

Titanium tetrafluoride ammonia, $\text{TiF}_4 \cdot 2\text{NH}_3$.

Sol. in H_2O ; decomp. in aq. solution on boiling. (Ruff, B. 1903, 36. 1781.)

Titanium monohydroxide, TiO_2H_2 .

Ppt. (Wöhler, A. 73. 49.)

$\text{Ti}_2\text{O}_4\text{H}_2$. Not attacked by cold conc. acids; sl. attacked on warming. Insol. in cold or hot $\text{KOH} + \text{Aq}$. (Winkler, B. 1890, 23. 2659.)

Titanium sequihydroxide, $\text{Ti}_2\text{O}_3 \cdot x\text{H}_2\text{O}$.

Decomposes very quickly with H_2O , forming titanium dihydroxide.

TiO_2H_2 . (Polidori, Z. anorg. 1899, 19. 306.)

Titanium dihydroxide.

See Titanic acid.

Titanium hydroxychloride, $\text{TiCl}_3(\text{OH})$.

Deliquescent. Easily sol. in H_2O and alcohol. Sol. in ether.

$\text{TiCl}_3(\text{OH})_2 + 1\frac{1}{2}\text{H}_2\text{O}$. Deliquescent. Sol. in H_2O , alcohol, and ether. Aqueous solution decomp. by boiling.

$\text{TiCl}(\text{OH})_3 + \text{H}_2\text{O}$. Nearly insol. in H_2O . Insol. in alcohol and ether. (König and v. der Pfordten, B. 21. 1708.)

See also Titanium oxychloride.

Titanium diiodide, TiI_2 .

Very hygroscopic; insol. in organic solvents; sol. in conc. HF and boiling HCl ; decomp. by

lies, H_2SO_4 and HNO_3 . (Defacqs, 3, 147. 66.)

triiodide, $\text{TiI}_3 + 6\text{H}_2\text{O}$.

Hydroscopic. (Stähler, B. 1904, 37.)

tetraiodide, TiI_4 .

on air, and dissolves rapidly in H_2O on addition of heat. Solution decomposes on heating. (Weber.)

nitride, Ti_3N_4 .

Slightly sol. in warm $\text{HNO}_3 + \text{Aq}$. More sol. in aqua regia. (Rose.)

Insol. in dil. acids. Decomp. by hot conc. acid by conc. HNO_3 , especially when added, and by boiling $\text{KOH} + \text{Aq}$. (Eisner, B. 1908, 41. 2252.)

Decomp. by H_2O and dil. acids.

Insol. in all ordinary indifferent organic solvents. (Ruff, B. 1912, 45. 1369.)

Insol. in H_2O . (Wöhler.)

According to Guerin (C. R. 82. 972.)

monoxide, TiO .

Prepared by, C. R. 1895, 120. 290.)

sesquioxide, Ti_2O_3 .

Insol. in HCl or $\text{HNO}_3 + \text{Aq}$. Difficultly sol. in H_2SO_4 . (Ebelmen, A. ch. (3) 20. 392.) Insol. in moist, insol. in H_2O or $\text{NH}_4\text{OH} + \text{Aq}$, slowly decomp. to TiO_2 . Sol. in oxygen on heating quickly decomp. (Berzelius.)

dioxide, TiO_2 .

Amorphous. Insol. in H_2O , HCl , or dil. Aq , even when heated for a long time.

Insol. in conc. H_2SO_4 by long digestion.

Strongly ignited at 1000° , is practically insol. in conc. H_2SO_4 and HF .

Less strongly ignited (by heating metatitanic acid to 700°) it is easily sol. in. (Bornemann and Schirrmeister, 10, II. 1870.)

Amorphous TiO_2 is very difficultly sol. in HF . (Gibson, J. Am. Chem. Soc. 1896, 18.)

Insolubility of ignited TiO_2 in H_2SO_4 is increased by H_2O_2 . (Weiss and Landecker, 1909, 64. 71.)

Insolubility in H_2SO_4 is increased by addition of H_2O_2 . H_2O_2 brings TiO_2 quickly and completely into solution in the presence of OH^- , NH_4Cl , NaOH , Na_2CO_3 , and H_2O_2 . (Weiss and Landecker, Z. anorg. Chem. 71.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1880, 830.)

Minerals. Rutile, Brookite, and Anatase. Solubility as above.

so Titanic acid.

Titanium oxide, Ti_2O_3 .

(Deville, C. R. 53. 163.)

True formula is Ti_7O_{12} . (v. der Pfordten, A. 237. 201.)

Titanium peroxide, TiO_4 .

Sol. in acids. Solution in H_2SO_4 is very stable, but the HCl solution decomposes very easily. (Weber, B. 15. 2599; Piccini, B. 15. 2221; Classen, B. 21. 370.)

Titanium oxychloride, TiO_2 , $\text{TiOCl}_2 + 8\text{H}_2\text{O}$.

Sol. in much H_2O . (Merz, Bull. Soc. 1867. 401.)

$\text{Ti}_2\text{O}_3\text{Cl}_2$. Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$ with separation of TiO_2 .

See also Titanium hydroxychloride.

Titanium oxyfluoride.

Insol. in H_2O . (Berzelius.)

Titanium oxyfluoride with MF.

See Fluoxypertitanate, M.

Titanium phosphide, TiP .

Sl. sol. in boiling aqua regia.

Insol. in dil. or conc. acids and alkalies.

Sl. attacked by fuming HNO_3 in sealed tube at 250° – 300° . (Gewecke, A. 1908, 361. 84.)

Titanium phosphochloride.

See Phosphorus titanium chloride.

Titanium silicide, TiSi_2 .

Sol. in HF ; insol. in other min. acids.

Slowly sol. in 10% $\text{KOH} + \text{Aq}$. (Hönigschmid, C. R. 1906, 143. 226.)

Titanium monosulphide, TiS .

Insol. in alkalies. Difficultly sol. in nitric acid and aqua regia.

Insol. in HF . (v. der Pfordten, A. 234. 257.)

Titanium disulphide, TiS_2 .

Decomp. slowly on moist air. Insol. in HCl or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Ebelmen.)

Sol. in aqua regia or $\text{HNO}_3 + \text{Aq}$. Decomp. by $\text{KOH} + \text{Aq}$ or $\text{NaOH} + \text{Aq}$. Insol. in $\text{KSH} + \text{Aq}$. (Rose.)

Sol. in HF at 100° . (v. der Pfordten, A. 234. 257.)

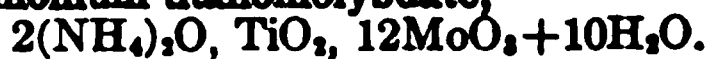
Titanium sesquisulphide, Ti_2S_3 .

Insol. in caustic alkalies + Aq . Sol. in HF at a high temp. Insol. in aqua regia. (v. der Pfordten, A. 234. 257.)

Titanomolybdic acid, TiO_2 , $12\text{MoO}_3 + 22\text{H}_2\text{O}$.

Very sol. in H_2O .

Sol. in ether. (Péchar, C. R. 1893, 117. 790.)

Ammonium titanomolybdate,

Sol. in H_2O and acids; completely insol. in solutions of ammonium salts. (Pèchard.)

Potassium titanomolybdate,

Efflorescent.

Sol. in H_2O . (Pèchard.)

Titanododecatungstic acid, $\text{H}_2\text{TiW}_{12}\text{O}_{42} + x\text{H}_2\text{O}$.

(Lecarme, Bull. Soc. (2) 36. 17.)

Titanotungstic acid or Titanoduodecatungstic acid, $\text{H}_2\text{TiW}_{12}\text{O}_{42} + x\text{H}_2\text{O}$

(Lecarme, Bull. Soc. (2) 36. 17.)

Titanous acid.**Sodium titanite, $\text{Na}_2\text{TiO}_3 = 3\text{Na}_2\text{O}, \text{Ti}_2\text{O}_3$.**

Sol. in dil. acids. (Koenig and v. der Pfordten, B. 22. 2075.)

Titanyl compounds.

See Titanium oxy-compounds.

Triamine cobaltic compounds.

See Dichrocoaltic compounds.

Trithionic acid, $\text{H}_2\text{S}_3\text{O}_6$.

Known only in aqueous solution.

Solution in H_2O gradually decomposes in the cold, rapidly at 80° . Not decomp. if very dilute or in presence of acids, except HNO_3 , HClO_3 , and HIO_3 . (Fordos and Gélis, A. ch. (3) 28. 451.)

Trithionates.

The trithionates are all sol. in H_2O , and very easily decomposed.

Ammonium trithionate, $(\text{NH}_4)_2\text{S}_3\text{O}_6$.

Very deliquescent and unstable.

Very sol. in H_2O .

Insol. in abs. alcohol. (Divers and Ogawa, Chem. Soc. 1900, 77. 337.)

Barium trithionate, $\text{BaS}_3\text{O}_6 + 2\text{H}_2\text{O}$.

Very sol. in H_2O . Precipitated from aqueous solution by large excess of alcohol. Aqueous solution is very unstable. (Kessler, Pogg. 74. 250.)

Lead trithionate, PbS_3O_6 .

Very sl. sol. in H_2O . Sol. in $\text{Na}_2\text{S}_2\text{O}_3 + \text{Aq}$. (Fogh, C. R. 110. 524.)

Potassium trithionate, $\text{K}_2\text{S}_3\text{O}_6$.

Sol. in H_2O . Insol. in alcohol. (Kessler, Pogg. 74. 270.)

Sol. in H_2O with decomp.

Insol. in alcohol. (Langlois, A. 1841, 40. 102.)

Sodium trithionate, $\text{Na}_2\text{S}_3\text{O}_6$.

Very sol. in H_2O .

+ $3\text{H}_2\text{O}$. (Villiers, C. R. 106. 1356.)

Thallous trithionate, $\text{Tl}_2\text{S}_3\text{O}_6$.

Sol. in H_2O . (Bevan, C. N. 33. 294.)

Zinc trithionate.

Sol. in H_2O , but decomposes upon warming the solution. (Fordos and Gélis, C. R. 16. 1070.)

Tungsten, W.

Metallic. Not attacked by heating with fuming HNO_3 , aqua regia, or other acids, or by boiling $\text{KOH} + \text{Aq}$. Sol. in $\text{KOH} + \text{Aq}$ and $\text{NaClO} + \text{Aq}$. (v. Usler, A. 94. 255.)

Not easily acted upon by moist air, if no CO_2 present. Sol. in a mixture of HF and HNO_3 . Very slowly sol. in H_2SO_4 , HCl and HF . (Moissan, C. R. 1896, 123. 15.)

Very slowly attacked by HNO_3 , H_2SO_4 , HCl and even CrO_3 . A mixture of CrO_3 and H_2SO_4 dissolved 1.67 g. in 16 hrs. from a fine wire and 1.36 g. in 14 hours. (Fink, Met. Chem. Eng. 1910, 8. 341.)

Compact tungsten is not attacked by dil., and only sl. dissolved by conc. H_2SO_4 . Not attacked by dil. or conc. HCl . HNO_3 and $\text{HNO}_3 + \text{HCl}$ attack slowly by long heating, forming thin layer of WO_3 . Slowly sol. in $\text{HNO}_3 + \text{HF}$. (Weiss, Z. anorg. 1910, 65. 339.)

Aluminothermic tungsten is insol. in acids and in aqua regia. Sol. in fused KOH (Stavenhagen, B. 1899, 32. 1515.)

Insol. in HCl of any concentration at room temp. and only very sl. sol. at 110° . After being in contact with hot conc. HCl (sp. gr. 1.15) for 175 hrs. the metal lost 0.5% of its weight. Sl. sol. in dil. HCl at 110° .

Insol. in conc. H_2SO_4 at room temp. and in dil. H_2SO_4 at 110° . Somewhat sol. in conc. H_2SO_4 at high temp.

Insol. in conc. HNO_3 , and hot or cold HF .

Sl. sol. in aqua regia.

Very sol. in $\text{HF} + \text{HNO}_3$. (Ruder, J. Am. Chem. Soc. 1912, 34. 387.)

Insol. in aqua regia and acids; sol. in fused KOH . (Stavenhagen, B. 1899, 32. 1514.)

Insol. in $\text{KOH} + \text{Aq}$.

Sol. in fused KOH .

Slowly sol. in fused Na_2CO_3 , K_2CO_3 , or mixture of the two.

Somewhat sol. in $\text{NaOCl} + \text{Aq}$. (Ruder, J. Am. Chem. Soc. 1912, 34. 388.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.)

Crystalline. Insol. in H_2O , HCl , or H_2SO_4 . Oxidised by HNO_3 or aqua regia. (D'Elhujar.)

Sol. in boiling $\text{KOH} + \text{Aq}$. (Riche, A. ch. (3) 50. 5.)

Amorphous. Easily oxidised by $\text{HNO}_3 + \text{Aq}$. (Zettnow.)

Tungsten amide.

See Tungsten nitride.

an arsenide, WAs_2 .

in H_2O and other solvents. Not d by boiling HF or HNO_3 . Sol. in $HF + HNO_3$ and in hot aqua regia. Not d by hot $KOH + Aq$ or $NaOH + Aq$. p. by fused KOH or $NaOH$. (Defacqz, C. R. 1901, **132**. 139.)

an boride, WB_2 .

y attacked by conc. acids; vigorously d by aqua regia. (Tucker and Moody, Soc. 1902, **81**. 16.)

an dibromide, WBr_2 .

y sol. in H_2O , the rest decomposing to d HBr.

an pentabromide, WBr_5 .

mp. by moist air or H_2O . Sol. in alkalis + Aq.

hygroscopic. Fumes in the air.

mp. by H_2O .

n HF, or conc. HCl. Sl. sol. in fuming H_2SO_4 . Readily attacked by fused or alkalis + Aq. Sol. in CCl_4 , $CHCl_3$, abs. alcohol, ether, essence of tere and benzene. (Defacqz, C. R. **28**. 1232.)

an hexabromide, WBr_6 .

mp. by H_2O and in the air.

in $NH_4OH + Aq$. (Smith, J. Am. Soc. 1897, **18**. 1100.)

an bromochloride, WCl_5, WBr_5 .

mp. by H_2O . Sol. in HF. Decomp. O_2 or H_2SO_4 . Violently attacked by alkali or alkali + Aq. Sol. in most solvents.

$3WBr_6$. Properties like those of WBr_6 . (Defacqz, C. R. 1899, **129**. 516.)

mp. by H_2O . Sol. in 40% HF + Aq.

HCl + Aq gives a sl. ppt. of WO_3 . p. by HNO_3 and by H_2SO_4 . Sol. in alcohol, ether, CS_2 , C_6H_6 and glycerine. CCl_4 only on warming. Nearly insol. in turpentine. (Defacqz.)

an bronze.**state tungsten oxide, barium.****state tungsten oxide, barium potas-****state tungsten oxide, barium sodium.****state tungsten oxide, calcium potas-****state tungsten oxide, calcium sodium.****state tungsten oxide, lithium.****state tungsten oxide, lithium potas-****state tungsten oxide, potassium.****state tungsten oxide, potassium so-****Tungstate tungsten oxide, potassium stron-****tium.****Tungstate tungsten oxide, sodium.****Tungstate tungsten oxide, sodium stron-****tium.****Tungsten carbide, W_2C .**

Sol. in boiling HNO_3 ; very slowly acted upon by other acids. (Moissan, C. R. 1896, **123**. 16.)

WC. Insol. in dil. acids; only sl. sol. in H_2SO_4 and conc. HNO_3 ; sol. in fused $KClO_3$ and KNO_3 . (Williams, C. R. 198, **126**. 1724.)

Tungsten dichloride, WCl_2 .

Decomp. on the air or with H_2O . (Roscoe.)

Tungsten tetrachloride, WCl_4 .

Deliquescent. Partly sol. in H_2O , with subsequent decomposition. (Roscoe.)

Tungsten pentachloride, WCl_5 .

Very deliquescent. Decomp. with H_2O with hissing and evolution of heat and separation of W_2O_5 .

Very sl. sol. in CS_2 . (Roscoe.)

Tungsten hexachloride, WCl_6 .

Not decomp. by moist air or H_2O . Decomp. by alcohol. Very sol. in CS_2 . (Roscoe.)

Easily sol. in PCl_5 . (Teclu, A. **187**. 255.)

Tungsten chloride nitrogen sulphide, WCl_4, N_4S_4 .

(Davis, Chem. Soc. 1906, **89**. (2) 1575.)

Tungsten chloroarsenide, W_2AsCl_3 .

Hygroscopic; decomp. by H_2O and acids; sol. in aq. solution of alkalis; insol. in anhydrous organic solvents. (Defacqz, C. R. 1901, **132**. 139.)

Tungsten chlorosulphide, $W_2S_7Cl_3$.

Decomp. by H_2O .

Sol. in S_2Cl_2 . (Smith and Oberholtzer, Z. anorg. 1894, **5**. 68.)

$WCl_6, 3WS_2$. Decomp. by H_2O . Insol. in CS_2 , alcohol and C_6H_6 . (Defacqz, A. ch. 1901, (7) **22**. 266.)

Tungsten hexafluoride, WF_6 .

Fumes in the air.

Decomp. by H_2O . Easily sol. in aq. alkalis. (Ruff, B. 1905, **38**. 747.)

Tungsten diiodide, WI_2 .

Not decomp. by H_2O . (Roscoe, A. **162**. 366.)

Insol. in H_2O , CS_2 and alcohol. Decomp. by boiling H_2O , HNO_3 , H_2SO_4 and aqua regia; sol. in fused KOH , and alkali carbonates. (Defacqz, C. R. 1898, **126**. 938.)

Tungsten tetraiodide, WI_4 .

Insol. in H_2O , ether, chloroform and turpentine; sol. in abs. alcohol; decomp. when boiled with H_2O ; sol. with decomp. in dil. HCl and H_2SO_4 , in HNO_3 and aqua regia, and in alkali hydroxides and carbonates fused or in aq. solution. (Defacqz, C. R. 1898, 127. 511.)

Tritungsten nitride, W_3N_3 .

(Uhrlaub.)

W_3N_3 . Insol. in HNO_3 , dil. H_2SO_4 and $NaOH + Aq$. (Rideal, Chem. Soc. 1889, 55. 44.)

Tungsten nitride amide, $W_3N_3H_4 = 2WN_2, W(NH_2)_2$.

Not attacked by acids or caustic alkalis + Aq . (Wöhler, A. 73. 191.)

Tungsten nitride amide oxide, $W_3N_3H_4O_4 = 3WN_2, W_2(NH_2)_2, 2WO_3$.

Not attacked by acids or alkalis. (Wöhler.)

Tungsten monoxide, WO .

Insol. in H_2O . Not attacked by HCl , HF , H_2SO_4 , or $KOH + Aq$. $HNO_3 + Aq$ or aqua regia convert it into WO_3 . (Headden, Sill. Am. J. 145. 280.)

Tungsten dioxide, WO_2 .

(a) When prepared in the dry way, is attacked only by aqua regia, which oxidises to WO_3 .

(b) When moist, is sol. in HCl or $H_2SO_4 + Aq$, also in $KOH + Aq$. Insol. in $NH_4OH + Aq$. (Riche, A. ch. (3) 50. 5.)

Cryst. Insol. in HCl , H_2SO_4 and conc. aq. alkalis; sol. in HNO_3 . (Hallopeau, C. R. 1898, 127. 135.)

Tungsten oxide, blue.

W_2O_6 (Riche, A. ch. (3) 50. 33); W_3O_8 (v. Uslar); W_4O_{11} (Gmelin).

All are probably the same substance. Not attacked by boiling HNO_3 or aqua regia. Slowly sol. in boiling $KOH + Aq$.

Tungsten trioxide, WO_3 .

Insol. in H_2O or acids. Sl. sol. in dil. $KOH + Aq$, $NaOH + Aq$, $Na_2CO_3 + Aq$, or $H_2CO_3 + Aq$, but easily sol. in conc. boiling solutions of same. $NH_4OH + Aq$ when boiling has a solvent action.

Insol. in conc. and dil. H_2SO_4 . (Desi, J. Am. Chem. Soc. 1897, 19. 214.)

Min. *Tungstite*. Insol. in acids. Sol. in $NH_4OH + Aq$.

Tungsten oxide, W_2O_5 .

Sol. in alkalis. (Desi, J. Am. Chem. Soc. 1897, 19. 214.)

W_2O_5 . Insol. in acids and alkalis. (Desi, J. Am. Chem. Soc. 1897, 19. 228.)

+ H_2O . Like $W_3O_{14} + H_2O$. (Allen and Gottschalk, Am. Ch. J. 1902, 27. 336.)

W_4O_8 . (Desi, J. Am. Chem. Soc. 1897, 19. 219.)

W_5O_{11} . (Desi.)

$W_5O_{14} + H_2O$. Insol. in H_2O containing a little HCl .

Slowly attacked by cold, conc. $MOH + Aq$. (Allen and Gottschalk, Am. Ch. J. 1902, 27. 333.)

Tungsten trioxide ammonia, $WO_3, 3NH_3$.

(Rosenheim and Jacobsen, Z. anorg. 1906, 50. 306.)

Tungsten oxybromide, etc.

See Tungstyl bromide, etc.

Tungsten monophosphide, WP .

Not attacked by HF or HCl .

Sol. in warm $HNO_3 + HF$. Slowly attacked by hot HNO_3 .

Not attacked by $KOH + Aq$ or $NaOH + Aq$. (Defacqz, C. R. 1901, 132. 34.)

Tungsten diphosphide, WP_2 .

Insol. in H_2O and in most organic solvents; insol. in HCl and HF ; sol. in a mixture of HF and HNO_3 in the cold, and in aqua regia on warming. (Defacqz, C. R. 1900, 130. 916)

Tungsten phosphide, W_4P_3 .

Not attacked by any acid, not even by aqua regia. (Wöhler and Wright, A. 79. 24.)

W_3P_4 . (Wöhler and Wright.)

Tungsten diselenide, WSe_2 .

(Uelsmann.)

Tungsten triselenide, WSe_3 .

Easily sol. in alkali sulphides or selenides + Aq . (Uelsmann, Jahrb. f. Ch. 1880. 92.)

Tungsten silicide.

Sol. in HF .

Only very sl. sol. in other acids. (Warren, C. N. 1898, 78. 319.)

WSi_2 . Not attacked by ordinary acids and scarcely by warm aqua regia, but violently attacked by $HNO_3 + HF$. Sl. attacked by 10% alkalis + Aq . (Hönigschmid, M. 1907. 28. 1017.)

Not attacked by dil. or conc. HCl , HF , HNO_3 or H_2SO_4 , nor by aqua regia.

Attacked by $HNO_3 + HF$ or by fused alkalis. (Defacqz, C. R. 1907, 144. 850.)

WSi_3 . Violently attacked by $HNO_3 + HF$. Not attacked by HNO_3 , H_2SO_4 , HCl or HF . (Frlley, Rev. Mét. 1911, 8. 509.)

W_2Si_3 . Insol. in acids including HF ; sol. in a mixture of HF and HNO_3 ; sol. in fused alkali hydroxides and carbonates. (Vigouroux, C. R. 1898, 127. 394.)

Tungsten disulphide, WS_2 .Oxidised by $HNO_3 + Aq.$ (Berzelius.)

Insol. in min. acids.

Sol. in a mixture of HF and HNO_3 and in fused alkalis and alkali carbonates. (Defacqz, C. R. 1899, 128. 611.)**Tungsten trisulphide, WS_3 .**Somewhat sol. in cold, abundantly in hot H_2O , but separated out by the addition of salts, especially NH_4Cl , or acids. Sol. in alkali sulphides, and hydrosulphides + $Aq.$ Sol. in caustic alkalis, and alkali carbonates + $Aq.$ Slowly sol. in $NH_4OH + Aq.$ in the cold.**Tungstic acid, H_2WO_4 .**Insol. in H_2O . Sol. in HF . Insol. in tungstates + $Aq.$ 44.7% H_2WO_4 is sol. in 50% $HF + Aq.$ at 25°.55.3% H_2WO_4 is sol. in 50% $HF + Aq.$ at 50°.100 g. sat. $H_2WO_4 + HCl + Aq.$ contain 0.68 g. H_2WO_4 at 80°.9.8 % H_2WO_4 is sol. in sat. alcoholic HCl at 75°.Insol. in alcoholic solutions of HBr and HI . (Rosenheim, Chem. Soc. 1911, 100. (2) 402.)Freshly pptd. tungstic acid dissolves in H_2O_2 . (Kellner, Dissert. 1909.)Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 830.) H_2WO_4 . Precipitate. Sl. sol. in H_2O and aqueous solutions of the tungstates. Sol. in 250-300 pts. H_2O . When freshly pptd., sol. in alkali hydrates or carbonates + $Aq.$ (Anthon, J. pr. 9. 6.)**Metatungstic acid, $H_2W_{12}O_{41} + 9H_2O$.**Sol. in H_2O . Solution may be boiled and evaporated to a syrupy consistency, when it suddenly gelatinises and ordinary tungstic acid is precipitated.Sol. in H_2O . When heated to 50°, it becomes insol. in H_2O . (Soboleff, Z. anorg. 1896, 12. 28.)**Solubility in H_2O at t° .**

t°	100 ccm. H_2O dissolve g. of the cryst. acid	Sp. gr. of the solution
0	41.46	1.6025
22	88.57	2.5239
43.5	111.87	3.6503

(Soboleff.)

Sp. gr. of solution of metatungstic acid at 17.5° containing:

2.79	12.68	27.61	43.75% WO_3 .
1.0257	1.1275	1.3274	1.6343

(Scheibler, J. pr. 83. 273.)

Sp. gr. of aqueous solution calculated by $M =$ Mendelejeff, and $G =$ (Gerlach (Z. anal. 27. 300)), containing:

	5	10	15	20	25% WO_3 .
M	1.047	1.098	1.153	1.214	1.285
G	1.0469	1.0980	1.1544	1.2172	1.2873

	30	35	40	45	50% WO_3 .
M	1.366	1.458	1.555	1.581 (?)	
G	1.3660	1.4540	1.5527	1.6630	1.7860

Solubility in ether at t° .

t°	100 ccm. ether dissolve g. of the cryst. acid
0	83.456
7.8	88.389
18.2	99.66
24.3	110.76

(Soboleff, Z. anorg. 1896, 12. 32.)

Colloidal. Sol. in H_2O . Not precipitated by acids or alcohol. Can be evaporated to dryness and heated to 200°, and still remains sol. in H_2O . Sol. in $\frac{1}{4}$ pt. of H_2O .

Sp. gr. of aqueous solution containing:

5	20	50	66.5	79.8% WO_3 .
1.0475	1.2168	1.8001	2.596	3.243

(Graham, Chem. Soc. 17. 318.)

Perhaps *paratungstic acid*, $H_{10}W_{12}O_{41}$. (Klein, Bull. Soc. (2) 36. 547.)**Tungstates.**Few normal tungstates are sol. in H_2O , even some of the K and NH_4 salts are very sl. sol. Most of the metatungstates, however, are easily sol. in H_2O .Tungstates insol. in H_2O are usually insol. in dil. acids.**Aluminum tungstate, $Al_2(WO_4)_3 + 8H_2O$.**Precipitate. Insol. in H_2O and $Na_2WO_4 + Aq.$ Sol. in $(NH_4)_2Al_2(SO_4)_4 + Aq.$, $NaOH + Aq.$, $NH_4OH + Aq.$ Easily sol. in H_3PO_4 , $H_2C_2O_4$, and $H_2C_4H_4O_6 + Aq.$ (Lotz, A. 83. 65.)Sol. in 1500 pts. H_2O at 15°. (Lefort, C. R. 87. 748.) $Al_2O_3, 4WO_3 + 9H_2O$. Sol. in 400 pts. H_2O at 15°. (Lefort, C. R. 87. 748.) $Al_2O_3, 5WO_3 + 6H_2O$. Sol. in H_2O , from which it is pptd. by alcohol. (Lefort.)Formula according to Lefort is $Al_2O_3, 3WO_3 + 3H_2O, 2WO_3$.See also *Aluminicotungstic acid*.**Aluminum paratungstate, $5Al_2O_3, 36WO_3 + 46H_2O = Al_2O_3, 7WO_3 + 9H_2O$ (?)**

Easily sol. in an alum solution. (Lotz, A. 83. 65.)

Aluminum ammonium tungstate, $3(NH_4)_2O, Al_2O_3, 9WO_3 + 4H_2O$.Sol. in conc. HNO_3 and in conc. HCl . (Balke and Smith, J. Am. Chem. Soc. 1903, 25. 1230.)

Aluminum ammonium antimony tungstate.

See **Aluminicoantimoniotungstate, ammonium.**

Aluminum antimony tungstate.

See **Aluminicoantimoniotungstic acid.**

Aluminum zinc tungstate, Al_2O_3 , ZnO , 9WO_3 , $+20\text{H}_2\text{O}$.

Very sol. in H_2O . (Daniels, J. Am. Chem. Soc. 1908, **30**. 1850.)

$2\text{Al}_2\text{O}_3$, 3ZnO , $18\text{WO}_3 + 16\text{H}_2\text{O}$. Sol. in much H_2O .

Sol. in very dil. mineral acids or in acetic acid. (Daniels.)

Ammonium tungstate, $(\text{NH}_4)_2\text{WO}_4$.

Known only in solution.

$(\text{NH}_4)_4\text{W}_2\text{O}_{11} + 3\text{H}_2\text{O} = 2(\text{NH}_4)_2\text{O}$, $3\text{WO}_3 + 3\text{H}_2\text{O}$. Sol. in H_2O with decomp. Decomp. on air with evolution of NH_3 , and formation of *paratungstate*. Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Marignac, A. ch. (3) **69**. 23.)

$(\text{NH}_4)_4\text{W}_2\text{O}_{17} + 5\text{H}_2\text{O} = 2(\text{NH}_4)_2\text{O}$, $5\text{WO}_3 + 5\text{H}_2\text{O}$. Sol. at ordinary temp. in 26-29 pts. H_2O with partial decomposition. (Marignac.) $+2\frac{1}{2}\text{H}_2\text{O}$, $+3\text{H}_2\text{O}$, $+4\text{H}_2\text{O}$, $+4\frac{1}{2}\text{H}_2\text{O}$, and $+5\text{H}_2\text{O}$. (Pinagel, Dissert, 1904.)

$(\text{NH}_4)_6\text{W}_3\text{O}_{27} + 8\text{H}_2\text{O} = 3(\text{NH}_4)_2\text{O}$, $8\text{WO}_3 + 8\text{H}_2\text{O}$. Sol. in H_2O . (Marignac.)

Colloidal. $(\text{NH}_4)_2\text{O}$, $6\text{WO}_3 + 4$ or $6\text{H}_2\text{O}$. Miscible with water in nearly all proportions. (Taylor, J. Am. Chem. Soc. 1902, **24**. 632.)

Ammonium metatungstate, $(\text{NH}_4)_2\text{W}_2\text{O}_{11}$.

$+6\text{H}_2\text{O}$. (Marignac, A. ch. (4) **3**. 74.)

$+8\text{H}_2\text{O}$. Efflorescent. Very sol. in H_2O .

1 pt. dissolves at 15° in 0.84 pt. H_2O . (Lotz.)

1 pt. dissolves at ordinary temp. in 0.35 pt. H_2O . (Riche.)

Solubility increases rapidly with the temperature.

Saturated solution at 40° is solid on cooling.

Sl. sol. in ordinary, insol. in absolute alcohol. (Lotz.) Insol. in ether. (Riche.)

$[(\text{NH}_4)_2\text{W}_2\text{O}_{10} + 5\text{H}_2\text{O}]$ of Marguerite.]

$(\text{NH}_4)_6\text{W}_3\text{O}_{21} + 17\text{H}_2\text{O} = 3(\text{NH}_4)_2\text{O}$, $16\text{WO}_3 + 17\text{H}_2\text{O}$. Very efflorescent. Decomp. by dissolving in pure H_2O . (Marignac, A. ch. (4) **3**. 75.)

Ammonium paratungstate, $(\text{NH}_4)_{10}\text{W}_{12}\text{O}_{41} = 5(\text{NH}_4)_2\text{O}$, 12WO_3 .

(Marignac, A. ch. (3) **69**. 25.)

According to Lotz (A. **91**. 49) and Scheibler (J. pr. **80**. 208), formula is $(\text{NH}_4)_6\text{W}_7\text{O}_{24} = 3(\text{NH}_4)_2\text{O}$, 7WO_3 .

$+5\text{H}_2\text{O}$. (Scheibler, J. pr. **48**. 232.)

$+11\text{H}_2\text{O}$. Sol. in 25-28 pts. cold H_2O . (Anthon.)

Sol. in 26.1 pts. H_2O at 10.7° , and 5.8 pts. at 100° . (Lotz.)

Sol. in 33.3 pts. cold H_2O , and 9.6 pts. at 100° . (Riche.)

Sol. in 22-38 pts. H_2O at $15-18^\circ$. The solution gradually decomposes, with the formation of a more soluble salt. (Marignac.)

Not much more sol. in $\text{NH}_4\text{OH} + \text{Aq}$ than in H_2O . Insol. in alcohol. (Anthon.)

Sol. in H_2O_2 . (Kellner, Dissert, 1900.)

Ammonium bismuth tungstate.

See **Bismuthicotungstate, ammonium.**

Ammonium cadmium paratungstate,

$3(\text{NH}_4)_2\text{O}$, 12CdO , $35\text{WO}_3 + 35\text{H}_2\text{O}$.

Ppt. Sol. in H_2O acidulated with HNO_3 . (Lotz, A. **91**. 49.)

Ammonium cerium tungstate.

See **Cericotungstate, ammonium.**

Ammonium cobaltous tungstate, $8(\text{NH}_4)_2\text{O}$, 2CoO , $15\text{WO}_3 + 3\text{H}_2\text{O}$.

(Carnot, C. R. **109**. 147.)

Ammonium hydroxylamine tungstate, $\text{NH}_4\text{OWO}_4\text{NH}_4$.

Sol. in H_2O . (Hofmann, Z. anorg. **1898**. **16**. 465.)

Ammonium iron (ferric) tungstate, $5(\text{NH}_4)_2\text{O}$, Fe_2O_3 , $5\text{WO}_3 + 5\text{H}_2\text{O}$.

Sol. in H_2O . (Borck.)

Ammonium lanthanum tungstate.

See **Lanthanicotungstate, ammonium.**

Ammonium magnesium paratungstate, $2(\text{NH}_4)_2\text{O}$, 3MgO , $12\text{WO}_3 + 24\text{H}_2\text{O}$.

Very slightly sol. in H_2O . (Marignac, A. ch. (3) **69**. 58.)

$(\text{NH}_4)_2\text{O}$, 2MgO , $7\text{WO}_3 + 10\text{H}_2\text{O}$. Very sl. sol. in H_2O ; sol. in H_2O acidulated with HNO_3 . (Lotz.)

Ammonium mercuric tungstate, $(\text{NH}_4)_2\text{WO}_4$, $\text{HgWO}_4 + \text{H}_2\text{O}$.

Insol. in H_2O . Decomp. by acids or alkalis. (Anthon.)

Ammonium neodymium tungstate.

See **Neodymicotungstate, ammonium.**

Ammonium nickel tungstate.

See **Nickelicotungstate, ammonium.**

Ammonium potassium paratungstate, $5\text{K}(\text{NH}_4)\text{O}$, $12\text{WO}_3 + 11\text{H}_2\text{O}$.

Sol. in boiling H_2O ; sl. sol. in cold H_2O . (Hallopeau, C. R. **1896**, **123**. 180.)

Ammonium potassium sodium paratungstate, $5(\text{K}, \text{Na}, \text{NH}_4)_2\text{O}$, $12\text{WO}_3 + 13\text{H}_2\text{O}$, where $\text{K} : \text{Na} : \text{NH}_4 = 3 : 3 : 4$.

$10(\text{K}, \text{Na}, \text{NH}_4)_2\text{O}$, $24\text{WO}_3 + 26\text{H}_2\text{O}$, where $\text{K} : \text{Na} : \text{NH}_4 = 3 : 3 : 4$. (Laurent.)

Ammonium sodium paratungstate, $4(\text{NH}_4)_2\text{O}$, Na_2O , $12\text{WO}_3 + 5\text{H}_2\text{O}$.

Can be crystallised from H_2O without decomp. (Lotz, A. 91. 57.)

+ $14\text{H}_2\text{O}$. Sol. in warm H_2O . (Hallopeau, C. R. 1896, 123. 181.)

$(\text{NH}_4)_2\text{O}$, $4\text{Na}_2\text{O}$, $12\text{WO}_3 + 25\text{H}_2\text{O}$. Sl. sol. in H_2O . (Hallopeau, C. R. 1895, 120. 1344.)

$5\text{Na}_2\text{O}$, $15(\text{NH}_4)_2\text{O}$, $48\text{WO}_3 + 48\text{H}_2\text{O}$. (Marignac, A. ch. (3) 69. 53.)

$2\text{Na}_2\text{O}$, $3(\text{NH}_4)_2\text{O}$, $12\text{WO}_3 + 15\text{H}_2\text{O}$. (Marignac.)

$3(\text{NH}_4)_2\text{O}$, $2\text{Na}_2\text{O}$, $12\text{WO}_3 + 15\text{H}_2\text{O}$. $3(\text{NH}_4)_2\text{O}$, $3\text{Na}_2\text{O}$, $16\text{WO}_3 + 22\text{H}_2\text{O}$. Sol. in H_2O without decomp. (Hallopeau, C. R. 1896, 123. 181.)

$3\text{Na}_2\text{O}$, $4(\text{NH}_4)_2\text{O}$, $16\text{WO}_3 + 18\text{H}_2\text{O}$. (Gibbs, Am. Ch. J. 7. 236.)

Is $2\text{Na}_2\text{O}$, $3(\text{NH}_4)_2\text{O}$, $12\text{WO}_3 + 13\text{H}_2\text{O}$, according to Knorre (B. 19. 823.)

Very sol. in hot H_2O . (Knorre, B. 1886, 19. 823.)

$(\text{NH}_4)_2\text{O}$, $3\text{Na}_2\text{O}$, $16\text{WO}_3 + 38\text{H}_2\text{O}$. (Wyrouboff, Bull. Soc. Min. 1892, 15. 85.)

$6(\text{NH}_4)_2\text{O}$, $2\text{Na}_2\text{O}$, $20\text{WO}_3 + 24\text{H}_2\text{O}$. Can be cryst from boiling H_2O . (Baragiola, Dissert, 1902.)

$4\text{Na}_2\text{O}$, $16(\text{NH}_4)_2\text{O}$, $50\text{WO}_3 + 50\text{H}_2\text{O}$. Sl. sol. in cold H_2O . (Gibbs, Proc. Am. Acad. 15. 12.)

Ammonium zinc paratungstate, $(\text{NH}_4)_2\text{O}$, 2ZnO , $7\text{WO}_3 + 13\text{H}_2\text{O}$.

Sl. sol. in boiling H_2O , but more easily on addition of oxalic, tartaric, phosphoric, or dil. nitric acids, or of ammonium tungstate. (Lotz, A. 91. 49.)

Ammonium zirconium tungstate.

See Zirconotungstate, ammonium.

Ammonium metatungstate nitrate.

See Nitrate metatungstate, ammonium.

Ammonium tungstate vanadate.

See Vanadiotungstate, ammonium.

Antimony tungstate, Sb_2O_3 , $5\text{WO}_3 + 4\text{H}_2\text{O}$.

Sol. in H_2O without decomp. (Lefort.)

Sb_2O_3 , $6\text{WO}_3 + 8\text{H}_2\text{O}$. Ppt.

See also Antimoniotungstic acid.

Barium tungstate, BaWO_4 .

Anhydrous. Insol. in H_2O . Decomp. by boiling $\text{HNO}_3 + \text{Aq}$. (Geuther and Forsberg, A. 120. 270.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Insol. in H_2O or boiling $\text{H}_3\text{PO}_4 + \text{Aq}$. Sol. in boiling, less sol. in cold $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. (Anthon.)

+ $2\frac{1}{2}\text{H}_2\text{O}$. Insol. precipitate. (Scheibler.)

Pptd. BaWO_4 is attacked by dil. acids. More sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$ than in H_2O . (Smith and Bradbury, B. 24. 2930.)

Barium ditungstate, $\text{BaW}_2\text{O}_7 + \text{H}_2\text{O}$ (?).

Nearly insol. in H_2O . 100 ccm. H_2O dissolve about 0.05 g. at 15° . (Lefort, A. ch. (5) 15. 325.)

Barium tritungstate, $\text{BaW}_3\text{O}_{10} + 4\text{H}_2\text{O}$ (?).

Sol. in about 300 pts. H_2O at 15° . Decomp. by boiling H_2O into an insol. salt. (Lefort, C. R. 88. 798.)

+ $6\text{H}_2\text{O}$. (Scheibler.)

Barium metatungstate, $\text{BaW}_4\text{O}_{13} + 9\text{H}_2\text{O}$.

Efflorescent. Quite sol. in hot H_2O . Partly decomp. by cold H_2O into BaW_2O_7 and WO_3 , which recombine on heating. (Scheibler, J. pr. 80. 204.)

Barium tungstate, $\text{BaW}_2\text{O}_7 + 8\text{H}_2\text{O}$.

Insol. in H_2O or $\text{HCl} + \text{Aq}$. (Zettnow.)

$\text{BaW}_2\text{O}_{10}$. Barium bronze. (Hallopeau, A. ch. 1900, (7) 19. 121.)

Barium paratungstate, $\text{Ba}_2\text{W}_{12}\text{O}_{41} + 14\text{H}_2\text{O}$, or $\text{Ba}_2\text{W}_7\text{O}_{24} + 8\text{H}_2\text{O}$.

Insol. in cold H_2O ; when freshly pptd. is sl. sol. in $\text{HNO}_3 + \text{Aq}$. (Lotz, A. 91. 60.)

Sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wackenroder.)

+ $27\text{H}_2\text{O} = \text{Ba}_2\text{W}_7\text{O}_{24} + 16\text{H}_2\text{O}$. Insol. in cold, sl. sol. in hot H_2O . (Knorre, B. 18. 327.)

Barium potassium tungstate tungsten oxide, $\text{BaW}_4\text{O}_{12}$, $5\text{K}_2\text{W}_2\text{O}_{12}$.

(Engels, Z. anorg. 1903, 37. 136.)

Barium silver metatungstate.

(Scheibler.)

Barium sodium paratungstate, 2BaO , $3\text{Na}_2\text{O}$, $12\text{WO}_3 + 24\text{H}_2\text{O}$. (Marignac), or BaO , $2\text{Na}_2\text{O}$, $7\text{WO}_3 + 14\text{H}_2\text{O}$ (Scheibler).

Insol. in H_2O .

Barium sodium tungstate tungsten oxide, $2\text{BaW}_4\text{O}_{12}$, $3\text{Na}_2\text{W}_2\text{O}_{12}$.

$\text{BaW}_4\text{O}_{12}$, $5\text{Na}_2\text{W}_2\text{O}_{12}$. (Engels, Z. anorg. 1903, 37. 131.)

Bismuth tungstate, Bi_2O_3 , $6\text{WO}_3 + 8\text{H}_2\text{O}$.

Very sol. in H_2O with decomp. Pptd. by alcohol from aqueous solution. (Lefort, C. R. 87. 748.)

Cadmium tungstate, CdWO_4 .

Anhydrous.

+ H_2O . Sol. in about 2000 pts. H_2O . (Lefort.)

+ $2\text{H}_2\text{O}$. Insol. in H_2O . Sol. in hot phosphoric or oxalic acids, or in $\text{NH}_4\text{OH} + \text{Aq}$. (Anthon, J. pr. 9. 341.)

Sol. in $\text{KCN} + \text{Aq}$. (Smith and Bradbury, B. 24. 2390.)

Cadmium ditungstate, $\text{CdW}_2\text{O}_7 + 3\text{H}_2\text{O}$ (?).

Sol. in about 500 pts. H_2O at 15° . (Lefort, A. ch. (5) 15. 346.)

Cadmium tritungstate, $\text{CdW}_3\text{O}_{10} + 4\text{H}_2\text{O}$ (?).
(Lefort.)**Cadmium metatungstate, CdO , $4\text{WO}_3 + 10\text{H}_2\text{O}$.**

Not efflorescent. (Scheibler, J. pr. 83. 273.)
Somewhat less sol. in H_2O than the Mn salt.
(Wyruboff, Bull. Soc. Min. 1892, 15. 84.)

Cadmium paratungstate, $\text{Cd}_3\text{W}_7\text{O}_{24} + 16\text{H}_2\text{O}$.

Ppt. (Gonzalez.)
Insol. in H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$, and hot H_3PO_4 , $\text{H}_2\text{C}_2\text{O}_4$, or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

Cadmium sodium paratungstate, 2CdO , Na_2O , $7\text{WO}_3 + 18\text{H}_2\text{O}$.

Difficultly sol. in cold H_2O . (Knorre, B. 19. 824.)

Calcium tungstate, CaWO_4 .

Insol. in H_2O or dil. acids. Sol. in about 500 pts. H_2O . (Lefort.)

Decomp. by $\text{KOH} + \text{Aq}$. (Anthon.)
When freshly pptd., sol. in $\text{NH}_4\text{Cl} + \text{Aq}$. (Wackenroder.)

Sol. in Mg, and NH_4 salts, also $\text{Na}_2\text{WO}_4 + \text{Aq}$. (Sonstadt, C. N. 11. 97.)

Min. *Scheelite*. Decomp. by HCl or $\text{HNO}_3 + \text{Aq}$, with separation of WO_3 .

Calcium ditungstate, $\text{CaW}_2\text{O}_7 + 3\text{H}_2\text{O}$ (?).

Sol. in 30 pts. H_2O at 15° . (Lefort, A. ch. (5) 15. 328.)

Calcium tritungstate, $\text{CaW}_3\text{O}_{10} + 6\text{H}_2\text{O}$ (?).

Sol. in cold H_2O . (Lefort.)

Calcium metatungstate, $\text{CaW}_4\text{O}_{13} + 10\text{H}_2\text{O}$.

Easily sol. in H_2O . (Scheibler.)

Calcium paratungstate, $\text{Ca}_3\text{W}_7\text{O}_{24} + 18\text{H}_2\text{O}$ (or $\text{Ca}_5\text{W}_{12}\text{O}_{41} + 30\text{H}_2\text{O}$).

Much more sol. than Sr or Ba salt. (Knorre B. 18. 328.)

Easily sol. in H_2O_2 . (Knorre, B. 1885, 18. 326.)

Calcium potassium tungstate tungsten oxide, $\text{CaW}_4\text{O}_{12}$, $5\text{K}_2\text{W}_4\text{O}_{12}$.

(Engels, Z. anorg. 1903, 37.149.)

Calcium sodium paratungstate, 2CaO , $3\text{Na}_2\text{O}$, $12\text{WO}_3 + 3\text{H}_2\text{O}$.

(Gonzalez, J. pr. (2) 36. 44.)

Calcium sodium tungstate tungsten oxide, $\text{CaW}_4\text{O}_{12}$, $5\text{Na}_2\text{W}_4\text{O}_{12}$.

Engels Z. anorg. 1903, 37. 145.)

Cerium tungstate, $\text{Ce}_2(\text{WO}_4)_3 + \text{H}_2\text{O}$.

Precipitate. (Cossa and Zecchino, (ch. it. 10. 225.)

Cerium metatungstate, Ce_2O_3 , $12\text{WO}_3 + 30\text{H}_2\text{O}$.

Permanent. Sol. in H_2O . (Scheibler.)

Cerium sodium tungstate, $\text{Ce}_2\text{Na}_2(\text{WO}_4)_3$.

Insol. in H_2O . Slowly sol. in dil. a. easily in $\text{HCl} + \text{Aq}$. (Högbom, Bull. Soc. 42. 2.)

$\text{Ce}_2(\text{WO}_4)_3$, $3\text{Na}_2\text{WO}_4$. (Didier, C. R. 823.)

Cerium tungstate chloride, $3\text{Ce}_2(\text{WO}_4)_3 + 2\text{CeCl}_3$.

(Didier, C. R. 102. 823.)

Chromic tungstate, basic, Cr_2O_3 , $2\text{W}_2\text{O}_7 + 5\text{H}_2\text{O}$.

Sol. in 400 pts. H_2O at 15° . (Lefort, (87. 748.)

Chromic tungstate, $\text{Cr}_2(\text{WO}_4)_3 + 7$, and $13\text{H}_2\text{O}$.

Sol. in $\text{CrCl}_3 + \text{Aq}$, and in phosph. oxalic, or tartaric acids $+ \text{Aq}$. (Lotz.)

$+ 3\text{H}_2\text{O}$. (Lefort, C. R. 87. 748.)
 Cr_2O_3 , $4\text{WO}_3 + 6\text{H}_2\text{O}$. Sol. in about 50 H_2O at 15° . (Lefort.)

Cr_2O_3 , 5WO_3 . Not attacked by aqua r. (Smith and Oberholtzer, Z. anorg. 5. 63.)

Chromic paratungstate, $\text{Cr}_2\text{W}_7\text{O}_{24} + 9\text{H}_2\text{O}$.

Insol. in H_2O or NH_4 paratungstate; sol. in $\text{CrCl}_3 + \text{Aq}$. (Lotz.)

Cobaltous tungstate, CoWO_4 .

Anhydrous. Insol. in H_2O and acids. $+ 2\text{H}_2\text{O}$. Insol. in H_2O and cold HN . Aq. Sl. sol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. Completely in warm H_3PO_4 , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{NH}_4\text{OH} + \text{Aq}$. (Anthon, J. pr. 9. 344.)

Sol. in about 500 pts. H_2O . (Lefort.)

Cobaltous ditungstate, CoW_2O_7 (?).

$+ 3\text{H}_2\text{O}$. Insol. in H_2O . Sl. sol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. Completely sol. in H_3PO_4 , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{NH}_4\text{OH} + \text{Aq}$. (Anthon.) $+ 5\text{H}_2\text{O}$. Sol. in about 100 pts. H_2O . (Lefort.)

$+ 8\text{H}_2\text{O}$ (?). (Lefort.)

Cobaltous tritungstate, $\text{CoW}_3\text{O}_{10} + 4\text{H}_2\text{O}$.

Sol. in H_2O . (Lefort, C. R. 88. 798.)

Cobaltous metatungstate, $\text{CoW}_4\text{O}_{13} + 9\text{H}_2\text{O}$.

Sol. in H_2O . (Scheibler, J. pr. 83. 273.)

Cobaltous paratungstate, $\text{Co}_3\text{W}_7\text{O}_{24} + 25\text{H}_2\text{O}$.

(Gonzalez, J. pr. (2) 36. 44.)

Cobaltous sodium paratungstate, 2CoO , $3\text{Na}_2\text{O}$, $12\text{WO}_3 + 30\text{H}_2\text{O}$.
(Gonzalez.)

Cupric tungstate, CuWO_4 .
+ $2\text{H}_2\text{O}$. Insol. in H_2O . Sol. in H_3PO_4 , $\text{H}_2\text{C}_2\text{O}_4$, or $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. (Anthon.)
100 ccm. H_2O at 15° dissolve 0.1 g. (Lefort.)

Cupric ditungstate, CuW_2O_7 (?).
+ $4\text{H}_2\text{O}$. Insol. in H_2O and HNO_3 . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Anthon, J. pr. 9. 346.)
+ $5\text{H}_2\text{O}$. Sol. in about 300 pts. H_2O . (Lefort.)

Cupric metatungstate, $\text{CuW}_4\text{O}_{13} + 11\text{H}_2\text{O}$.
Sol. in H_2O . (Scheibler.)

Cupric paratungstate, $\text{Cu}_3\text{W}_7\text{O}_{24} + 19\text{H}_2\text{O}$.
Insol. in H_2O . (Knorre, B. 19. 826.)

Cuprocupric tungstate, Cu_2WO_4 , 2CuWO_4 .
Insol. in H_2O . (Zettnow, Pogg. 130. 255.)

Cupric sodium paratungstate, $\text{Cu}_3\text{Na}_5(\text{W}_7\text{O}_{24})_2 + 32\text{H}_2\text{O}$.
Ppt. (Knorre, B. 19. 826.)
 CuO , $4\text{Na}_2\text{O}$, $12\text{WO}_3 + 32\text{H}_2\text{O}$. Ppt. (Gonzalez, J. pr. (2) 36. 52.)

Cupric tungstate ammonia, CuWO_4 , $2\text{NH}_3 + \text{H}_2\text{O}$.
(Schiff, A. 123. 39.)

CuWO_4 , 4NH_3 . Gives off NH_3 at ord. temp. Sol. in H_2O . Sol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Briggs, Chem. Soc. 1904, 85. 676.)

CuO , 4WO_3 , $6\text{NH}_3 + 8\text{H}_2\text{O}$. Insol. in H_2O . Nearly insol. in dil. $\text{NH}_4\text{OH} + \text{Aq}$. (Briggs, Chem. Soc. 1904, 85. 676.)

Didymium tungstate, $\text{Di}_2(\text{WO}_4)_3$.
Precipitate. (Frerichs and Smith, A. 191. 355.)

Didymium metatungstate.
Sol. in H_2O . (Scheibler.)

Didymium sodium tungstate, $\text{DiNa}_3(\text{WO}_4)_3$.
Insol. in H_2O . Slowly sol. in dil. acids. Sol. in conc. $\text{HCl} + \text{Aq}$.
 $\text{DiNa}(\text{WO}_4)_2$. As above. (Högbom, Bull. Soc. (2) 42. 2.)

Erbium sodium tungstate, $\text{Na}_3\text{Er}_4(\text{WO}_4)_3$.
Insol. in H_2O . (Högbom.)

Glucinum metatungstate.
Very sol. in H_2O .

Indium tungstate, $\text{In}_2(\text{WO}_4)_3 + 8\text{H}_2\text{O}$.
Insol. in H_2O . Decomp. by acids. (Renz, Dissert. 1902.)

Iron (ferrous) tungstate, FeWO_4 .

Min. *Ferberite*, *Reinite*.
+ $3\text{H}_2\text{O}$. Insol. in H_2O . Sol. in cold H_2SO_4 , HCl , or $\text{HNO}_3 + \text{Aq}$. Decomp. by boiling acids with separation of WO_3 . Sol. in boiling $\text{H}_3\text{PO}_4 + \text{Aq}$ or warm $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. (Anthon, J. pr. 9. 343.)
+ $x\text{H}_2\text{O}$. Very unstable. (Lefort, A. ch. (5) 15. 314.)

Iron (ferrous) ditungstate, FeW_2O_7 (?).

Insol. in H_2O . Sol. in hot $\text{H}_3\text{PO}_4 + \text{Aq}$ or $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. Decomp. by dil. $\text{HCl} + \text{Aq}$ or by $\text{KOH} + \text{Aq}$. (Ebelmen, C. R. 17. 1198.)
+ $x\text{H}_2\text{O}$. Very unstable. (Lefort.)

Iron (ferrous) tritungstate, $\text{FeW}_3\text{O}_{10} + 4\text{H}_2\text{O}$ (?).

Ppt. Decomp. by cold, more rapidly by hot H_2O . (Lefort.)

Iron (ferrous) metatungstate.

Sol. in H_2O . (Scheibler, J. pr. 83. 315.)

Iron (ferric) tungstate, basic, Fe_2O_3 , $2\text{WO}_3 + 4\text{H}_2\text{O}$.

Sol. in about 50 pts. H_2O . (Lefort.)
 $2\text{Fe}_2\text{O}_3$, $3\text{WO}_3 + 6\text{H}_2\text{O}$. Sol. in about 300 pts. H_2O at 15° . (Lefort.)

Iron (ferric) tritungstate (?), Fe_2O_3 , $4\text{WO}_3 + 4\text{H}_2\text{O} = \text{Fe}_2\text{O}_3$, $3\text{WO}_3 + \text{WO}_3$, $4\text{H}_2\text{O}$ (?).

Sol. in H_2O without decomp. (Lefort.)

Iron (ferric) metatungstate.

Sol. in H_2O . (Scheibler, J. pr. 83. 273.)

Iron (ferrous) manganous tungstate, 7FeWO_4 , MnWO_4 .

(Geuther and Forsberg, A. 120. 277.)
 4FeWO_4 , MnWO_4 . (G. and F.)
 3FeWO_4 , MnWO_4 . Partially sol. in conc. $\text{HCl} + \text{Aq}$. (G. and F.)
 3FeWO_4 , 2MnWO_4 . (G. and F.)
 FeWO_4 , MnWO_4 . (Zettnow, Pogg. 130. 250.)

FeWO_4 , 2MnWO_4 . (G. and F.)
 FeWO_4 , 7MnWO_4 . (G. and F.)
 $x\text{FeWO}_4$, $y\text{MnWO}_4$. Min. *Wolframite*. Sol. in $\text{HCl} + \text{Aq}$, and boiling $\text{H}_3\text{PO}_4 + \text{Aq}$.

Lanthanum tungstate, $\text{La}_2(\text{WO}_4)_3$.

Precipitate.

Lanthanum metatungstate.

Sol. in H_2O . (Scheibler.)

Lanthanum silver tungstate.

See *Lanthanicotungstate, silver*.

Lanthanum sodium tungstate, $\text{Na}_3\text{La}_2(\text{WO}_4)_7$.

Insol. in H_2O . Slowly sol. in dil. acids. Sol. in $\text{HCl} + \text{Aq}$.
 $\text{La}_4\text{Na}_6(\text{WO}_4)_9$. As above. (Högbom, Bull. Soc. (2) 42. 2.)

Lead tungstate, PbWO_4 .

Insol. in H_2O or cold $\text{HNO}_3 + \text{Aq.}$ Sol. in $\text{KOH} + \text{Aq.}$ Decomp. by hot $\text{HNO}_3 + \text{Aq.}$ (Anthon, J. pr. 9. 342.)

Sol. in about 4000 pts. H_2O . (Lefort.)

Min. *Scheelenite*, *Stolzite*. Sol. in $\text{KOH} + \text{Aq.}$; decomp. by HNO_3 .

Absolutely insol. in $\text{NH}_4\text{NO}_3 + \text{Aq.}$ (Smith and Bradbury, B. 24. 2930.)

Lead ditungstate, $\text{PbW}_2\text{O}_7 + 2\text{H}_2\text{O}$ (?).

Sol. in about 80 pts. H_2O at 15° . (Lefort.)

Lead tritungstate, $\text{PbW}_3\text{O}_{10} + 2\text{H}_2\text{O}$ (?).

Ppt. (Lefort.)

Lead metatungstate, $\text{PbW}_4\text{O}_{13} + 5\text{H}_2\text{O}$.

Sl. sol. in cold, more in hot H_2O . Sol. in hot $\text{HNO}_3 + \text{Aq.}$ (Scheibler, J. pr. 83. 318.)

Lead paratungstate, $\text{Pb}_3\text{W}_7\text{O}_{24}$.

Insol. in H_2O , dil. $\text{HNO}_3 + \text{Aq.}$, $(\text{NH}_4)_2\text{WO}_4 + \text{Aq.}$, or $\text{Pb}(\text{NO}_3)_2 + \text{Aq.}$ Sol. in $\text{NaOH} + \text{Aq.}$ or boiling $\text{H}_3\text{PO}_4 + \text{Aq.}$ (Lotz, A. 91. 49.)

Lead sodium paratungstate, PbO , $4\text{Na}_2\text{O}$, $12\text{WO}_3 + 28\text{H}_2\text{O}$.

(Gonzalez.)

Lithium tungstate, Li_2WO_4 .

Rather easily sol. in H_2O . (Gmelin.)

Lithium metatungstate, $\text{Li}_2\text{W}_4\text{O}_{13}$.

Insol. in H_2O . (Knorre, J. pr. (2) 27. 94.)
 $+x\text{H}_2\text{O}$. Syrup. (Scheibler.)

**Lithium paratungstate, $\text{Li}_{10}\text{W}_{12}\text{O}_{41} + 33\text{H}_2\text{O}$
(or $\text{Li}_6\text{W}_7\text{O}_{24} + 19\text{H}_2\text{O}$).**

According to Scheibler, more sol. than the paratungstates of the other alkali metals.

Lithium tungstate tungsten oxide, $\text{Li}_2\text{W}_6\text{O}_{18}$.

Lithium bronze. Insol. in H_2O .

Lithium potassium tungstate tungsten oxide, $\text{Li}_2\text{W}_6\text{O}_{18}$, $3\text{K}_2\text{W}_4\text{O}_{12}$.

Lithium potassium bronze. Insol. in H_2O . (Feit, B. 21. 135.)

Lithium sodium tungstate, $\text{Li}_2\text{WO}_4 + 3\text{H}_2\text{O}$, $3(\text{Na}_2\text{WO}_4 + 3\text{H}_2\text{O})$.

(Traube, N. Jahrb. Miner, 1894, I. 190.)

Magnesium tungstate, MgWO_4 .

Anhydrous. Insol. in H_2O . Gradually decomp. by boiling conc. $\text{HNO}_3 + \text{Aq.}$ (Geuther and Forsberg, A. 120. 272.)

$+3\text{H}_2\text{O}$. Very sol. in H_2O ; nearly insol. in alcohol. (Lefort, A. ch. (5) 15. 329.)

$+7\text{H}_2\text{O}$. Slowly sol. in cold, very easily in hot H_2O . (Ullik, W. A. B. 56. 2. 152.)

Magnesium ditungstate, $\text{MgW}_2\text{O}_7 + 8\text{H}_2\text{O}$ (?).

Sol. in about 100 pts. H_2O . (Lefort.)

Magnesium tritungstate, $\text{MgW}_3\text{O}_{10} + 4\text{H}_2\text{O}$ (?).

Easily sol. in H_2O with gradual decomp. (Lefort.)

Magnesium metatungstate, $\text{MgW}_4\text{O}_{13} + 8\text{H}_2\text{O}$.

Sol. in H_2O . (Scheibler.)

Magnesium paratungstate, $\text{Mg}_3\text{W}_7\text{O}_{24} + 24\text{H}_2\text{O}$.

Very difficultly sol. in cold, somewhat sol. in hot H_2O . (Knorre, B. 19. 825.)

Magnesium potassium tungstate, MgWO_4 , K_2WO_4 .

$+2\text{H}_2\text{O}$. Very sl. sol. in H_2O . (Ullik.)
 $+6\text{H}_2\text{O}$. Precipitate.

Magnesium potassium paratungstate, $5(2/3\text{K}_2\text{O}, 1/3\text{MgO}), 12\text{WO}_3 + 24\text{H}_2\text{O}$.

Insol. in cold, sol. in hot H_2O . (Hallopeau, C. R. 1898, 127. 621.)

Magnesium sodium paratungstate, 3MgO , $3\text{Na}_2\text{O}$, $14\text{WO}_3 + 33\text{H}_2\text{O}$.

Nearly insol. in H_2O . (Knorre, B. 19. 825.)

Manganous tungstate, MnWO_4 .

Min. *Hubnerite*. Partially sol. in $\text{HCl} + \text{Aq.}$

$+2\text{H}_2\text{O}$. Insol. in H_2O ; sol. in warm H_3PO_4 and $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq.}$; sl. sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq.}$ Insol. in cold $\text{HCl} + \text{Aq.}$ (Anthon.)
 $+ \text{H}_2\text{O}$. Sol. in about 2500 pts. H_2O at 15° . (Lefort.)

Manganous ditungstate, $\text{MnW}_2\text{O}_7 + 3\text{H}_2\text{O}$ (?).

Sol. in about 450 pts. H_2O at 15° . (Lefort, A. ch. (5) 15. 333.)

Manganous tritungstate, $\text{MnW}_3\text{O}_{10} + 5\text{H}_2\text{O}$ (?).

Decomp. by H_2O into $\text{MnW}_4\text{O}_{13}$ and $\text{MnW}_4\text{O}_{13}$. (Lefort, A. ch. (5) 17. 484.)

Manganous metatungstate, $\text{MnW}_4\text{O}_{13} + 10\text{H}_2\text{O}$.

Very sol. in H_2O . (Wyrouboff, Bull. Soc. Min. 1892, 15. 82.)

Manganous paratungstate, 5MnO , $12\text{WO}_3 + 34\text{H}_2\text{O}$.

(Gonzalez, J. pr. (2) 36. 44.)

$\text{Mn}_3\text{W}_7\text{O}_{24} + 11\text{H}_2\text{O}$. When recently pptd., sol. in a small amt. of H_2O acidulated with HNO_3 . (Lotz.)

Manganous potassium tungstate, 2MnO , $3\text{K}_2\text{O}$, $12\text{WO}_3 + 16\text{H}_2\text{O}$.

Completely insol. in H_2O . (Hallopeau, Bull. Soc. 1898, (3) 19. 955.)

nous sodium paratungstate, $3\text{Na}_2\text{O}$, InO , $14\text{WO}_3 + 36\text{H}_2\text{O}$.

$n\text{H}_2\text{O}$. (Knorre, B. 19. 826.)

nic sodium tungstate.

Permanganotungstate, sodium.

ous tungstate, Hg_2WO_4 .

$\text{Sol. in H}_2\text{O}$. (Anthon.)

Impossible to obtain pure, as it is decomp.

O , $3\text{WO}_3 + 8\text{H}_2\text{O}$. Sol. in 100 pts. 15° . (Lefort.)

ous metatungstate, $\text{Hg}_2\text{W}_4\text{O}_{13} + \text{H}_2\text{O}$.

(Scheibler, J. pr. 83. 319.)

ic tungstate, HgWO_4 .

$\text{Sol. in H}_2\text{O}$ and very unstable. (Lefort, (5) 15. 356.)

O , 2WO_3 . Insol. in H_2O . (Anthon.)

O , 3WO_3 . Insol. in H_2O . (Anthon.)

O , $5\text{WO}_3 + 5\text{H}_2\text{O}$. Sol. in about 250 O at 15° . (Lefort.)

O , $5\text{WO}_3 + 7\text{H}_2\text{O}$. Decomp. by hot or O . (Lefort, C. R. 88. 798.)

ic tritungstate, $\text{HgW}_3\text{O}_{10} + 7\text{H}_2\text{O}$ (?).

n about 120 pts. H_2O at 15° . (Lefort, (5) 15. 360.)

lenum tungstate.

$\text{Sol. in H}_2\text{O}$. Insol. in $\text{NH}_4\text{Cl} + \text{Aq}$ alcohol of 0.87 sp. gr. (Berzelius.)

nium tungstate, $\text{Nd}_2(\text{WO}_4)_3$.

$\text{Sol. in H}_2\text{O}$. 1 pt. is sol. in 52630 O at 22° ; 59580 pts. at 65° ; 66040 pts. (Hitchcock, J. Am. Chem. Soc. 1895, .)

tungstate, NiWO_4 .

$\text{Sol. in H}_2\text{O}$. Sol. in about 1000 pts. H_2O at 15° .

$\text{Sol. in H}_2\text{O}$. Insol. in H_2O or $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. Boiling $\text{H}_3\text{PO}_4 + \text{Aq}$, $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, or $n\text{NH}_4\text{OH} + \text{Aq}$. (Anthon.)

ditungstate, $\text{NiW}_2\text{O}_7 + 5\text{H}_2\text{O}$ (?).

n about 250 pts. H_2O . (Lefort.)

tritungstate, $\text{NiW}_3\text{O}_{10} + 4\text{H}_2\text{O}$ (?).

$n\text{H}_2\text{O}$. Pptd. by alcohol. Decomp. or warm H_2O after above pptn. (Le-

metatungstate, $\text{NiW}_4\text{O}_{13} + 8\text{H}_2\text{O}$.

$n\text{H}_2\text{O}$. (Scheibler, J. pr. 83. 273.)

paratungstate, $\text{Ni}_3\text{W}_7\text{O}_{24} + 14\text{H}_2\text{O}$.

$\text{Sol. in H}_2\text{O}$. Sl. sol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$. Slightly sol. in warm H_3PO_4 or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Anthon.)

Potassium tungstate, K_2WO_4 .

Anhydrous. Rather deliquescent. Easily sol. in H_2O .

$+ \text{H}_2\text{O}$. Easily sol. in H_2O . Insol. in alcohol.

$+ 2\text{H}_2\text{O}$. Very sol. in H_2O with absorption of heat.

1 pt. dissolves in 1.94 pts. cold, and 0.66 pt. boiling H_2O . Alcohol does not mix with conc. aq. solution, but slowly separates out the salt from it. Acids, even H_2SO_4 , $\text{HC}_2\text{H}_3\text{O}_2$, or $\text{H}_2\text{C}_2\text{O}_4$, separate out WO_3 from solution. (Riche, A. ch. (3) 50. 45.)

Potassium ditungstate, $\text{K}_2\text{W}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Sol. in about 8 pts. H_2O at 15° , but on heating is converted into—

$+ 3\text{H}_2\text{O}$. 100 pts. H_2O dissolve only 2-3 pts. at 15° . (Lefort, A. ch. (5) 9. 102.)

Potassium tritungstate, $\text{K}_2\text{W}_3\text{O}_{10} + 2\text{H}_2\text{O}$.

Sol. in 5-6 pts. H_2O at 15° . Can be recryst. from hot H_2O . (Lefort, A. ch. (5) 9. 105.)

Potassium metatungstate, $\text{K}_2\text{W}_4\text{O}_{13} + 5\text{H}_2\text{O}$.

Not efflorescent. Easily sol. in H_2O . (Marignac.)

($\text{K}_4\text{W}_5\text{O}_{17} + 8\text{H}_2\text{O}$ of Margueritte.)

$+ 8\text{H}_2\text{O}$. Extremely efflorescent. (Scheibler.)

Potassium octotungstate, $\text{K}_2\text{W}_8\text{O}_{26}$.

Insol. in H_2O . (Knorre, J. pr. (2) 27. 49.)

Potassium tungstate, $\text{K}_8\text{W}_{10}\text{O}_{34} + 9\text{H}_2\text{O} = 4\text{K}_2\text{O}$, $10\text{WO}_3 + 9\text{H}_2\text{O}$.

Properties resemble the *paratungstate*. (Gibbs, Proc. Am. Acad. 15. 11.)

$+ 8\text{H}_2\text{O} = \text{K}_4\text{W}_5\text{O}_{17} + 4\text{H}_2\text{O}$. Sol. in 15 pts. H_2O at 15° , but decomposed by heating into $\text{K}_2\text{W}_2\text{O}_7$ and $\text{K}_2\text{W}_3\text{O}_{10}$. (Lefort, A. ch. (5) 9. 104.)

$\text{K}_{10}\text{W}_{14}\text{O}_{47}$. Very difficulty sol. in cold, appreciably sol. in hot H_2O , probably with decomposition. (Knorre.)

Potassium paratungstate, $\text{K}_{10}\text{W}_{12}\text{O}_{41} + 11\text{H}_2\text{O}$ (or $\text{K}_6\text{W}_7\text{O}_{24} + 6\text{H}_2\text{O}$, according to Lots and Scheibler.)

Much more sol. in hot than cold H_2O . (Anthon.)

Sol. in 100 pts. H_2O at 18° , in 8.5 pts. at 100° . (Anthon.)

Sol. in 46.5 pts. cold, and 15.15 pts. boiling H_2O . (Riche.)

By shaking the crystals several days at 20° , 1 pt. dissolves in 71 pts. H_2O . If the salt is treated with boiling water, more goes into solution the longer it is boiled, until after several days' boiling 1 pt. of the salt dissolved in 5.52 pts. H_2O at 18° . Kept in a closed flask, this solution contained after 26 days 1 pt. of salt to 11.9 pts. H_2O ; after 153 days, 1 pt. of salt to 15.6 pts. H_2O ; after 334 days, 1 pt. of salt to 15.6 pts. H_2O . Insol. in alcohol. (Marignac.)

$+ 8\text{H}_2\text{O}$.

Potassium sodium tungstate, K_2WO_4 , $2Na_2WO_4 + 14H_2O$.

Easily sol. in hot and cold H_2O . (Ullik, W. A. B. **56**, 2. 150.)

Deliquescent. Sol. in 1 pt. cold, and $\frac{1}{2}$ pt. hot H_2O . (Anthon.)

Potassium sodium paratungstate, Na_2O , $4K_2O$, $12WO_3 + 15H_2O$.

Sol. in H_2O . (Marignac.)

$\frac{8}{11}Na_2O$, $\frac{3}{11}K_2O$, $12WO_3 + 25H_2O$. Sol. in H_2O . (Marignac.)

Potassium strontium tungstate tungsten oxide, $5K_2W_4O_{12}$, SrW_4O_{12} .

(Engels, Z. anorg. 1903, **37**. 143.)

Potassium uranous tungstate.

See Uranosotungstate, potassium.

Potassium zirconium tungstate.

See Zirconotungstate, potassium.

Potassium tungstate tungsten oxide, K_2WO_4 , W_2O_6 .

Potassium tungsten bronze. (Scheibler, J. pr. 83. 321.)

Formula is $K_2W_4O_{12}$. Not attacked by acids, and only very sl. by alkalis. (Knorre, J. pr. (2) **27**. 49.)

K_2WO_4 , $4WO_3$. Not attacked by acids, even HF , or by alkalis + Aq. Insol. in alcohol. (Zettnow, Pogg, **130**. 262.)

Does not exist. (Knorre.)

Potassium sodium tungstate tungsten oxide, $5K_2W_4O_{12} + 2Na_4W_4O_{16}$.

Potassium sodium tungsten bronze. Properties as potassium bronze.

$3K_2W_4O_{12}$, $2Na_4W_4O_{16}$. As above. (Knorre, J. pr. (2) **27**. 49.)

Praseodymium tungstate, $Pr_2(WO_4)_3$.

Very sl. sol. in H_2O .

Insol. in H_2O at 20° ; at 75° , 1 pt. is sol. in 23,300 pts. H_2O . (Hitchcock, J. Am. Chem. Soc. 1895, **17**. 529.)

Rubidium metatungstate, Rb_2O , $4WO_3 + 8H_2O$.

Sol. in about 10 pts. cold H_2O .

Moderately sol. in warm H_2O . (Wyruboff, Bull. Soc. Min. 1892, **15**. 69.)

Rubidium pentatungstate, $Rb_2W_5O_{18}$.

Almost insol. in hot H_2O . When finely powdered, it is sol. in alkali carbonates + Aq. (Schaeffer, Z. anorg. 1904, **38**. 163.)

Rubidium octotungstate, $Rb_2W_8O_{26}$.

Insol. in H_2O , acids, and alkalis. (Schaeffer, Z. anorg. 1904, **38**. 103.)

Rubidium paratungstate, $5Rb_2O$, $12WO_3 + 18H_2O$.

Very sl. sol. in H_2O . (Schaeffer, Z. anorg. 1904, **38**. 173.)

Samarium metatungstate, Sm_2O_3 , $12WO_3 + 35H_2O$.

Easily sol. in H_2O . (Cleve.)

Samarium sodium tungstate, $Na_2Sm_2WO_6$.

Insol. in H_2O . Slowly sol. in dil acids easily in conc. HCl + Aq. (Höglom, Bull. Soc. (2) **42**. 2.)

Silver (argentous) tungstate, Ag_2O , $2WO_3$.

HNO_3 + Aq separates WO_3 . KOH + Aq dissolves out WO_3 and separates Ag_2O (Wöhler and Rautenberg, A. **114**. 120.)

Does not exist. (Muthmann, B. **20**. 583.)

Silver tungstate, Ag_2WO_4 .

Sol. in about 2000 pts. H_2O at 15° . Easily decomp. by $NaCl$ + Aq or HNO_3 + Aq. (Lefort.)

$Ag_2W_2O_7$. Insol. in H_2O . Nearly insol. in HCl , H_2O_2 or H_3PO_4 + Aq. More sol. in KOH , NH_4OH + Aq. or $H_2C_2O_4$ + Aq. (Anthon, J. pr. **9**. 347.)

+ H_2O . Sol. in about 5000 pts. H_2O at 15° . (Lefort.)

Silver metatungstate, $Ag_2W_4O_{12} + 3H_2O$.

Sl. sol. in H_2O . (Scheibler, J. pr. **83**. 31.)
Nearly insol. in H_2O . (Rosenheim, Z. anorg. 1911, **69**. 250.)

Silver paratungstate, $Ag_{10}W_{12}O_{41} + 8H_2O$.

(Gonzalez, J. pr. (2) **36**. 44.)

Silver tungstate ammonia, Ag_2WO_4 , $4NH_3$.

Sol. in H_2O with rapid decomp. (Widmann, Bull. Soc. (2) **20**. 64.)

Sodium tungstate, $Na_2WO_4 + 2H_2O$.

Sol. in 4 pts. cold, and 2 pts. boiling H_2O . (Vauquelin and Hecht.)

Sol. in 1.1 pts. cold, and 0.5 pt. boiling H_2O . (Anthon.)

Sol. in 2.44 pts. H_2O at 0° ; 1.81 pts. at 15° ; 0.81 pt. at 100° . (Riche.)

Solubility in H_2O at t° .

t°	% Na_2WO_4	Mols. H_2O to 1 mol. Na_2WO_4	Mols. of anhydrous salt to 100 mols. H_2O
-3.5	41.67	22.87	4.37
+0.5	41.73	22.80	4.39
21.0	42.27	22.30	4.48
43.5	43.98	20.80	4.81
80.5	47.65	17.95	5.57
100.0	49.31	16.79	5.95

(Funk, B. 1900, **33**. 3701.)

See also + $10H_2O$.

Sp. gr. of $\text{Na}_2\text{WO}_4 + \text{Aq}$ at 24.5° containing:		
5	10	15 % $\text{Na}_2\text{WO}_4 + 2\text{H}_2\text{O}$,
1.036	1.075	1.119
20	25	30 % $\text{Na}_2\text{WO}_4 + 2\text{H}_2\text{O}$,
1.166	1.215	1.274
35	40	44 % $\text{Na}_2\text{WO}_4 + 2\text{H}_2\text{O}$.
1.349	1.430	1.492
(Franz, J. pr. (2) 4. 238.)		

Sp. gr. of $\text{Na}_2\text{WO}_4 + \text{Aq}$ at 25° .

Sp. gr. at 20°	Per cent Na_2WO_4	Per cent $\text{Na}_2\text{WO}_4 \cdot 2\text{H}_2\text{O}$
1.02016	2.21	2.48
1.03945	4.26	4.78
1.04292	4.59	5.15
1.05831	6.25	7.01
1.07449	7.83	8.79
1.08209	8.61	9.66
1.09687	10.08	11.31
1.12114	12.30	13.81
1.13036	13.16	14.77
1.14392	14.44	16.21
1.16896	16.56	18.62
1.19154	18.52	20.79
1.19938	19.10	21.44
1.20787	19.74	22.16
1.21720	20.59	23.11
1.25041	23.16	25.99
1.25083	23.30	26.15
1.26234	24.05	27.00
1.28143	25.46	28.58
1.33993	29.50	33.11
1.38826	32.68	36.68
1.41072	33.91	38.06
1.47193	37.30	41.87
1.48481	38.20	42.87
1.48595	38.43	43.14

(Pawlewski, B. 1900, **33**. 1224.)

$\text{Na}_2\text{WO}_4 + \text{Aq}$ is pptd. by HCl , HNO_3 , or $\text{H}_2\text{SO}_4 + \text{Aq}$, but not by H_2SO_3 , HI , HCN , oxalic, or tartaric acids + Aq , but pptn. by the former acids is not prevented by presence of the latter, but when heated with $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, or in presence of $\text{H}_3\text{PO}_4 + \text{Aq}$, mineral acids cause no ppt. (Zettnow, Pogg, **130**. 16.)

Much more sol. in H_2O_2 than in H_2O . (Kellner, Dissert, **1909**.)

Sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 829.)

Insol. in alcohol. (Riche, A. ch. (3) **50**. 52.)

Insol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

+ $10\text{H}_2\text{O}$.

Solubility in H_2O at t° .

t°	% Na_2WO_4	Mols. H_2O to 1 mol. Na_2WO_4	Mols. anhydrous salt to 100 mols. H_2O
-5.	30.60	37.04	2.70
-4.0	31.87	34.92	2.86
-3.5	32.98	33.19	3.01
-2.0	34.52	30.90	3.23
0.0	36.54	28.37	3.52
+3.0	39.20	25.33	3.95
+5.0	41.02	23.48	4.26

(Funk, B. 1900, **33**. 3701.)

Sodium ditungstate, $\text{Na}_2\text{W}_2\text{O}_7$.

Sol. in H_2O by heating several hours to $130-150^\circ$. (Knorre, J. pr. (2) **27**. 80.)

+ $6\text{H}_2\text{O}$. Sol. in 13 pts. H_2O at 15° . (Lefort, C. R. **88**. 798.)

Sodium tritungstate, $\text{Na}_2\text{W}_3\text{O}_{10} + 4\text{H}_2\text{O}$.

Sol. in 1 pt. H_2O . Decomp. on standing into sol. *tetratungstate* and insol. *ditungstate*. (Lefort, C. R. **88**. 798.)

Neither this nor the other *tritungstates* of Lefort exist, according to Knorre (J. pr. (2) **27**. 49.)

Sodium metatungstate, $\text{Na}_2\text{W}_4\text{O}_{13}$.

Anhydrous. Insol. in H_2O .

+ $10\text{H}_2\text{O}$. Sol. at 13° in 0.935 pt. H_2O to form a solution of 3.02 sp. gr. (Scheibler.)

Sol. at 19° in 0.195 pt. H_2O . (Forcher.)

Precipitated by alcohol.

Sodium pentatungstate, $\text{Na}_2\text{W}_5\text{O}_{16}$.

Sl. sol. in H_2O by heating 3 hours at 150° . (Knorre, J. pr. (2) **27**. 49.)

Sodium octatungstate, $\text{Na}_2\text{W}_8\text{O}_{26}$.

Insol. in H_2O . Very difficultly attacked by acids and alkalies. (Knorre.)

+ $12\text{H}_2\text{O}$. Easily sol. in cold H_2O , and can be recryst. without decomp. (Ulrik, W. A. B. **56**, 2. 157.)

$3\text{Na}_2\text{O}$, $8\text{WO}_3 + 17\text{H}_2\text{O}$. Very efflorescent. Very sol. in hot H_2O . (Wells, J. Am. Chem. Soc. 1907, **29**. 112.)

Sodium tungstate, $\text{Na}_4\text{W}_7\text{O}_{27}$ (?).

+ $16\text{H}_2\text{O}$ (?). (Marignac, A. ch. (3) **69**. 51.)

+ $21\text{H}_2\text{O}$ (?). Much more sol. and much more rapidly than the paratungstate. (Marignac.)

$\text{Na}_4\text{W}_3\text{O}_{11} + 7\text{H}_2\text{O}$ (?). Mixture of $\text{Na}_2\text{W}_4\text{O}_{13}$ and Na_2WO_4 . (Knorre, J. pr. (2) **27**. 49.)

$\text{Na}_4\text{W}_5\text{O}_{17} + 11\text{H}_2\text{O}$. Efflorescent. Sol. in H_2O . (Marignac.)

100 pts. H_2O dissolve 16 pts. at 15° . (Lefort, A. ch. (5) **9**. 97.)

Formula is $4\text{Na}_2\text{O}$, $10\text{WO}_3 + 23\text{H}_2\text{O}$, according to Gibbs (Proc. Am. Acad. **16**. 5.)

Sodium paratungstate, $\text{Na}_{10}\text{W}_{12}\text{O}_{41} + 21\text{H}_2\text{O}$.
 $+ 25\text{H}_2\text{O}$.

$+ 28\text{H}_2\text{O} = 3\text{Na}_2\text{W}_7\text{O}_{24} + 16\text{H}_2\text{O}$, according to Lotz and Scheibler.

Sol. in 8 pts. cold H_2O (Anthon); in 12.6 pts. at 22° . (Forcher.)

Sol. in about 12 pts. H_2O . (Marignac.)

The aqueous solution saturated at $35-40^\circ$ contained to 1 pt. of the salt, after:

1	12	77	227	410 days,
at 18°	18°	18°	16°	20°
9.25	11.26	10.92	11.90	11.74 pts. H_2O .

The solution saturated by very long boiling, after a part of the salt had crystallised out, contained, after:

1	2	12 days,
0.68	0.91	2.59 pts. H_2O to 1 pt. salt,
72	222	405 days,
6.88	9.75	8.80 pts. H_2O to 1 pt. salt.
(Marignac.)		

Decomp. by boiling with H_2O . (Knorre, B. 18. 2362.)

Sodium strontium paratungstate, Na_2O , 4SrO , $12\text{WO}_3 + 29\text{H}_2\text{O}$.

(Gonzalez, J. pr. (2) 36. 44.)

Sodium strontium tungstate tungsten oxide, $5\text{Na}_2\text{W}_7\text{O}_{24}$, $\text{SrW}_4\text{O}_{12}$.

$12\text{Na}_2\text{W}_7\text{O}_{24}$, $\text{SrW}_4\text{O}_{12}$. (Engels, Z. anorg. 1903, 37. 138.)

Sodium thorium tungstate, $\text{Na}_4\text{Th}(\text{WO}_4)_4$.

Insol. in H_2O . Slowly sol. in dil. acids, easily in conc. $\text{HCl} + \text{Aq}$. (Högbom, Bull. Soc. (2) 42. 2.)

Sodium ytterbium tungstate, Yb_2O_3 , $9\text{Na}_2\text{O}$, 12WO_3 .

Insol. in H_2O . (Cleve, Z. anorg. 1902, 32. 154.)

$2\text{Yb}_2\text{O}_3$, $4\text{Na}_2\text{O}$, 7WO_3 . Ppt. (Cleve.)

Sodium yttrium tungstate, $\text{Na}_3\text{Y}_2(\text{WO}_4)_7$.

Insol. in H_2O , and very slowly attacked by dil. acids. (Högbom, Bull. Soc. (2) 42. 2.)

Sodium zinc paratungstate, Na_2O , 2ZnO , $7\text{WO}_3 + 15\text{H}_2\text{O}$.

Difficultly sol. in cold, more sol. in hot H_2O . (Knorre, B. 19. 823.)

$+ 21\text{H}_2\text{O}$. (Knorre.)

Sodium tungstate tungsten oxide, Na_2WO_4 , W_2O_6 .

Yellow tungsten bronze. Gradually deliquesces on air. Not decomp. by any acid, even aqua regia, except HF , or by alkalies. (Wöhler, Pogg. 2. 350.)

Correct formula is $\text{Na}_2\text{W}_6\text{O}_{14}$, according to Phillip (B. 15. 499.)

Sol. in ammoniacal silver solution with separation of Ag. Easily sol. in boiling alkaline potassium ferricyanide + Aq. (Phillip, B. 12. 2234.)

Na_2WO_4 , $2\text{W}_2\text{O}_6$. *Blue tungsten bronze*. Not attacked by acids or alkalies. (Scheibler.)

Correct formula is $\text{Na}_2\text{W}_6\text{O}_{14}$, according to Phillip (B. 15. 506.)

Sol. in ammoniacal silver solution with separation of Ag.

$\text{Na}_4\text{W}_8\text{O}_{18}$. Properties as above. (Phillip, B. 15. 499.)

$\text{Na}_2\text{W}_2\text{O}_6$. Properties as above. (Phillip.)

Strontium tungstate, SrWO_4 .

Precipitate. (Schultze.)

Sol. in about 700 pts. H_2O . (Lefort.)

Strontium ditungstate, $\text{SrW}_2\text{O}_7 + 3\text{H}_2\text{O}$ (?).

100 ccm. H_2O dissolve 0.35 g. at 15° . (Lefort, A. ch. (5) 15. 326.)

Strontium tritungstate, $\text{SrW}_3\text{O}_{10} + 5\text{H}_2\text{O}$ (?).

Sol. in H_2O with decomp. into SrW_2O_7 and $\text{SrW}_4\text{O}_{13}$. (Lefort, A. ch. (5) 17. 477.)

Strontium metatungstate, $\text{SrW}_4\text{O}_{13} + 8\text{H}_2\text{O}$.

Solubility as calcium *metatungstate*. (Scheibler.)

Extraordinarily sol. in H_2O . (Wyruboff, Bull. Soc. Min. 1892, 15. 63.)

Strontium paratungstate, $\text{Sr}_2\text{W}_7\text{O}_{24} + 16\text{H}_2\text{O}$, or $\text{Sr}_3\text{W}_{12}\text{O}_{41} + 27\text{H}_2\text{O}$.

Insol. in cold, sl. sol. in hot H_2O . (Knorre, B. 18. 327.)

Thallous tungstate, Tl_2WO_4 .

Very sl. sol. in H_2O . Sol. in hot $\text{Na}_2\text{CO}_3 + \text{Aq}$. (Flemming, J. B. 1868. 250.)

Thallous metatungstate, $\text{Tl}_2\text{W}_4\text{O}_{13} + 3\text{H}_2\text{O}$.

Nearly insol. in H_2O . (Rosenheim, Z. anorg. 1911, 69. 251.)

Thallous paratungstate, $5\text{Tl}_2\text{O}$, 12WO_3 .

Insol. in H_2O .

Sol. in $\text{Na}_2\text{CO}_3 + \text{Aq}$. and $\text{KOH} + \text{Aq}$. decomposed by mineral acids. (Schaeffer, Z. anorg. 1904, 38. 171.)

Thallous hydrogen tungstate, TlHWO_4 .

Insol. in H_2O . Difficultly sol. in $\text{NH}_4\text{OH} + \text{Aq}$. Easily sol. in boiling alkali carbonates or hydrates + Aq. (Oettinger, J. B. 1864. 254.)

Thorium tungstate.

Precipitate. (Berzelius.)

Insol. in H_2O .

Tin (stannous) tungstate, $\text{SnWO}_4 + 6\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in oxalic acid and in $\text{KOH} + \text{Aq}$. Slowly sol. in hot $\text{H}_3\text{PO}_4 + \text{Aq}$. (Anthon, J. pr. 9. 341.)

Tin (stannic) tungstate, $9\text{SnO}_2, 13\text{WO}_3$.

Insol. in ammonium tungstate + Aq. Sol. in tin salts + Aq, also in phosphoric, oxalic, or tartaric acids + Aq. (Lotz, A. 91. 49.)

Tungsten tungstate, $\text{WO}_2, \text{WO}_3 = \text{W}_2\text{O}_5$.

See Tungsten oxide, W_2O_5 .

Uranous tungstate, $\text{UO}_2, 3\text{WO}_3 + 6\text{H}_2\text{O}$.

Decomp. by $\text{NaOH} + \text{Aq}$ or $\text{HNO}_3 + \text{Aq}$. Sol. in $\text{HCl} + \text{Aq}$, but not in H_2SO_4 . (Rammelsberg.)

Uranyl tungstate, $\text{UO}_3, \text{WO}_3 + 2\text{H}_2\text{O}$.

Sol. in about 100 pts. H_2O . (Lefort, C. R. 87. 748.)

$\text{UO}_3, 3\text{WO}_3 + 5\text{H}_2\text{O}$ (?). Sol. in about 200 pts. H_2O . (Lefort.)

Vanadium tungstate.

Sl. sol. in H_2O .

Ytterbium tungstate basic, $(\text{YbO})_2\text{WO}_4$:

Ppt. (Cleve, Z. anorg. 1902, 32. 153.)

Ytterbium metatungstate, $\text{Yb}_2\text{O}_3, 12\text{WO}_3 + 35\text{H}_2\text{O}$.

Very sol. in H_2O . (Cleve.)

Yttrium tungstate, $\text{Y}_2(\text{WO}_4)_3 + 6\text{H}_2\text{O}$.

Very sl. sol. in H_2O , but more sol. in $\text{Na}_2\text{WO}_4 + \text{Aq}$. (Berlin.)

Zinc tungstate, ZnWO_4 .

Insol. in H_2O . (Geuther and Forsberg, A. 120. 270.)

+ H_2O . Sol. in 500 pts. H_2O .

Zinc ditungstate, $\text{ZnW}_2\text{O}_7 + 3\text{H}_2\text{O}$ (?).

Sol. in 10 pts. H_2O at 15° , but solution soon decomposes. (Lefort.)

Zinc tritungstate, $\text{ZnW}_3\text{O}_{10} + 5\text{H}_2\text{O}$.

Insol. in boiling H_2O . Sol. in $\text{ZnSO}_4 + \text{Aq}$, or $\text{Na}_4\text{W}_6\text{O}_{17} + \text{Aq}$. (Gibbs.)

Zinc metatungstate, $\text{ZnW}_4\text{O}_{13} + 10\text{H}_2\text{O}$.

Easily sol. in H_2O . Loses crystal H_2O by ignition, and becomes insol. in H_2O . (Scheibler, J. pr. 83. 273.)

+ $8\text{H}_2\text{O}$. More sol. in H_2O than magnesium comp. (Wyruboff, Bull. Soc. Min. 1892, 15. 72.)

Zinc tungstate, $\text{Zn}_4\text{W}_{10}\text{O}_{34} + 18\text{H}_2\text{O} = 4\text{ZnO}, 10\text{WO}_3 + 18\text{H}_2\text{O}$.

Insol. in H_2O . Sol. in excess of zinc sulphate or of sodium tungstate + Aq. (Gibbs, Proc. Am. Acad. 15. 14.)

+ $29\text{H}_2\text{O}$. (Gibbs.)

Zinc paratungstate, $5\text{ZnO}, 12\text{WO}_3 + 37\text{H}_2\text{O}$.

(Gonzalez, J. pr. (2) 36. 44.)

Zinc tungstate, $\text{Zn}_9\text{W}_{22}\text{O}_{76} + 66\text{H}_2\text{O} = 9\text{ZnO}, 22\text{WO}_3 + 66\text{H}_2\text{O}$.

Insol. in H_2O . (Gibbs.)

Zinc tungstate ammonia, $\text{ZnWO}_4, 4\text{NH}_3 + 3\text{H}_2\text{O}$.

Decomp. in the air. (Briggs, Chem. Soc. 1904, 85. 677.)

Pertungstic acid.

See Pertungstic acid.

Tungstoarsenic acid.

See Arseniotungstic acid.

Tungstoboric acid.

See Borotungstic acid.

Tungstocyanhydric acid, $\text{H}_4\text{W}(\text{CN})_8 + 6\text{H}_2\text{O}$.

Hydroscopic.

Sol. in H_2O and abs. alcohol. Insol. in ether, benzene etc. (Olsson, Z. anorg. 1914, 88. 71.)

Ammonium tungstocyanide, $(\text{NH}_4)_4\text{W}(\text{CN})_8$.

Easily sol. in H_2O . Aqueous solution decomp. slowly.

Insol. in organic solvents. (Olsson, Z. anorg. 1914, 88. 62.)

Cadmium tungstocyanide, $\text{Cd}_2\text{W}(\text{CN})_8 + 8\text{H}_2\text{O}$.

Nearly insol. in H_2O . Sl. sol. in dil. HCl . Sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$. Insol. in organic solvents. (Olsson, Z. anorg. 1914, 88. 68.)

Cæsium tungstocyanide, $\text{Cs}_4\text{W}(\text{CN})_8$.

Easily sol. in H_2O forming stable solutions. Insol. in alcohol and other organic solvents (Olsson.)

Calcium tungstocyanide, $\text{Ca}_2\text{W}(\text{CN})_8 + 8\text{H}_2\text{O}$.

Easily sol. in H_2O . Aqueous solution decomp. slowly.

Insol. in organic solvents. (Olsson.)

Lead tungstocyanide, $\text{Pb}_2\text{W}(\text{CN})_8 + 4\text{H}_2\text{O}$.

Sol. in H_2O . Solution decomp. after short time.

Insol. in organic solvents. (Olsson.)

Magnesium tungstocyanide, $\text{Mg}_2\text{W}(\text{CN})_8 + 6\text{H}_2\text{O}$.

Easily sol. in H_2O . Aqueous solution decomp. on heating.

Insol. in organic solvents. (Olsson.)

Manganous tungstocyanide, $\text{Mn}_2\text{W}(\text{CN})_8 + 8\text{H}_2\text{O}$.

Insol. in H_2O and in acids.

Insol. in organic solvents. (Olsson.)

Potassium tungstocyanide, $K_4W(CN)_8 + 2H_2O$.

Easily sol. in H_2O from which it can be cryst. 10 ccm. H_2O dissolve 13-14 g. salt at 18° .

Insol. in alcohol, ether and other organic solvents. (Olsson.)

Rubidium tungstocyanide, $Rb_4W(CN)_8 + 3H_2O$.

Easily sol. in H_2O . Can be cryst. from H_2O . Insol. in alcohol and other organic solvents. (Olsson.)

Silver tungstocyanide, $Ag_4W(CN)_8$.

Insol. in H_2O .

Insol. in acids. Decomp. by dil. HCl .

Sol. in hot conc. $NH_4OH + Aq$.

Insol. in organic solvents. (Olsson.)

Sodium tungstocyanide, $Na_4W(CN)_8 + 2\frac{1}{2}H_2O$.

Hydroscopic.

Easily sol. in H_2O .

Insol. in organic solvents. (Olsson.)

Strontium tungstocyanide, $Sr_2W(CN)_8 + 8H_2O, +9H_2O$.

Easily sol. in H_2O . Aqueous solution decomp. on standing.

Insol. in organic solvents. (Olsson.)

Thallium tungstocyanide, $Tl_4W(CN)_8$.

Difficultly sol. in cold H_2O , more sol. in hot H_2O .

Insol. in organic solvents. (Olsson.)

Zinc tungstocyanide, $Zn_2W(CN)_8 + 4H_2O$.

Insol. in H_2O , and acids.

Sol. in conc. $NH_4OH + Aq$. (Olsson.)

Metatungstoiodic acid.**Ammonium metatungstoiodate, $2(NH_4)_2O, 2I_2O_5, 4WO_3 + 12H_2O$.**

Very sl. sol. in H_2O . (Chrétien, A. ch. 1898, (7) 15. 431.)

Potassium tungstoiodate, $K_2H_3WIO_8$.

(Blomstrand, J. pr. (2) 40. 327.)

$2K_2O, 2I_2O_5, 4WO_3 + 8H_2O$.

5.13 g. are sol. in 1 l. H_2O at 15° ; 8.25 g. at 100° . (Chrétien, A. ch. 1898, (7) 15. 431.)

Tungstoperiodic acid.**Ammonium sodium tungstoperiodate, $2(NH_4)_2O, Na_2O, I_2O_7, 2WO_3 + 16H_2O$.**

Ppt. (Rosenheim, A. 1899, 308. 64.)

Barium tungstoperiodate, $5BaO, I_2O_7, 12WO_3 + 12H_2O$.

Ppt. (Rosenheim.)

Potassium tungstoperiodate, $5K_2O, I_2O_7, 12WO_3 + 8H_2O$.

Sol. in H_2O . (Rosenheim.)

Sodium tungstoperiodate, $3Na_2O, I_2O_7, 2WO_3 + 4H_2O$.

Ppt.

$5Na_2O, I_2O_7, 12WO_3 + 16H_2O$. Sol H_2O . (Rosenheim.)

Strontium tungstoperiodate, $5SrO, I_2O_7, 12WO_3 + 28H_2O$.

Sol. in H_2O . (Rosenheim.)

Tungstophosphoric acid, See Phosphotungstic acid.**Tungstosilicic acid, See Silicotungstic acid.****Tungstotungstic acid.****Lithium tungstotungstate, $Li_2O, WO_2, 3WO_3$.**

Insol. in boiling H_2O and conc. l (Hallopeau, C. R. 1898, 127. 514.)

Potassium tungstotungstate, $K_2O, WO_2, 3WO_3$.

Insol. in hot H_2O , and conc. HCl . Hallopeau, Bull. Soc. 1899, (3) 21. 267.)

Tungstous acid.**Sodium tungstite, $Na_2W_2O_6$.**

See Tungstate tungsten oxide, sodium.

Tungstovanadic acid.

See Vanadiotungstic acid.

Tungstyl dibromide, WO_2Br_2 .

Not decomp. by cold H_2O . (Roscoe.)

Tungstyl tetrabromide, $WOBr_4$.

Extremely deliquescent. Decompose once in moist air or with H_2O .

Tungstyl dichloride, WO_2Cl_2 .

Not decomp. by cold, and but slowly boiling H_2O . Sol. in alkalis and amm

Tungstyl tetrachloride, $WOCl_4$.

Easily decomp. by H_2O or moist air.

Very sol. in CS_2 and S_2Cl_2 . Sl. sol. in benzene. (Smith, J. Am. Chem. Soc. 1896, 18. 1008.)

Tungstyl tetrafluoride, WOF_4 .

Sol. in H_2O with decomp. Very hygroscopic.

Insol. in carbon tetrachloride.

Sl. sol. in carbon bisulphide, dry benzene and ether.

Easily sol. in chloroform and absolute alcohol. (Ruff, Z. anorg. 1907, **52**. 265.)

Tungstyl tetrafluoride ammonia, $2\text{WOF}_4 \cdot \text{NH}_3$.

Sol. in H_2O with decomp.

Insol. in liquid NH_3 . (Ruff, Z. anorg. 1907, **52**. 266.)

Ultramarine blue, $2\text{Na}_2\text{Al}_2\text{Si}_2\text{O}_8 \cdot \text{Na}_2\text{S}_2$ (?)

Not attacked by solutions of alkalis or $\text{NH}_4\text{OH} + \text{Aq}$. Decomp. by acids or acid salts + Aq. Decomp. by alum + Aq.

Ultramarine green, $\text{Na}_2\text{Al}_2\text{Si}_2\text{O}_8 \cdot \text{Na}_2\text{S}$ (?)

Decomp. by mineral acids. Not attacked by alkalis. Decomp. by alum + Aq.

Ultramarine white, $2\text{Na}_2\text{Al}_2\text{Si}_2\text{O}_8 \cdot \text{Na}_2\text{S}$ (?)

Uranic acid, H_2UO_4 .

Insol. in H_2O . Sol. in acids. Very sol. in cold dil. $\text{HNO}_3 + \text{Aq}$. Sl. sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$. Insol. in KOH , NaOH , or $\text{NH}_4\text{OH} + \text{Aq}$. Easily sol. in $(\text{NH}_4)_2\text{CO}_3$, KHCO_3 , and $\text{NaHCO}_3 + \text{Aq}$; less in $\text{K}_2\text{CO}_3 + \text{Aq}$. (Ebelmen.)

Easily sol. in malic and tartaric acids to form complex compds. (Itzig, B. 1901, **34**. 3822.)

H_4UO_6 . Insol. in H_2O ; sol. in acids. (Ebelmen.)

Uranates.

Insol. in H_2O ; sol. in acids.

Ammonium uranate.

Sl. sol. in pure H_2O ; insol. in H_2O containing NH_4Cl or NH_4OH .

Sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. (Peligot, A. ch. (3) **5**. 11.)

$(\text{NH}_4)_2\text{O}$, $4\text{UO}_3 + 7\text{H}_2\text{O}$. (Grubler, Dissert, 1908.)

$(\text{NH}_4)_2\text{O}$, $6\text{UO}_3 + 10\text{H}_2\text{O}$. Insol. in cold and hot H_2O and alkalis + Aq. Very sol. in H_2SO_4 , HCl and acetic acid + Aq. (Zehenter, M. 1900, **21**. 235.)

Barium uranate, BaUO_4 .

Insol. in H_2O . Sol. in dil. acids.

BaU_2O_7 . As above. (Ditte, C. R. **95**. 988.)

$\text{BaU}_3\text{O}_{10} + 4\frac{1}{2}\text{H}_2\text{O}$. Nearly insol. in H_2O , $\text{KOH} + \text{Aq}$. and alcohol.

Easily sol. in cold dil. HCl or HNO_3 and in hot acetic acid. (Zehenter, M. 1904, **25**. 200.)

$\text{Ba}_2\text{U}_3\text{O}_{17} + 8\text{H}_2\text{O}$. Nearly insol. in hot or cold H_2O , $\text{KOH} + \text{Aq}$. and alcohol.

Easily sol. in cold dil. HCl or HNO_3 and in hot acetic acid. (Zehenter.)

$\text{Ba}_2\text{U}_7\text{O}_{23} + 11\text{H}_2\text{O}$. Same properties as $\text{BaU}_3\text{O}_{10}$. (Zehenter.)

Bismuth uranate, $\text{Bi}_2\text{O}_3 \cdot \text{UO}_3 + \text{H}_2\text{O}$.

Min. *Uranosphaerite*.

Calcium uranate, CaUO_4 .

Insol. in H_2O ; sol. in dil. acids. (Ditte, C. R. **95**. 988.)

CaU_2O_7 . Insol. in H_2O ; sol. in dil. acids. (Ditte.)

Cobalt uranate.

Insol. in H_2O ; sol. in $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$. (Persoz, J. pr. **3**. 216.)

Sol. in $\text{HNO}_3 + \text{Aq}$; insol. in $\text{KNO}_3 + \text{Aq}$. (Ebelmen, A. ch. (3) **5**. 222.)

Cupric uranate, CuU_2O_7 .

Insol. in H_2O . (Debray, A. ch. (3) **61**. 451.)

Lead uranate, PbUO_4 .

If ignited, very difficultly sol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Wertheim, J. pr. **29**. 228.)

Insol. in $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2 + \text{Aq}$. (Persoz.) 3PbO , 2UO_3 . Sol. in dil. $\text{HNO}_3 + \text{Aq}$. (Ditte, A. ch. (6) **1**. 338.)

$\text{PbU}_3\text{O}_{10}$. Insol. in H_2O . Sol. in HNO_3 . Insol. in $\text{KOH} + \text{Aq}$, NH_4OH and cold acetic acid. Sol. in hot acetic acid. (Zehenter, M. 1904, **25**. 215.)

$\text{Pb}_4\text{U}_5\text{O}_{19} + 4\text{H}_2\text{O}$. Insol. in hot or cold H_2O . Sol. in HNO_3 . Insol. in $\text{KOH} + \text{Aq}$, NH_4OH , alcohol and ether. Sl. sol. in cold, more easily sol. in hot acetic acid. (Zehenter.)

Lithium uranate, Li_2UO_4 .

Insol. in H_2O , but decomp. thereby. Sol. in dil. acids.

Magnesium uranate, MgUO_4 .

Insol. in H_2O . Nearly insol. in cold $\text{HCl} + \text{Aq}$. Slowly sol. in $\text{HCl} + \text{Aq}$ on warming, and more rapidly by addition of a little $\text{HNO}_3 + \text{Aq}$. (Ditte.)

MgU_2O_7 . Ppt. (Berzelius.)

Neodymium uranate, $\text{Nd}_2(\text{U}_3\text{O}_{10})_3 + 18\text{H}_2\text{O}$.

Ppt. (Orloff, Ch. Z. 1907, **31**. 1119.)

Potassium uranate, K_2UO_4 (?)

Insol. in H_2O ; sol. in dil. acids, etc., exactly as Na_2UO_4 . (Ditte.)

$\text{K}_2\text{U}_2\text{O}_7 + 6\text{H}_2\text{O}$. Insol. in H_2O . Sol. in dil. acids, even acetic acid. (Zimmermann, B. **14**. 440.)

Insol. in $\text{K}_2\text{CO}_3 + \text{Aq}$, but easily sol. in alkali hydrogen carbonates + Aq. Sol. in $\text{HCl} + \text{Aq}$. (Ebelmen, A. ch. (3) **5**. 220.)

K_2O , $4\text{UO}_3 + 5\text{H}_2\text{O}$. (Zehenter, M. 1900, **21**. 235.)

K_2O , $6\text{UO}_3 + 6\text{H}_2\text{O}$. Insol. in H_2O . (Drenckmann, Zeit. ges. Nat. **17**. 113.)

+ $10\text{H}_2\text{O}$. Nearly insol. in cold and hot H_2O . Easily sol. in hot acetic acid, dil. H_2SO_4 , HCl and HNO_3 . Insol. in $\text{KOH} + \text{Aq}$, alcohol and ether. ((Zehenter, M. 1900, **21**. 235.))

Potassium hydroxylamine uranate,
 $\text{UO}_4(\text{NH}_4\text{O})(\text{NH}_2\text{OK}) + \text{H}_2\text{O}.$

Sl. sol. in H_2O ; insol. in alcohol. (Hofmann, A. 1899, **307**. 318.)

Rubidium uranate, RbUO_4 .

Insol. in H_2O . (Ditte, A. ch. (6) **1**. 338.)

Silver uranate, $\text{Ag}_2\text{U}_2\text{O}_7$.

Insol. in H_2O . Easily sol. in acids. (Alibegoff, A. **233**. 117.)

Sodium uranate, Na_2UO_4 (?).

Insol. in H_2O ; sol. in dil. acids. Sol. in alkali carbonates + Aq. (Ditte.)

$\text{Na}_2\text{U}_2\text{O}_7 + 6\text{H}_2\text{O}$. Insol. in H_2O . Sol. in dil. acids. (Stolba, Z. anal. **3**. 74.)

Na_2O , 3UO_3 . Insol. in H_2O . Easily sol. in very dil. acids. (Drenckmann.)

Na_2O , $5\text{UO}_3 + 5\text{H}_2\text{O}$. Insol. in H_2O , alcohol, NH_4OH , KOH + Aq. Sol. in HCl , HNO_3 , H_2SO_4 . Sl. sol. even on boiling in conc. acetic acid. (Zehenter, M. 1900, **21**. 235.)

Sodium hydroxylamine uranate,
 $\text{UO}_4(\text{NH}_4\text{O})(\text{NH}_2\text{ONa}) + \text{H}_2\text{O}.$

Sol. in H_2O . (Hofmann, A. 1899, **307**. 319.)

$\text{UO}_4(\text{NH}_2\text{ONa})_2 + 6\text{H}_2\text{O}$. Very sol. in H_2O . (Hofmann.)

Strontium uranate, SrUO_4 .

Insol. in H_2O . Sol. in dil. acids.

SrU_2O_7 . As above. (Ditte, C. R. **95**. 988.)

Very sl. sol. in H_2O . Sol. in all acids especially oxalic.

+ H_2O . Very sl. sol. in H_2O . Sol. in all acids especially oxalic. (J. C. C. **1896**, II. 512.)

Thallous uranate.

Ppt. (Bolton, Am. Chemist, **1872**, **2**. 456.)

Zinc uranate.

Insol. in H_2O ; sol. in $\text{Pb}(\text{C}_2\text{H}_3\text{O}_2)_2$ + Aq. (Persoz, J. pr. **3**. 216.) Sol. in HNO_3 + Aq; insol. in KNO_3 , and NH_4NO_3 + Aq. (Ebelmen, A. ch. (3) **5**. 221.)

Peruranic acid.

See Peruranic acid.

Uranium, U.

Not attacked by H_2O . Slowly decomp. by cold dil. H_2SO_4 + Aq, rapidly on warming. Easily sol. in dil. or conc. HCl + Aq. Fused U is slightly attacked by conc. or fuming HNO_3 , or conc. H_2SO_4 . Amorphous U, however, is easily attacked thereby. Not attacked by acetic acid, KOH , NaOH , or NH_4OH + Aq. (Zimmermann, B. **15**. 849.)

When finely divided, it is decomp. by H_2O slowly at ordinary temps. and rapidly at 100° . (Moissan, C. R. 1896, **122**. 1091.)

Uranium antimonide, U_2Sb_2 .

Violently attacked by conc. HNO_3 . (Colani, C. R. 1903, **137**. 383.)

Uranium arsenide, U_2As_2 .

Violently attacked by conc. HNO_3 . (Colani, C. R. 1903 **137**. 383.)

Uranium boride, UB_2 .

Sol. in HNO_3 and HF . Decomp. by fused alkalis. (Wedekind, B. 1913, **46**. 1204.)

Uranium tribromide, UBr_3 .

Very hygroscopic. Sol. in H_2O with hissing. (Alibegoff, A. **233**. 117.)

Uranium tetrabromide, UBr_4 .

Anhydrous. Very deliquescent. Sol. in H_2O with hissing. (Hermann.)

Insol. in alcohol. (v. Unruh, Dissert. **1900**.)

Sol. in acetone. (Eidmann, C. C. **1899**, II. 1014); methyl acetate (Naumann, B. 1909, **42**. 3790); ethyl acetate. (Naumann, B. 1904, **37**. 3601.)

+ $8\text{H}_2\text{O}$. Very deliquescent, and sol. in H_2O . (Rammelsberg.)

Uranium carbide, UC_2 .

(Ruff and Heinzelmann, Z. anorg. 1911, **71**. 72.)

Attacked slowly by H_2O . Slowly attacked by cold dil. HCl , H_2SO_4 or HNO_3 + Aq. Conc. acids, except HNO_3 , react sl. in the cold, violently on heating. (Moissan, Bull. Soc. 1897, (3) **17**. 12.)

Sol. in fused KNO_3 and KClO_3 ; sol. in dil. acids in the cold and in conc. acids on heating; decomp. by H_2O . (Moissan, C. R. 1896, **122**. 276.)

Uranium trichloride, UCl_3 .

Very sol. in H_2O . (Peligot.)

Very unstable. (Zimmermann.)

Very hygroscopic. Sol. in H_2O with decomp. Sol. in conc. HCl and solution is much more stable than aqueous one. (Rosenheim and Loebel, Z. anorg. 1908, **57**. 234.)

Uranium tetrachloride, UCl_4 .

Anhydrous. Extremely deliquescent.

Sol. in H_2O with evolution of heat. Decomp. on boiling. Sol. in NH_4Cl + Aq without decomp.

HCl increases its solubility in H_2O . (Aloy, Dissert. **1901**.)

Sol. in alcohol, acetone, acetic ether, benzoic ether. Insol. in ether, CHCl_3 and C_6H_6 . (Loebel.)

Sol. in ethyl acetate. (Naumann, B. 1904, **37**. 3601.)

Uranium pentachloride, UCl_5 .

Deliquescent. Sol. in H_2O with evolution of heat and decomposition. (Roscoe, B. **7**. 1131.)

acetic acid, acetic ether, benzaldehyde, benzyl alcohol (trace), aniline (trace), xylidine and p-toluidine (ing).

aniline, ligroin, pyridine, quinoline, ether, thioamyl ether and CS₂. Dissert. 1904.)

abs. alcohol. Insol. in ether, C₆H₆, benzene, ethylene bromide. Sl. sol. in CHCl₃. Sol. in benzoic ether, and trichloroacetic acid. Best solvents are acetate and benzonitrile. Sol. in many organic compounds containing oxygen. Dissert. 1907.)

Hexafluoride, UF₆ + 2H₂O.

Giolitti and Agamennone, C. C. 130.)

Tetrafluoride, UF₄.

1 H₂O. Very sl. sol. in dil. acids. In conc. H₂SO₄, and slowly in warm H₂O + Aq. (Bolton, J. B. 1866. 209.)

Hexafluoride, UF₆.

1. in H₂O. (Ditte, A. ch. (6) 1.

in the air.

hygroscopic; sol. in H₂O. (Ruff, B. 495.)

hygroscopic. Sol. in H₂O. Nearly insol. in paraffine oil. Sol. in all tetrachlorethane (best solvent), CH₂Cl₂, and nitrobenzene. (v. Unruh, 309.)

by H₂O, alcohol and ether. Insol. in CS₂.

CHCl₃, CCl₄, nitrobenzene and (Ruff, Z. anorg. 1911, 72. 81.)

Hydrogen fluoride, UF₆, 8HF (?).

I₂O. (Ditte.)

2, HF + H₂O. (Smithells)

Hydroxide, UO₂, xH₂O.

sol. in dil. acids.

1 alkali hydrates and carbonates (Berzelius.)

alkali carbonates + Aq. (Rammels-

Sol. in dil. acids. (Aloy, Bull. (3) 21. 613.)

Uranic hydroxide, U₃O₈, 6H₂O (?).

sol. in acids.

by (NH₄)₂CO₃ + Aq, which displaces UO₃. (Berzelius.)

Iodide.

Uranic acid.

Tetraiodide, UI₄.

H₂O. (Guichard, C. R. 1907, 145.

Uranium iodide.

Sol. in ethyl acetate.. (Naumann, B. 1904 37. 3601.)

Uranium nitride, U₃N₄.

(Colani, C. R. 1903, 137. 383.)

Uranium suboxide, UO (?).

(Guyard, Bull. Soc. (2) 1. 89.)

Does not exist. (Zimmermann, A. 213. 301.)

U₂O₃(?). Ppt. Decomp. by H₂O and in the air. (Peligot.)

Uranium dioxide (Uranous oxide), UO₂.

Insol. in dil. HCl or H₂SO₄ + Aq.

Sol. in conc. H₂SO₄, and easily in HNO₃ + Aq. (Peligot.)

Insol. in NH₄Cl + Aq. (Rose.)

Only sl. sol. in H₂SO₄, but a considerable amount is converted into the sulphate which is nearly insol. in H₂SO₄.

Slowly sol. in HCl, the amount dissolved in a given time varying widely with the method of preparation of the oxide. (Colani, C. R. 1912, 155. 1251.)

Sl. more sol. in HNO₃ than in aqua regia. (Raynaud, Bull. Soc. 1912, (4) 11. 802.)

Very sol. in conc. HNO₃; less sol. in dil. HNO₃. 1 gram is sol. in 3100 grams HCl (1.17) at 17°; 4650 grams HBr (1.52) at 17°; 2200 grams H₂SO₄ (1.79) at 17°; 12,000 grams acetic acid at 19°. (Raynaud, C. R. 1911, 153. 1481.)

Sl. attacked by liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 830.)

Min. *Uraninite*. Easily sol. in warm HNO₃ + Aq. Not attacked by HCl + Aq.

Uranium trioxide (Uranic oxide), UO₃.

Sol. in HNO₃ + Aq. (Peligot.)

Insol. in boiling K tartrate + Aq. (Kahlenberg and Hillyer, Am. Ch. J. 1894, 16. 102.)

Sol. in oleic acid. (Gibbons, Arch. Pharm. 1883, 221. 621.)

See *Uranic acid*.

Uranium tetroxide, UO₄.

Decomp. by HCl + Aq. (Fairley, Chem. Soc. 31. 133.)

+ 2H₂O. Very hygroscopic. (Zimmermann.)

+ 3H₂O.

Uranium pentoxide, U₂O₅.

Sol. in acids. (Peligot.)

Mixture of UO₃ and U₂O₅. (Rammelsberg, Pogg. 59. 5.)

Mixture of UO₂ and U₂O₅. (Zimmermann, A. 232. 273.)

Uranouranic oxide, U₃O₈.

Green uranium oxide. Very slowly and slightly sol. in dil. HCl or H₂SO₄ + Aq; more easily when conc. Completely sol. in boiling H₂SO₄. Easily sol. in HNO₃ + Aq.

Uranous oxychloride, $\text{UO}_2, \text{UCl}_4 + \text{H}_2\text{O}$.

Moderately sol. in H_2O . (Aloy, Dissert. 1901.)

$2\text{UO}_2, \text{UCl}_4 + \text{H}_2\text{O}$. Very sol. in H_2O and alcohol. (Orloff, C. C. 1903, II. 484.)
 $+13\text{H}_2\text{O}$. Very sol. in H_2O and alcohol. (Orloff.)

$4\text{UO}_2, \text{UCl}_4$. Very sol. in H_2O and alcohol. (Orloff.)

$5\text{UO}_2, \text{UCl}_4 + 10\text{H}_2\text{O}$. Insol. in H_2O . (Aloy, Dissert, 1901.)

Uranous oxyfluoride, $\text{UOF}_2 + 2\text{H}_2\text{O}$.

(Giolitti and Agamennone, C. C. 1905, I. 1130.)

Uranous oxysulphide, $\text{U}_2\text{O}_3\text{S}_4 = \text{UO}_2, 2\text{US}_2$.

Slightly attacked by dil., easily by conc. $\text{HCl} + \text{Aq}$. Sol. in cold $\text{HNO}_3 + \text{Aq}$. (Hermann, J. B. 1861. 258.)

Uranic oxy-compounds.

See Uranyl compounds.

Uranium phosphide, U_3P_2 .

Violently attacked by conc. HNO_3 . (Colani, C. R. 1903, 137. 383.)

U_3P_4 . Slowly attacked by H_2O , not by dil. $\text{HCl} + \text{Aq}$. More easily attacked by conc. $\text{HCl} + \text{Aq}$. Quickly decomp. by boiling conc. HNO_3 and $\text{HNO}_3 + \text{HCl}$. (Colani, A. ch. 1907, (8) 12. 59.)

Uranium selenide, USe .

Spontaneously inflammable. Sol. in fuming HNO_3 . (Colani, C. R. 1903, 137. 383.)

USe_2 . As US_2 . (Colani, C. C. 1903, II. 707.)

U_2Se_3 . (Colani.)

Uranium disilicide, USi_2 .

Sol. in cold or hot conc. HF ; insol. in HCl , HNO_3 , H_2SO_4 and aqua regia. (Defacqz, C. R. 1908, 147. 1051.)

Uranium monosulphide, US .

(Alibegoff, A. 233. 117.)

Uranium sesquisulphide, U_2S_3 .

Not attacked by HCl or dil. $\text{HNO}_3 + \text{Aq}$. Oxidised by fuming H_2SO_4 or aqua regia. (Alibegoff, A. 233. 117.)

Uranium disulphide, US_2 .

Insol. in cold or boiling dil. $\text{HCl} + \text{Aq}$. Sol. in cold conc. $\text{HCl} + \text{Aq}$. Decomp. by $\text{HNO}_3 + \text{Aq}$. (Hermann, J. B. 1861. 258.)

Uranium telluride, U_4Te_3 .

Violently attacked by conc. HNO_3 . (Colani, C. R. 1903, 137. 383.)

Uranosotungstic acid.**Potassium uranosotungstate, $9\text{K}_2\text{O}, 6\text{UO}_2, 8\text{WO}_3 + 34\text{H}_2\text{O}$.**

Insol. in H_2O and in HCl . (Gibbs, Am. Ch. J. 1895, 17. 175.)

Sodium uranosotungstate, $12\text{Na}_2\text{O}, 6\text{UO}_2, 8\text{WO}_3 + 25\text{H}_2\text{O}$.

Insol. in cold H_2O . (Gibbs.)

Uranyl bromide, UO_2Br_2 .

Sol. in H_2O . (de Coninck, C. C. 1903, I. 693.)

Sol. in ether. (v. Unruh, Dissert. 1909.)
 $+7\text{H}_2\text{O}$. Deliquescent. Sol. in H_2O .

Uranyl bromide ammonia, $\text{UO}_2\text{Br}_2, 2\text{NH}_3$.

(v. Unruh, Dissert. 1909.)

$\text{UO}_2\text{Br}_2, 3\text{NH}_3$. (v. Unruh.)

$\text{UO}_2\text{Br}_2, 4\text{NH}_3$. (v. Unruh.)

Uranyl chloride, UO_2Cl_2 .

Anhydrous. Very deliquescent. Sol. in H_2O , alcohol, and ether.

Very sol. in H_2O .

Sp. gr. of $\text{UO}_2\text{Cl}_2 + \text{Aq}$ at t° .

t°	% UO_2Cl_2	Sp. gr.
14.6	1	1.0056
16.3	2	1.0112
13.7	3	1.0161
13.1	4	1.0215
14.2	5	1.0260
15.2	6	1.0313
14.3	7	1.0366
14.5	8	1.0418
15.0	9	1.0469
14.8	10	1.0517

(de Coninck, A. ch. 1904, (8) 3. 500.)

Sol. in conc. HCl , conc. HNO_3 and in selenic acid.

Decomp. by H_2SO_4 . (de Coninck, A. ch. 1904, (8) 3. 504.)

Sol. in methyl acetate (Naumann, B. 1909, 42. 3790); acetone. (Naumann, B. 1904, 37. 4328.)

$+ \text{H}_2\text{O}$. Sol. in H_2O , alcohol, and ether.
 $+ 3\text{H}_2\text{O}$. Deliquescent.

Very sol. in H_2O . 1 pt. is sol. in 0.134 pts. H_2O at 18° and solution, which is saturated, contains 76.2% UO_2Cl_2 or 88.2% $\text{UO}_2\text{Cl}_2 + 3\text{H}_2\text{O}$. Sp. gr. of solution = 2.740. The solubility increases with rise in temp.

Sol. in alcohol and in ether. (Mylius and Dietz, B. 1901, 34. 2775.)

Uranyl hydrogen chloride, $\text{UO}_2\text{Cl}_2, \text{HCl} + 2\text{H}_2\text{O}$.

Cryst. at -10° from sat. solution of UO_2Cl_2 in $\text{HCl} + \text{Aq}$.

Fumes in the air. (Aloy, Bull. Soc. 1901, (3) **25**. 154.)

Uranyl chloride ammonia, $\text{UO}_2(\text{NH}_4\text{Cl})_2$.

Decomp. by H_2O . (Regelsberger, A. **227**. 119.)

$\text{UO}_2(\text{NH}_4\text{NH}_4\text{Cl})\text{NH}_4\text{Cl}$. Decomp. by H_2O . (Regelsberger.)

$\text{UO}_2(\text{NH}_4\text{NH}_4\text{Cl})_2$. Decomp. by H_2O . (Regelsberger.)

Uranyl fluoride, UO_2F_2 .

Very sol. in H_2O . (Smithells, Chem. Soc. **43**. 125.)

Insol. in H_2O or dil. acids. Very sl. sol. in $\text{HF} + \text{Aq}$. Sol. in $\text{H}_2\text{SO}_4 + \text{aqua regia}$. (Ditte, A. ch. (6) **1**. 339.)

Insol. in ether and amyl alcohol. (v. Unruh, Dissert, **1909**.)

UOF_4 . Very sol. in H_2O . (Ditte, C. R. **91**. 115.)

True composition is UO_2F_2 . (Smithells.)

Uranyl hydrogen fluoride, $\text{UO}_2\text{F}_2, \text{HF} + \text{H}_2\text{O}$.

Very sol. in H_2O . (Smithells, Chem. Soc. **43**. 131.)

Uranyl fluoride ammonia, $\text{UO}_2\text{F}_2, 2\text{NH}_3$.

(v. Unruh, Dissert. **1909**.)

$\text{UO}_2\text{F}_2, 3\text{NH}_3$. (v. Unruh.)

$\text{UO}_2\text{F}_2, 4\text{NH}_3$. (v. Unruh.)

Uranyl iodide, UO_2I_2 .

Ppt. Deliquescent. (Aloy, A. ch. 1910, (7) **24**. 417.)

Deliquescent.

Sol. in alcohol, ether and benzene. (Aloy, Dissert, **1901**.)

Sol. in acetone. (Eidmann, C. C. **1899**, II. 1014; Naumann, B. 1904, **37**. 4328); methyl acetate. (Naumann, B. 1909, **42**. 3790.)

Uranyl iodide ammonia, $\text{UO}_2\text{I}_2, 2\text{NH}_3$.

(Aloy, Dissert. **1901**.)

$\text{UO}_2\text{I}_2, 3\text{NH}_3$. (Aloy.)

$\text{UO}_2\text{I}_2, 4\text{NH}_3$. (Aloy.)

Uranyl selenide, UO_2Se .

Very slowly decomp. by H_2O .

Easily sol. in cold HCl . Violently attacked by cold HNO_3 . Not attacked by dil. alkalies. (Milbauer, Z. anorg. 1904, **42**. 450.)

Uranyl sulphide, UO_2S .

Sl. sol. in pure H_2O . Sol. in dil., insol. in absolute alcohol. Sol. in conc. $\text{HCl} + \text{Aq}$, also in dil. acids. Decomp. by caustic alkalies $+ \text{Aq}$. Partly sol. in $(\text{NH}_4)_2\text{S} + \text{Aq}$.

Metavanadic acid, HVO_3 .

Insol. in H_2O ; sol. in acids and alkalies.

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 830.)

$+ \frac{1}{2}\text{H}_2\text{O}$.

See Vanadium pentoxide.

Pyrovanadic acid, $\text{H}_4\text{V}_2\text{O}_7$.

Insol. in H_2O . Sol. in acids and alkalies.

Vanadates.

The alkali, Ba, and Pb metavanadates are sl. sol. in H_2O , the others are more easily sol. Insol. in alcohol.

Aluminum metavanadate.

Very sl. sol. in H_2O . (Berzelius.)

Aluminum divanadate.

Very sl. sol. in H_2O . (Berzelius.)

Ammonium metavanadate, $(\text{NH}_4)\text{VO}_3$.

(a) Very slowly and sparingly sol. in cold H_2O . Easily sol. in hot H_2O . (Berzelius.)

Easily sol. in H_2O at about 70° . Very sl. sol. at above and below that temperature. (Guyard, Bull. Soc. (2) **25**. 355.)

10 g. dissolve in 1 litre cold, and 63 g. in 1 litre hot H_2O with partial decomp. (Ditte, C. R. **102**. 918.)

Solubility in H_2O at t° .

t°	Solubility, mol. per litre
18	0.03715
25	0.05189
35	0.08980
45	0.13406
55	0.17041
70	0.25994

(Meyer, Z. Elektrochem, 1909, **15**. 266.)

Solubility in $\text{NH}_4\text{OH} + \text{Aq}$ at t° .

$\text{NH}_4\text{OH} + \text{Aq}$	t°	Solubility, mol. per litre
0.0677-N	18	0.04763
0.2452-N		0.06798
0.5872-N		0.1029
0.0677-N	25	0.06026
0.2452-N		0.07303
0.5872-N		0.1080

(Meyer, Z. Elektrochem, 1909, **15**. 268.)

Extremely sl. sol. in sat. $\text{NH}_4\text{Cl} + \text{Aq}$. (v. Hauer.)

Insol. in sat. $\text{NH}_4\text{Cl} + \text{Aq}$.

Solubility in salts + Aq at t°.		
Salt solution	t°	Solubility, mol. per litre
0.05-N NH ₄ Cl + Aq	18	0.01419
	25	0.02246
	35	0.04445
	45	0.07575
	55	0.09544
0.1-N NH ₄ Cl + Aq	18	0.00356
	25	0.00995
	35	0.02347
	45	0.04507
	55	0.06314
0.05-N NH ₄ NO ₃ + Aq	18	0.01433
	25	0.02364
0.1-N NH ₄ NO ₃ + Aq	18	0.00497
	25	0.01050

(Meyer, Z. Elektrochem, 1909, 15. 267.)

Insol. in alcohol. (v. Hauer.)

(b) Sol. in cold H₂O, from which it is pptd. by alcohol. (Berzelius.)

Ammonium divanadate, (NH₄)₂V₄O₁₁ + 4H₂O.

Sol. in H₂O, from which it is precipitated by sat. NH₄Cl + Aq or alcohol. (v. Hauer, W. A. B. 21. 337.)

Correct formula is (NH₄)₃V₇O₁₀ + 2H₂O, according to Rammelsberg (B. A. B. 1883. 3.) + 3H₂O. Very sol. in H₂O. (Ditte, C. R. 102. 918.)

Ammonium trivanadate, (NH₄)₃V₆O₁₆.

Anhydrous. Nearly insol. in hot or cold H₂O. (Norblad, B. 8. 126.)

1.5 g. dissolve in 1 litre of boiling H₂O. (Ditte, C. R. 102. 918.)

+ 5H₂O. Very sl. sol. in H₂O. (Ditte.)

+ 6H₂O (?). Very sol. in H₂O. (v. Hauer, W. A. B. 39. 455.)

Could not be obtained. (Norblad; also Rammelsberg, B. A. B. 1883. 3.)

Ammonium vanadate, (NH₄)₃V₇O₁₀ + 2H₂O.

Correct formula of v. Hauer's divanadate, according to Rammelsberg (B. A. B. 1883. 3.)

Sl. sol. in H₂O.

Ammonium sesquivanadate, (NH₄)₄V₆O₁₇ + 4 or 6H₂O.

Very sol. in H₂O. (Ditte, C. R. 102. 918.)

Ammonium pentavanadate, (NH₄)₄V₁₀O₂₇ + 10H₂O.

Sol. in H₂O. (Rammelsberg, B. A. B. 1883. 3.)

Ammonium hydroxylamine vanadate, VO₂N₂H₁₀.

Rapidly decomp. by H₂O. (Hofmann and Kohlschütter, Z. anorg. 1898, 16. 472.)

HVO₃, 3NH₃O, 2NH₃ = VO₂N₂H₁₀. Rapidly decomp. by H₂O. (Hofmann and Kohlschütter.)

Ammonium potassium vanadate, K₂V₄O₁₂ (NH₄)₄V₆O₁₇ + 9H₂O.

Sol. in H₂O. (Ditte, C. R. 104. 1844.)

Ammonium sodium vanadate, Na₂V₄O₁₂ (NH₄)₄V₆O₁₇ + 15H₂O.

Sol. in H₂O. (Ditte, C. R. 104. 1841.)

Ammonium uranyl vanadate, (NH₄)₂O, 2UO₂ V₂O₅ + H₂O.

Insol. in H₂O, NH₄OH + Aq, or dil. HC₂H₃O₂ + Aq. (Carnot, C. R. 104. 1850.)

Barium metavanadate, Ba(VO₃)₂ + H₂O.

Somewhat sol. in H₂O before ignition. Sol. in conc. H₂SO₄. (Berzelius.)

Barium pyrovanadate, Ba₂V₂O₇.

Somewhat sol. in H₂O. (Roscoe.)

Barium vanadate, Ba₂V₆O₁₇ + 14H₂O.

(Ditte, C. R. 104. 1705.)

Ba₂V₁₀O₂₈ + 19H₂O. 1 pt. is sol. in 500 pts. H₂O at 20-25°. Much more sol. in hot, but decomp. by boiling H₂O. (v. Hauer, W. A. B. 21. 344.)

Sol. in about 5000 pts. H₂O. (Manasse, C. C. 1886. 773.)

Ba₄V₁₀(O)₂₉ + 2H₂O. (Norblad.)

Bismuth vanadate, Bi₂(VO₄)₂.

Min. Pucherite. Sol. in HCl + Aq with evolution of Cl.

Cadmium vanadate, Cd(VO₃)₂.

(Ditte, C. R. 102. 918.)

CdV₆(O)₁₈ + 24H₂O. Sl. sol. in H₂O. (Ditte, C. R. 104. 1705.)

Cadmium potassium vanadate, CdK₂V₆O₁₇ + 9H₂O.

(Radau, A. 251. 148.)

Cd₂V₁₀(O)₂₈, K₆V₁₀(O)₂₈ + 27H₂O. 1000 pts. H₂O dissolve 5.4 pts. at 18°. (Radau.)

Cadmium vanadate bromide, 3Cd₂(VO₄)₃ CdBr₂.

Very sol. in dil. acids. (de Schulten, Bull. Soc. 1900, (3) 23. 160.)

Cadmium vanadate chloride, 3Cd₂(VO₄)₃ CdCl₂.

Very sol. in dil. acids. (de Schulten, Bull. Soc. 1900, (3) 23. 159.)

Cæsium metavanadate, CsVO₃.

(Chabrie, A. ch. 1902, (7) 26. 228.)

metavanadate, $\text{Ca}(\text{VO}_3)_2 + 4\text{H}_2\text{O}$.

more sol. than $\text{Sr}(\text{VO}_3)_2$, and solution not precipitated by alcohol. (Ber-

z. *O.* Sol. in H_2O_2 ; insol. in alcohol. (Berz., Z. anorg. 1898, 16. 304.)

pyrovanadate, $\text{Ca}_2\text{V}_2\text{O}_7 + 5\text{H}_2\text{O}$.

precipitate.

O. Very sol. in dil. acids. (Ditte, M. 1705.)

H_2O . (Roscoe.)

divanadate, $\text{CaV}_4\text{O}_{11} + 9\text{H}_2\text{O}$.

sol. in H_2O . (v. Hauer.)

fused is nearly insol. in H_2O . (v.

O. (Manasse, A. 240. 23.)

trivanadate, $\text{CaV}_6\text{O}_{17} + 12\text{H}_2\text{O}$.

sol. in H_2O . (Ditte, C. R. 104. 1705.)

vanadate, $\text{Ca}_3\text{V}_6\text{O}_{23} + 15\text{H}_2\text{O}$.

1 H_2O . (Manasse, A. 204. 23.)

$\text{V}_4\text{O}_{10} + 7\text{H}_2\text{O}$ (?). Sl. sol. in H_2O .

v. a mixture. (Manasse, A. 240. 23.)

$\text{V}_6\text{O}_{23} + 26\text{H}_2\text{O}$. Sol. in H_2O . (Man- 240. 23.)

copper vanadate, $(\text{Ca}, \text{Cu})_4\text{V}_2\text{O}_9 +$

Volborthite. Sol. in $\text{HNO}_3 + \text{Aq}$.

potassium vanadate, $\text{CaK}_3\text{V}_3\text{O}_{11} +$

H_2O . (Manasse, A. 240. 23.)

vanadate chloride, $\text{Ca}_3(\text{VO}_4)_2, \text{CaCl}_2$.
efeuille, C. R. 77. 896.)

m vanadate, CrVO_4 .

itely insol. in H_2O containing H_2O_2 and $\text{HC}_2\text{H}_3\text{O}_2$. (Carnot, C. R. 0.)

is metavanadate, $\text{Co}(\text{VO}_3)_2 + 3\text{H}_2\text{O}$.

sol. in H_2O . (Ditte, C. R. 104.

is potassium vanadate, $\text{CoKV}_5\text{O}_{14} +$

H_2O .
pts. H_2O dissolve 4.8 pts. of this salt. A. 251. 140.)

$\text{V}_{14}\text{O}_{39} + 21\text{H}_2\text{O}$. (Radau.)

metavanadate.

H_2O . (Berzelius.)

pyrovanadate, $\text{Cu}_2\text{V}_2\text{O}_7 + 3\text{H}_2\text{O}$.

hot H_2O . (Ditte, C. R. 104. 1705.)

not be obtained. (Radau, A. 251.

Cupric lead vanadate, $5(\text{Cu}, \text{Pb})\text{O}, \text{V}_2\text{O}_5 + 2\text{H}_2\text{O}$.

Min. *Moltramite*.

$3\text{CuO}, \text{V}_2\text{O}_5, 3(3\text{PbO}, \text{V}_2\text{O}_5), 6\text{CuO}, \text{H}_2\text{O} + 12\text{H}_2\text{O}$. Min. *Psittacinnite*.

Cupric potassium vanadate, $\text{CuKV}_5\text{O}_{14} + 17\text{H}_2\text{O}$.

Moderately sol. in warm H_2O . 100 pts. H_2O dissolve 11.1 pts. at 18° . (Radau, A. 251. 151.)

Didymium vanadate, $\text{Di}_2(\text{VO}_4)_2$.

Precipitate. (Cleve.)

$\text{Di}_2\text{V}_{10}\text{O}_{30} + 28\text{H}_2\text{O}$. Precipitate. (Cleve, Bull. Soc. (2) 43. 365.)

Glucinum metavanadate (?).

Difficultly sol. in H_2O . (Berzelius.)

Glucinum divanadate (?).

Difficultly sol. in H_2O . (Berzelius.)

Indium metavanadate, $\text{In}(\text{VO}_3)_2 + 2\text{H}_2\text{O}$.

Ppt. (Renz, Dissert. 1902.)

Iron (ferrous) metavanadate.

Ppt. Sol. in $\text{HCl} + \text{Aq}$. (Berzelius.)

Iron (ferric) metavanadate.

Somewhat sol. in H_2O . (Berzelius.)

Lead metavanadate, $\text{Pb}(\text{VO}_3)_2$.

Sl. sol. in H_2O . Easily sol. in warm dil. $\text{HNO}_3 + \text{Aq}$. Not completely decomp. by H_2SO_4 or by boiling with $\text{K}_2\text{CO}_3 + \text{Aq}$. (Berzelius.)

Min. *Dechenite*. Easily sol. in dil. $\text{HNO}_3 + \text{Aq}$, and decomp. by $\text{HCl} + \text{Aq}$.

Lead pyrovanadate, basic, $2\text{Pb}_2\text{V}_2\text{O}_7, \text{PbO}$.

Insol. in boiling H_2O or $\text{HC}_2\text{H}_3\text{O}_2$. De-comp. by $\text{HNO}_3 + \text{Aq}$ with separation of V_2O_5 , which dissolves on warming. (Roscoe.)

Lead pyrovanadate, $\text{Pb}_2\text{V}_2\text{O}_7$.

Sol. in warm dil. $\text{HNO}_3 + \text{Aq}$. (Ditte, C. R. 104. 1705.)

Min. *Descloizite*. Sol. in cold dil. $\text{HNO}_3 + \text{Aq}$.

Lead divanadate, $\text{PbV}_4\text{O}_{11}$.

(Ditte, C. R. 104. 1705.)

Lead orthovanadate, $\text{Pb}_2(\text{VO}_4)_2$.

Insol. in H_2O . (Roscoe, A. suppl. 8. 109.)

Lead zinc orthovanadate, $4\text{Pb}_2(\text{VO}_4)_2,$
 $3\text{Zn}_2(\text{VO}_4)_2$.

Min. *Eusynchite*. Easily sol. in $\text{HNO}_3 + \text{Aq}$.

Lead zinc vanadate, $(\text{Pb}, \text{Zn})_4\text{V}_2\text{O}_7 + \text{H}_2\text{O}$.

Min. *Descloizite*. Sol. in excess of $\text{HNO}_3 + \text{Aq}$.

Lead vanadate chloride, $3\text{Pb}_3(\text{VO}_4)_2, \text{PbCl}_2$.

Min. *Vanadinite*. Easily sol. in $\text{HNO}_3 + \text{Aq}$.

Lithium vanadate, basic, $\text{Li}_3\text{V}_2\text{O}_7 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Ditte, C. R. 104. 1168.)
 $\text{Li}_3\text{V}_2\text{O}_7 + \text{H}_2\text{O}$, and $14\text{H}_2\text{O}$. Sol. in H_2O . (Ditte.)

Lithium metavanadate, LiVO_3 .

Easily sol. in H_2O . (Berzelius.)
 $+2\text{H}_2\text{O}$. Quite easily sol. in H_2O . (Rammelsberg, B. A. B. 1883. 3.)

Lithium divanadate, $\text{Li}_2\text{V}_4\text{O}_{11} + 9\text{H}_2\text{O}$.

Very sol. in H_2O . (Norblad.)
 Correct formula is $\text{Li}_2\text{V}_5\text{O}_{14} + 12\text{H}_2\text{O}$. (Rammelsberg.)
 $+8$, or $12\text{H}_2\text{O}$. (Ditte, C. R. 104. 1168.)

Lithium orthovanadate, Li_3VO_4 .

Insol. in H_2O . (Rammelsberg, B. A. B. 1883. 3.)

Lithium pyrovanadate, $\text{Li}_4\text{V}_2\text{O}_7 + 4\text{H}_2\text{O}$.

Very sol. in H_2O . (Rammelsberg, B. 16. 1676.)
 $+3\text{H}_2\text{O}$. (Ditte, C. R. 104. 1168.)

Lithium vanadate, $\text{Li}_3\text{V}_5\text{O}_{14} + 7\text{H}_2\text{O}$.

Difficultly sol. in H_2O . (Rammelsberg.)
 $+12\text{H}_2\text{O}$. Very efflorescent. Correct formula for v. Hauer's divanadate. (Rammelsberg.)

$\text{Li}_4\text{V}_6\text{O}_{17} + 16\text{H}_2\text{O}$. Sol. in H_2O . (Ditte, C. R. 104. 1168.)

$+15\text{H}_2\text{O}$. (Rammelsberg.)

$+11\text{H}_2\text{O}$. (R.)

$+3\text{H}_2\text{O}$. (R.)

$\text{Li}_5\text{V}_6\text{O}_{18} + 15\text{H}_2\text{O}$. Not very easily sol. in H_2O . (Rammelsberg.)

$\text{Li}_5\text{V}_5\text{O}_{17} + 12\text{H}_2\text{O}$. Moderately sol. in H_2O . (Rammelsberg.)

$\text{Li}_{10}\text{V}_{12}\text{O}_{36} + 30\text{H}_2\text{O}$. Efflorescent. Very sol. in H_2O . (Rammelsberg.)

Magnesium metavanadate, $\text{Mg}(\text{VO}_3)_2$.

Very easily sol. in H_2O . (Berzelius.)
 $+6\text{H}_2\text{O}$. Very sol. in H_2O . (Ditte, C. R. 104. 1705.)

Magnesium divanadate, $\text{MgV}_4\text{O}_{11} + 8\text{H}_2\text{O}$.

Difficultly sol. in H_2O , but much more sol. than barium divanadate. (v. Hauer.)
 $+9\text{H}_2\text{O}$. (Ditte, C. R. 104. 1705.)

Magnesium trivanadate, $\text{Mg}_3\text{V}_6\text{O}_{17} + 4\frac{1}{2}\text{H}_2\text{O}$.

Very sl. sol. in H_2O . (Manasse, A. 240. 23.)

Magnesium vanadate, $\text{Mg}_3\text{V}_6\text{O}_{17} + 28\text{H}_2\text{O}$.

Sol. in H_2O . (Suguira and Baker, C. Soc. 35. 715.)

Manganous metavanadate, $\text{Mn}(\text{VO}_3)_2 + 4\text{H}_2\text{O}$.

Very sl. sol. in cold, somewhat more : hot H_2O . Easily sol. in dil. acids. (Rad. 251. 125.)

Sl. sol. in H_2O_2 ; insol. in alcohol. (Sci. Z. anorg. 1898, 16. 304.)

Manganous pyrovanadate, $\text{Mn}_2\text{V}_2\text{O}_7$.

Sl. sol. in hot dil. $\text{HNO}_3 + \text{Aq}$. (Ditte, 96. 1048.)

Manganous potassium vanadate, $\text{MnK} + 8\text{H}_2\text{O}$.

100 pts. H_2O dissolve 1.7 pts. salt a. Easily sol. in acids. (Radau, A. 251.

$3\text{Mn}_2\text{V}_2\text{O}_7, \text{K}_2\text{V}_2\text{O}_7 + 54\text{H}_2\text{O}$. (R.)

$7\text{Mn}(\text{VO}_3)_2, 2\text{KVO}_3 + 25\text{H}_2\text{O}$. (R.)

$11\text{Mn}(\text{VO}_3)_2, 2\text{KVO}_3 + 48\text{H}_2\text{O}$. (R.)

Mercuric vanadate.

Sl. sol. in H_2O .

Nickel vanadate, $\text{Ni}(\text{VO}_3)_2$.

Sol. in H_2O . (Ditte, C. R. 104. 1

Nickel orthovanadate, $\text{Ni}_2(\text{VO}_4)_3$.

Insol. in H_2O ; sol. in $\text{HNO}_3 + \text{Aq}$. (C. R. 96. 1049.)

Nickel divanadate, $\text{NiV}_4\text{O}_{11} + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Ditte, C. R. 104. 1705.

Nickel potassium vanadate, $5\text{Ni}(\text{VO}_3)_2 + 2\text{KVO}_3 + 25\text{H}_2\text{O}$.

$\text{Ni}_2\text{K}_2\text{V}_{10}\text{O}_{30} + 17\text{H}_2\text{O}$. Very sl. sol. in H_2O .

$\text{NiKV}_5\text{O}_{14} + 8\text{H}_2\text{O}$.

$2\text{Ni}_4\text{V}_{14}\text{O}_{38}, \text{K}_2\text{V}_{14}\text{O}_{38} + 69\text{H}_2\text{O}$. 1000 H_2O dissolve 1.7 pts. of salt at 17.5° . (R. A. 251. 137.)

Potassium vanadate, basic, $\text{K}_3\text{V}_2\text{O}_7 + 2\text{H}_2\text{O}$.

Sol. in H_2O . (Ditte, C. R. 104. 902.

Potassium metavanadate, KVO_3 .

Anhydrous. Slowly sol. in cold, easily in hot H_2O . Insol. in alcohol. (Berzelius.)

Completely sol. in a little cold H_2O . (Norblad.)

$+ \text{H}_2\text{O}$. Sol. in H_2O . (Rammelsberg.)

$+ 1\frac{1}{4}\text{H}_2\text{O}$. (Ditte.)

$+ 1\frac{1}{2}\text{H}_2\text{O}$. (Ditte.)

$+ 2\text{H}_2\text{O}$. (Ditte.)

$+ 3\text{H}_2\text{O}$. (Ditte, C. R. 104. 902.)

$+ 7\text{H}_2\text{O}$. (Rammelsberg.)

Potassium divanadate, $K_2V_2O_{11} + 4H_2O$.

Sol. in cold or lukewarm H_2O . Decomp. by hot H_2O . (Rammelsberg.)

+3 H_2O . (Berzelius.)

+3½ H_2O . Sol. in warm H_2O . (Norblad.)

+8 or 10 H_2O . (Ditte, C. R. 104. 902.)

+6 H_2O . (Ephraim, Z. anorg. 1903, 35. 76.)

Potassium trivanadate, $K_3V_3O_{18}$.

Anhydrous. Nearly insol. in H_2O . (Norblad.)

+6 H_2O . Insol. in cold or hot H_2O . (Norblad.)

+1, and 5 H_2O . (Ditte, C. R. 104. 902.)

Potassium orthovanadate, $K_3VO_4 + 4\frac{1}{2}$ or 6 H_2O .

Deliquescent. Sol. in H_2O . (Ditte, C. R. 104. 902.)

Decomp. by H_2O into $K_4V_2O_7$ and KOH. Rammelsberg, B. A. B. 1883. 3.)

Potassium pyrovanadate, $K_4V_2O_7 + 3H_2O$.

Deliquescent. Easily sol. in H_2O . Insol. in alcohol. (Norblad.)

+4 H_2O . (Ditte, C. R. 104. 902.)

Potassium vanadate, $K_3V_3O_{14} + 5H_2O$.

100 pts. H_2O dissolve 19.2 pts. at 17.5°. Radau, A. 251. 120.)

+4½ H_2O . (Radau.)

$K_4V_2O_7 + 2H_2O$. Slowly sol. in H_2O . Rammelsberg.)

+6 H_2O . (Ditte, C. R. 104. 902.)

+7 H_2O . (Friedheim, B. 23. 1526.)

$K_4V_{10}O_{27} + 12H_2O$. Very sol. in H_2O . Manasse, A. 240. 42.)

$K_{10}V_3O_{36} + 7H_2O$. Sol. in H_2O . (Rammelsberg.)

$K_3V_3O_{11} + 1\frac{1}{2}H_2O$. Very sl. sol. in H_2O . Ephraim, Z. anorg. 1903, 35. 75.)

$K_4V_{18}O_{47}$. (Ephraim, Z. anorg. 1903, 5. 78.)

Potassium sodium vanadate, $2(2K_2O, 3V_2O_5), 3(2Na_2O, 3V_2O_5) + 30H_2O$.

(Friedheim, Z. anorg. 1894, 5. 442.)

$2K_2O, 3V_2O_5, 4(2Na_2O, 3V_2O_5) + 35H_2O$. Efflorescent. (Friedheim, Z. anorg. 1894, 5. 441.)

Potassium strontium vanadate, $K_2Sr_3V_{14}O_{39} + 20H_2O$.

Sol. in H_2O . (Manasse, A. 240. 23.)

$K_2Sr_3V_{14}O_{39} + 30H_2O$. As above. (Manasse.)

$K_4Sr_3V_{14}O_{39} + 18H_2O$. As above. (Manasse.)

Potassium zinc vanadate, $KZnV_6O_{14} + 8H_2O$.

1000 pts. H_2O dissolve 4.1 pts of the salt. Radau, A. 251. 145.)

$2K_3V_{14}O_{39}, 3Zn_4V_{14}O_{39} + 90H_2O$. (Radau.)

Potassium vanadate cyanide, $K_4V_2O_7, 4KCN + 14H_2O$.

Easily decomp.

Insol. in alcohol. (Petersen, Z. anorg. 1904, 38. 343.)

Samarium vanadate, $Sm_2O_3, 5V_2O_5 + 28H_2O$.

(Cleve.)

+24 H_2O . (Cleve.)

Samarium orthovanadate.

Precipitate.

Silver metavanadate, $AgVO_3$.

Sol. in HNO_3 or dil. $NH_4OH + Aq$. (Berzelius.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, 20. 829.)

Silver orthovanadate, Ag_3VO_4 .

Ppt. Easily sol. in HNO_3 or $NH_4OH + Aq$. (Roscoe, Proc. Roy. Soc. 18. 316.)

Silver pyrovanadate, $Ag_4V_2O_7$.

Ppt. (Roscoe.)

Sol. in $NH_4OH + Aq$. (Ditte, C. R. 104. 1705.)

Silver vanadate, $Ag_3V_4O_{18}$.

Sol. in 21,414 pts. H_2O at 14°, and 13,617 pts. at 100°. (Carnelley, A. 166. 155.)

Silver vanadate ammonia, $6AgVO_3, 4NH_3 + 8H_2O$.

(Ditte, C. R. 104. 1705.)

Sodium vanadate, basic, $Na_3V_2O_7 + 26$ or $30H_2O$.

Very sol. in H_2O . (Ditte.)

Sodium metavanadate, $NaVO_3$.

Anhydrous. Slowly sol. in cold, very easily in hot H_2O . (Norblad.)

100 g. H_2O dissolve at:

25° 40° 60° 75°

21.10 26.23 32.97 38.83 g. $NaVO_3$.

(McAdam and Pierle, J. Am. Chem. Soc. 1912, 34. 606.)

+2 H_2O . Easily sol. in H_2O .

100 g. H_2O dissolve at:

25° 40° 60°

15.23 29.93 68.36 g. $NaVO_3$.

At 75° a value was obtained which showed that the solid phase had changed into the less sol. modification. (McAdam and Pierle, J. Am. Chem. Soc. 1912, 34. 607.)

+½ H_2O . (Ditte, C. R. 104. 1061.)

+3, 4, and 5 H_2O . (Ditte.)

Sodium divanadate, $Na_2V_2O_{11}$.

Anhydrous. Sl. sol. even in warm H_2O , but easily sol. on addition of acids.

+9H₂O. Easily sol. in cold H₂O. Insol. in alcohol. (Norblad.)
 +5H₂O. (Ditte, C. R. 104. 1061.)
 Not obtained by Rammelsberg (B. A. B. 1883. 3.)

Sodium trivanadate, Na₃V₃O₁₀+9H₂O.

Insol. in cold or hot H₂O. (Norblad.)
 Composition is Na₃V₁₀O₄₃+24H₂O. (Rammelsberg.)
 +3H₂O. (Ditte, C. R. 104. 1061.)

Sodium orthovanadate, Na₃VO₄+16H₂O.

Easily sol. in H₂O, but decomp. into Na₄V₂O₇ and KOH. Precipitated by an excess of alcohol. (Roscoe, A. suppl. 8. 102.)
 +7H₂O. (Hall, Chem. Soc. 51. 96.)
 +10., and 12H₂O. Less sol. in dil. NaOH + Aq than in H₂O. (Baker, A. 229. 286.)

Sodium pyrovanadate, Na₄V₂O₇+18H₂O.

Easily sol. in H₂O. Insol. in alcohol. (Norblad.)
 Sol. in alcohol. (Ditte, C. R. 104. 1061.)
 +8H₂O. (Ditte.)

Sodium sesquivanadate, Na₄V₃O₁₇.

Anhydrous. Insol. in H₂O or NH₄OH + Aq. (Rammelsberg.)
 +10H₂O. (Norblad.)
 +16H₂O. Efflorescent. (Rammelsberg.)
 +18H₂O. (Ditte.)

Sodium pentavanadate, Na₄V₁₀O₃₇+3½H₂O.
 Scarcely sol. in H₂O. (Rammelsberg.)

Sodium vanadate, Na₄V₄O₁₈+6H₂O.

Difficultly sol. in cold H₂O. (Carnelley, A. 166. 155.)
 +2H₂O. (Carnelley.)
 Na₄V₁₀O₄₃+24H₂O. Correct formula for Norblad's trivanadate. (Rammelsberg.)
 Na₂O, 4V₂O₅+7½H₂O. (Baragiola, Dissert. 1902.)
 +8½H₂O. (Baragiola.)
 3Na₂O, 5V₂O₅+22H₂O. (Prandtl and Lustig, Z. anorg. 1907, 53. 405.)
 4Na₂O, 7V₂O₅+33H₂O. (Friedheim, Z. anorg. 1894, 5. 443.)
 5Na₂O, 8V₂O₅+39H₂O. Sol. in H₂O. (Friedheim, Z. anorg. 1894, 5. 441.)

Sodium vanadate fluoride, 2Na₃VO₄, NaF+19H₂O.

Sol. in H₂O. (Rammelsberg, W. Ann. 20. 928.)

Strontium metavanadate, Sr(VO₃)₂+4H₂O.

Difficultly sol. in cold H₂O. (Norblad.)

Strontium divanadate, SrV₂O₁₁+9H₂O.

Sl. sol. in H₂O, but much more sol. than barium divanadate. (v. Hauer.)

Sol. in H₂O₂+Aq free from H₂SO₄. Insol. in alcohol. (Scheuer, Z. anorg. 1896, 16. 303.)

Strontium trivanadate, SrV₃O₁₄+14H₂O.

Sol. in H₂O, but decomposes slowly on boiling. Easily sol. in hot H₂O acidified with HC₂H₃O₂, and crystallizes therefrom without decomp. (v. Hauer, J. pr. 78. 156.)

Strontium tetravanadate, SrV₄O₂₁+11H₂O.

Sol. in hot H₂O with partial decomposition (Manasse, A. 240. 34.)

Strontium vanadate, Sr₂V₂O₂₂+14H₂O.

Sol. in H₂O. (Manasse, A. 240. 23.)
 Sr₂V₁₄O₂₈+30H₂O. Sol. in H₂O. (Norblad.)

Thallous metavanadate, TlVO₃.

Sol. in 11,534 pts. H₂O at 11°, and 473 pts. at 100°. (Carnelley.)

Thallous orthovanadate, Tl₃VO₄.

Sl. sol. in H₂O. Sol. in 999 pts. H₂O at 15°, and 574 p.s. at 100°. (Carnelley, Chem Soc. (2) 11. 323.)

Thallous pyrovanadate, Tl₄V₂O₇.

Sol. in 4996 pts. H₂O at 14°, and 3840 pts. H₂O at 100°. (Carnelley.)

Thallous vanadate, Tl₁₂V₃O₂₈.

Sol. in 3406 pts. H₂O at 14°, and 533 pts. at 100°. (Carnelley.)

Tl₁₂V₁₀O₃₁. Sol. in 9372 pts. H₂O at 11°, and 3366 pts. at 100°. (Carnelley.)

Tl₁₂V₁₄O₄₁. Ppt. (Carnelley.)

Thorium vanadate, Th₂O₁₂(VO)₄, 16V₂O₅+24H₂O (?).

Sol. in H₂O. (Cleve.)
 ThO₂, V₂O₅+6H₂O. Sol. in acids. (Volck, Z. anorg. 1894 6. 167.)

Uranyl vanadate, 2UO₃, V₂O₅, (CO₂)₂V₂O₇.

Insol. in H₂O. (Carnot, C. R. 104. 1850.)

Vanadium vanadate, 2VO₂, V₂O₅=V₂O₅.

Insol. in H₂O. Sol. in dil. H₂SO₄ or HNO₃ + Aq. (Rammelsberg.)

Slowly oxidised by HNO₃+Aq. Slowly sol. in NH₄OH+Aq. Easily sol. in HCl+Aq (Ditte, C. R. 101. 1487.)

+2½H₂O. (Brierley.)

2VO₂, 2V₂O₅+8H₂O. Insol. in H₂O (Brierley, Chem. Soc. 49. 31.)

Ytterbium vanadate, 3Yb₂O₃, 5V₂O₅+3H₂O.

Yb₂O₃, 15V₂O₅. Ppt. (Cleve, Z. anorg. 1902, 32. 150.)

Yttrium vanadate.

Precipitate. (Berzelius.)

vanadate, $\text{Zn}(\text{VO}_3)_2 + 2\text{H}_2\text{O}$.
insol. in H_2O . (Ditte, C. R. 104. 1705.)

pyrovanadate, $\text{Zn}_2\text{V}_2\text{O}_7$.
slightly sol. in H_2O . (Ditte, C. R. 108.)

vanadic acid.
Pervanadic acid.

metavanadic acid.

ammonium vanadicotungstate,
 $(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 8\text{WO}_3 + 10\text{H}_2\text{O}$.
sol. in H_2O .
insol. in organic solvents. (E. F. Smith,
Chem. Soc. 1903, 25. 1227.)

ammoniovanadic acid.

ammonium vanadicovanadate, $(\text{NH}_4)_2\text{O},$
 $\text{V}_2\text{O}_5, 4\text{V}_2\text{O}_5 + 8\text{H}_2\text{O}$.
sol. in cold and warm H_2O . (Gibbs,
ibid. 7. 209.)
 $(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 2\text{V}_2\text{O}_5 + 14\text{H}_2\text{O}$. Sol. in
(Brierley, Chem. Soc. 49. 30.)
 $(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 4\text{V}_2\text{O}_5 + 6\text{H}_2\text{O}$. Insol.
(Brierley.)

ammonium —, $2\text{K}_2\text{O}, 2\text{V}_2\text{O}_5, \text{V}_2\text{O}_5 + 6\text{H}_2\text{O}$.
insol. in hot H_2O . (Brierley, Chem. Soc.
ibid. 49. 30.)
 $(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 4\text{V}_2\text{O}_5 + \text{H}_2\text{O}$. Insol. in H_2O .
(Brierley.)
 $(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 8\text{V}_2\text{O}_5$. Insol. in H_2O .
insol. in conc. H_2SO_4 . (Prandtl, B. 1905, 38.)

ammonium —, $2\text{Na}_2\text{O}, 2\text{V}_2\text{O}_5, \text{V}_2\text{O}_5 + 13\text{H}_2\text{O}$.
slightly sol. in H_2O . Insol. in conc. solu-
tion of salts, especially acetate. (Brierley,
Chem. Soc. 49. 30.)
 $(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 5\text{V}_2\text{O}_5$. Insol. in H_2O . Sol.
in conc. H_2SO_4 . Not attacked by boiling
conc. HNO_3 . Slowly attacked by hot
 $\text{H}_2\text{O} + \text{Aq}$. (Prandtl, B. 1905, 38.)

diarsenic acid.
arseniovanadic acid.

iododic acid.
iodovanadic acid.

ammoniomolybdic acid, $8\text{MoO}_3, \text{V}_2\text{O}_5 +$
 H_2O .
sl. sol. in H_2O , and sl. sol. in boiling
+ Aq . (Ditte, C. R. 102. 757.)
could not be obtained. (Friedheim, B.
1903.)

ammonium vanadiomolybdate,
 $(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 2\text{MoO}_3 + 4\text{H}_2\text{O}$.
slightly insol. in H_2O . (Friedheim and
Castendyck, B. 1900, 33. 1615.)

$(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 2\text{MoO}_3 + 8\text{H}_2\text{O}$. Nearly
insol. in H_2O . (Friedheim and Castendyck.)
 $2(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 3\text{MoO}_3 + 6\text{H}_2\text{O}$. (Euler-
Chelpin, Dissert, 1895.)

$(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 3\text{MoO}_3 + 17\text{H}_2\text{O}$. (Euler-
Chelpin.)
 $2(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 4\text{MoO}_3 + 7\text{H}_2\text{O}$ and +
 $8\text{H}_2\text{O}$. (Euler-Chelpin.)
 $3(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 4\text{MoO}_3 + 7\text{H}_2\text{O}$. (Milch,
Dissert. Berlin, 1887.)
+ $9\text{H}_2\text{O}$. Sol. in H_2O . (Ditte, C. R. 102.
1019.)

+ $11\text{H}_2\text{O}$. Easily sol. in H_2O . Correct
composition of above compounds is =
 $(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5 + 2[(\text{NH}_4)_2\text{O}, 2\text{MoO}_3] + 11\text{H}_2\text{O}$.
(Friedheim, B. 24. 1173.)

Moderately sol. in H_2O and can be re-
cryst. therefrom. (Euler-Chelpin, Dissert,
1895.)

$2(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, 4\text{MoO}_3 + 11\text{H}_2\text{O}$. Near-
ly insol. in H_2O . (Friedheim and Casten-
dyck, B. 1900, 33. 1615.)

$2(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 5\text{MoO}_3$. Nearly insol.
in cold H_2O . (Euler-Chelpin, Dissert. 1895.)
+ $8\text{H}_2\text{O}$. Nearly insol. in H_2O . (Fried-
heim and Castendyck, B. 1900, 33. 1615.)

$3(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 5\text{MoO}_3 + 8\frac{1}{2}\text{H}_2\text{O}$. Very
easily sol. in H_2O . (Liebert, Dissert. 1891.)
 $4(\text{NH}_4)_2\text{O}, 12\text{V}_2\text{O}_5, 5\text{MoO}_3 + 24\text{H}_2\text{O}$. Prac-
tically insol. in H_2O . (Friedheim and Casten-
dyck, B. 1900, 33. 1615.)

$2(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 6\text{MoO}_3 + 5\text{H}_2\text{O}$. Sol.
in a large amount of H_2O . (Gibbs, Am. Ch.
J. 5. 361.)

+ $6\text{H}_2\text{O}$. Rather sl. sol. in H_2O .
Easily sol. in acids. (Liebert, Dissert,
1891.)

Composition is double the above formula,
or—

$4(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 12\text{MoO}_3 + 12\text{H}_2\text{O}$.
Rather difficultly sol. in H_2O . Composition
is $(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5 + 3[(\text{NH}_4)_2\text{O}, 4\text{MoO}_3]$.
(Friedheim)

$3(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 6\text{MoO}_3 + 7\text{H}_2\text{O}$. (Isen-
burg, Dissert, 1901.)

$5(\text{NH}_4)_2\text{O}, 4\text{V}_2\text{O}_5, 6\text{MoO}_3 + 12\text{H}_2\text{O}$ and
+ $14\text{H}_2\text{O}$. (Euler-Chelpin, Dissert, 1895.)

$5(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, 7\text{MoO}_3 + 13\text{H}_2\text{O}$ and
+ $16\text{H}_2\text{O}$. Sol. in H_2O and can be recryst.
therefrom. (Toggenburg, Dissert, 1902.)

$5(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, 8\text{MoO}_3 + 14\text{H}_2\text{O}$.
(Stamm, Dissert, 1905.)

$5(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 12\text{MoO}_3 + 10\text{H}_2\text{O}$. Quite
easily sol. in H_2O . Composition is $(\text{NH}_4)_2\text{O},$
 $2\text{V}_2\text{O}_5 + 4[(\text{NH}_4)_2\text{O}, 3\text{MoO}_3] + 10\text{H}_2\text{O}$.

$6(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, 12\text{MoO}_3 + 21\text{H}_2\text{O}$. Sol.
in H_2O . Composition is $(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5 +$
 $5(\text{NH}_4)_2\text{O}, 12\text{MoO}_3$. (F.)

$8(\text{NH}_4)_2\text{O}, 4\text{V}_2\text{O}_5, 13\text{MoO}_3 + 21\text{H}_2\text{O}$. Sol.
in H_2O . (Isenburg, Dissert, 1901.)

$8(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 18\text{MoO}_3 + 15\text{H}_2\text{O}$. De-
comp. by hot H_2O . (Gibbs.) Could not
be obtained. (Friedheim.)

$10(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, 24\text{MoO}_3 + 10\text{H}_2\text{O}$. Sol.
in H_2O . (Milch.) Could not be obtained.
(Friedheim.)

Ammonium barium vanadiomolybdate,
 $5(\text{NH}_4)_2\text{O}, 15\text{BaO}, 6\text{V}_2\text{O}_5, 36\text{MoO}_3.$

(Milch.)

$(\text{NH}_4)_2\text{O}, \text{BaO}, \text{V}_2\text{O}_5, \text{MoO}_3 + 6\text{H}_2\text{O}.$

(Euler-Chelpin, Dissert, 1895.)

$3(\text{NH}_4)_2\text{O}, \text{BaO}, 3\text{V}_2\text{O}_5, 5\text{MoO}_3 + 9\text{H}_2\text{O}.$

Sl. sol. in cold and hot H_2O . (Euler-Chelpin.)

Ammonium potassium —, $(\text{NH}_4)_2\text{O}, 3\text{K}_2\text{O}, 2\text{V}_2\text{O}_5, 4\text{MoO}_3 + 5\text{H}_2\text{O}.$

Decomp. by H_2O . (Euler-Chelpin,)

$(\text{NH}_4)_2\text{O}, 3\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 5\text{MoO}_3 + 9\text{H}_2\text{O}.$

(Euler-Chelpin.)

$\frac{1}{2}(\text{NH}_4)_2\text{O}, 3\frac{1}{2}\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 5\text{MoO}_3 + 16\text{H}_2\text{O}.$ Sol. in H_2O . (Jacoby, Dissert. 1900.)

Ammonium sodium —, $(\text{NH}_4)_2\text{O}, \text{Na}_2\text{O}, 2\text{V}_2\text{O}_5, 6\text{MoO}_3 + 12\text{H}_2\text{O}.$

(Euler-Chelpin, Dissert, 1895.)

Barium —, $\text{BaO}, \text{V}_2\text{O}_5, \text{MoO}_3 + 7\text{H}_2\text{O}.$

(Euler-Chelpin.)

Barium —, $3\text{BaO}, 2\text{V}_2\text{O}_5, 6\text{MoO}_3.$

(Milch.)

$+ 8\text{H}_2\text{O}.$ (Euler-Chelpin.)

$5\text{BaO}, 4\text{V}_2\text{O}_5, 6\text{MoO}_3 + 28\text{H}_2\text{O}.$ Sol. in hot H_2O . (Euler-Chelpin.)

$3\text{BaO}, \text{V}_2\text{O}_5, 8\text{MoO}_3 + 2\text{BaO}, \text{H}_2\text{O}, \text{V}_2\text{O}_5, 8\text{MoO}_3 + 28\text{H}_2\text{O}.$ Sol. in hot H_2O . (Gibbs, Am. Ch. J. 5. 361.)

$7\text{BaO}, 3\text{V}_2\text{O}_5, 18\text{MoO}_3 + 16\text{H}_2\text{O} = \text{BaO}, 3\text{V}_2\text{O}_5 + 6(\text{BaO}, 3\text{MoO}_3) + 16\text{H}_2\text{O}.$ Sl. sol. in $\text{H}_2\text{O}.$

$+ 36\text{H}_2\text{O}$ and $+ 48\text{H}_2\text{O}.$ (Liebert, Dissert. 1891.)

Potassium —, $\text{K}_2\text{O}, \text{V}_2\text{O}_5, 3\text{MoO}_3 + 15\text{H}_2\text{O}.$

Nearly insol. in cold H_2O . (Euler-Chelpin, Dissert, 1895.)

$3\text{K}_2\text{O}, 2\text{V}_2\text{O}_5, 4\text{MoO}_3 + 8\text{H}_2\text{O} = \text{K}_2\text{O}, 2\text{V}_2\text{O}_5 + 2(\text{K}_2\text{O}, 2\text{MoO}_3) + 8\text{H}_2\text{O}.$

Very sol. in H_2O . (Friedheim.)

$+ 7\text{H}_2\text{O}.$ Easily sol. in cold H_2O . (Euler-Chelpin.)

$+ 9\text{H}_2\text{O}.$ Easily sol. in cold H_2O . (Euler-Chelpin.)

$2\text{K}_2\text{O}, 2\text{V}_2\text{O}_5, 5\text{MoO}_3 + 10\text{H}_2\text{O}.$ Nearly insol. in cold, very sl. sol. in hot H_2O . (Euler-Chelpin.)

$3\text{K}_2\text{O}, \text{V}_2\text{O}_5, 6\text{MoO}_3 + 5\text{H}_2\text{O}.$ (Euler-Chelpin.)

$2\text{K}_2\text{O}, \text{V}_2\text{O}_5, 6\text{MoO}_3 + 6\text{H}_2\text{O}.$ Very sl. sol. in cold, easily sol. in hot H_2O . (Liebert.)

$4\text{K}_2\text{O}, 2\text{V}_2\text{O}_5, 12\text{MoO}_3 + 12\text{H}_2\text{O} = \text{K}_2\text{O}, 2\text{V}_2\text{O}_5 + 3(\text{K}_2\text{O}, 4\text{MoO}_3) + 12\text{H}_2\text{O}.$ Sl. sol. in H_2O . (Friedheim.)

$5\text{K}_2\text{O}, 2\text{V}_2\text{O}_5, 12\text{MoO}_3 + 12\text{H}_2\text{O} = \text{K}_2\text{O}, 2\text{V}_2\text{O}_5 + 4(\text{K}_2\text{O}, 3\text{MoO}_3) + 12\text{H}_2\text{O}.$ Rather sl. sol. in H_2O . (Friedheim.)

Sl. sol. in cold, easily sol. in hot H_2O . (Liebert, Dissert. 1891.)

$3\text{K}_2\text{O}, \text{V}_2\text{O}_5, 12\text{MoO}_3 + 15\text{H}_2\text{O}.$ (Liebert.)

Potassium sodium vanadiomolybdate, $\text{K}_2\text{O}, 4\text{Na}_2\text{O}, 2\text{V}_2\text{O}_5, 12\text{MoO}_3 + 18\text{H}_2\text{O}.$

(Euler-Chelpin, Dissert 1895.)

Sodium —, $2\text{Na}_2\text{O}, \text{V}_2\text{O}_5, 6\text{MoO}_3 + 16\text{H}_2\text{O}.$

(Euler-Chelpin, Dissert, 1895.)

Vanadiophosphoric acid.

See Phosphovanadic acid.

Vanadiouselenious acid, $3\text{V}_2\text{O}_5, 4\text{SeO}_2 + 4\text{H}_2\text{O}.$

$+ 6\text{H}_2\text{O}.$ Difficultly sol. in H_2O . Can be cryst. from $\text{H}_2\text{O}.$

$+ 10\text{H}_2\text{O}.$ Difficultly sol. in H_2O . Can be cryst. from H_2O . (Prandtl, B. 1905, 38. 1307.)

Ammonium vanadiouselenite,

$4(\text{NH}_4)_2\text{O}, 6\text{V}_2\text{O}_5, 5\text{SeO}_2 + 13\text{H}_2\text{O}.$

Sl. sol. in H_2O . Decomp. by boiling H_2O . $3(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{SeO}_2 + 2\text{H}_2\text{O}.$ Ppt. (Prandtl, B. 1905, 38. 1309.)

$(\text{NH}_4)_2\text{HV}_2\text{O}_{17}, 12\text{SeO}_2 + 2\text{H}_2\text{O}.$ Ppt. $+ 4\text{H}_2\text{O}.$ Ppt. (Prandtl, Z. anorg. 1911, 73. 231.)

Ammonium silver —, $2\frac{2}{3}(\text{NH}_4)_2\text{O}, 1\frac{1}{3}\text{Ag}_2\text{O}, 6\text{V}_2\text{O}_5, 5\text{SeO}_2 + 12\text{H}_2\text{O}, + 16\text{H}_2\text{O}$ and $+ 22\text{H}_2\text{O}.$

(Prandtl, Z. anorg. 1907, 53. 402.)

Lithium —, $4\text{Li}_2\text{O}, 6\text{V}_2\text{O}_5, 5\text{SeO}_2 + 30\text{H}_2\text{O}.$

Very sol. in H_2O . (Prandtl.)

Potassium —, $4\text{K}_2\text{O}, 6\text{V}_2\text{O}_5, 5\text{SeO}_2 + 13\text{H}_2\text{O}.$

$3\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{SeO}_2.$ (Prandtl, B. 1905, 38. 1309.)

Sodium —, $4\text{Na}_2\text{O}, 6\text{V}_2\text{O}_5, 5\text{SeO}_2 + 20\text{H}_2\text{O}.$

Very. sol. in H_2O . (Prandtl, Z. anorg. 1907, 53. 403.)

Vanadiousulphuric acid, $\text{V}_2\text{O}_5, 3\text{SO}_3 + 3\text{H}_2\text{O}.$

Deliquescent. Sol. in H_2O , but is decomp. by boiling. (Ditte, C. R. 102. 757.)

See Sulphate, vanadium.

Vanadiousulphurous acid.

Ammonium vanadiousulphite,

$3(\text{V}_2\text{O}_5, \text{SO}_2), (\text{NH}_4)_2\text{O}, \text{SO}_2 + 4\text{H}_2\text{O}.$

(Gain, C. R. 1907, 144. 1158.)

Cæsium —, $(\text{V}_2\text{O}_5, \text{SO}_2), 3(\text{Cs}_2\text{O}, \text{SO}_2) + 8\text{H}_2\text{O}.$

(Gain.)

Lithium —, $(\text{V}_2\text{O}_5, \text{SO}_2), 5(\text{Li}_2\text{O}, \text{SO}_2) + 8\text{H}_2\text{O}.$

(Gain.)

Potassium vanadiousulphite, $3\text{SO}_2, (\text{V}_2\text{O}_4), \text{SO}_2, (\text{K}_2\text{O}) + 5\text{H}_2\text{O}$.

(Gain.)

Rubidium —, $(\text{V}_2\text{O}_4, \text{SO}_2), 2(\text{Rb}_2\text{O}, \text{SO}_2) + 2\text{H}_2\text{O}$.

(Gain.)

Sodium —, $5(\text{V}_2\text{O}_4, \text{SO}_2), (\text{Na}_2\text{O}, \text{SO}_2) + 2\text{H}_2\text{O}$.

(Gain.)

Thallium —, $3(\text{V}_2\text{O}_4, \text{SO}_2), (\text{Tl}_2\text{O}, \text{SO}_2) + 8\text{H}_2\text{O}$.

(Gain.)

Vanadiotungstic acid, $6\text{H}_2\text{O}, \text{V}_2\text{O}_5, 10\text{WO}_3 + 16\text{H}_2\text{O}$.

Very sl. sol. in cold, more easily in hot H_2O . (Gibbs, Am. Ch. J. 5. 361.)

$6\text{H}_2\text{O}, \text{V}_2\text{O}_5, 16\text{WO}_3 + 30\text{H}_2\text{O}$. Readily sol. in H_2O . (Gibbs.)

$17\text{H}_2\text{O}, 4\text{V}_2\text{O}_5, 16\text{WO}_3 + 24\text{H}_2\text{O}$. Sl. sol. in cold, easily in hot H_2O . (Rosenheim, A. 251. 228.)

Aluminum sodium vanadiotungstate, $7\text{Al}_2\text{O}_3, 27\text{Na}_2\text{O}, 36\text{V}_2\text{O}_5, 144\text{WO}_3 + 504\text{H}_2\text{O} = 3(\text{Al}_2\text{O}_3, 9\text{Na}_2\text{O}, 48\text{WO}_3), 4(\text{Al}_2\text{O}_3, 9\text{V}_2\text{O}_5) + 504\text{H}_2\text{O}$.

Sol. in H_2O . (Rothenbach, B. 23. 3055.)

Ammonium —, $(\text{NH}_4)_2\text{O}, 3\text{V}_2\text{O}_5, \text{WO}_3 + 6\text{H}_2\text{O}$.

Sol. in H_2O . (Rammelsberg, B. 1. 158.) $2(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 4\text{WO}_3 + 4\text{H}_2\text{O}$. (Friedheim and Löwy, Z. anorg. 1894, 6. 24.)

$4(\text{NH}_4)_2\text{O}, 2\text{H}_2\text{O}, \text{V}_2\text{O}_5, 5\text{WO}_3 + 11\text{H}_2\text{O}$. Sol. in H_2O . (Gibbs, Am. Ch. J. 5. 361.)

$2(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_5, 5\text{WO}_3 + 10\text{H}_2\text{O}$. Sol. in H_2O . (Ditte, C. R. 102. 1019.)

$31(\text{NH}_4)_2\text{O}, 14\text{V}_2\text{O}_5, 60\text{WO}_3 + 58\text{H}_2\text{O} = 5[5(\text{NH}_4)_2\text{O}, 12\text{WO}_3], 2[3(\text{NH}_4)_2\text{O}, 7\text{V}_2\text{O}_5]$. Sol. in H_2O . (Rothenbach, B. 23. 3051.)

$7(\text{NH}_4)_2\text{O}, 4\text{V}_2\text{O}_5, 14\text{WO}_3 + 16\text{H}_2\text{O}$. Sol. in H_2O . (Rosenheim, A. 251. 197.)

$8(\text{NH}_4)_2\text{O}, 4\text{V}_2\text{O}_5, 16\text{WO}_3, 9\text{H}_2\text{O} + 4\text{H}_2\text{O}$. Efflorescent. Very sol. in H_2O . (Rosenheim, A. 251. 216.)

Barium —, $19\text{BaO}, 10\text{V}_2\text{O}_5, 36\text{WO}_3 + 94\text{H}_2\text{O} = 3(5\text{BaO}, 12\text{WO}_3), 2(2\text{BaO}, 5\text{V}_2\text{O}_5) + 94\text{H}_2\text{O}$.

Sl. sol. in H_2O . (Rothenbach, B. 23. 3052.) $8\text{BaO}, 4\text{V}_2\text{O}_5, 16\text{WO}_3, 9\text{H}_2\text{O} + 44\text{H}_2\text{O}$.

Efflorescent. Not very sol. in H_2O . (Rosenheim, A. 251. 218.)

Composition is $6\text{BaO}, 12\text{WO}_3, 3\text{V}_2\text{O}_5 + 39\text{H}_2\text{O}$. (Friedheim.)

$6\text{BaO}, 3\text{V}_2\text{O}_5, 12\text{WO}_3 + 34\text{H}_2\text{O}$. Not easily sol. in H_2O . (Friedheim and Löwy, Z. anorg. 1894, 6. 18.)

$4\text{BaO}, 4\text{V}_2\text{O}_5, 12\text{WO}_3 + 41\text{H}_2\text{O}$. Less sol.

than preceding salt. Decomp. by boiling or by mineral acids. (Rosenheim.)

Composition is $4\text{BaO}, 12\text{WO}_3, 3\text{V}_2\text{O}_5 + 30\text{H}_2\text{O}$. (Friedheim.)

Calcium vanadiotungstate, $2\text{CaO}, \text{V}_2\text{O}_5, 2\text{WO}_3 + 12\text{H}_2\text{O}$.

(Friedheim and Löwy, Z. anorg. 1894, 6. 20.)

Magnesium sodium —, $\text{MgO}, 6\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5, 12\text{WO}_3 + 42\text{H}_2\text{O} = 5\text{Na}_2\text{O}, 12\text{WO}_3 + \text{MgO}, \text{Na}_2\text{O}, 3\text{V}_2\text{O}_5 + 42\text{H}_2\text{O}$.

Sol. in H_2O . (Rothenbach, B. 23. 3054.)

Potassium —, $4\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 12\text{WO}_3 + 30\text{H}_2\text{O}$.

Sol. in H_2O .

Composition is potassium *metatungstate* vanadate, $3(\text{K}_2\text{O}, 4\text{WO}_3) + \text{K}_2\text{O}, 3\text{V}_2\text{O}_5 + 30\text{H}_2\text{O}$. (Friedheim, B. 23. 1515.)

$8\text{K}_2\text{O}, 4\text{V}_2\text{O}_5, 16\text{WO}_3, 9\text{H}_2\text{O} + 24\text{H}_2\text{O}$. Very efflorescent. Easily sol. in H_2O . (Rosenheim, A. 251. 214.)

Formula is $6\text{K}_2\text{O}, 12\text{WO}_3, 3\text{V}_2\text{O}_5 + 24\text{H}_2\text{O}$, which is a double salt, $5\text{K}_2\text{O}, 12\text{WO}_3 + \text{K}_2\text{O}, 3\text{V}_2\text{O}_5$. (Friedheim, B. 23. 1505.)

Potassium sodium —, $(5\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 22\text{H}_2\text{O}), 4(5\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 22\text{H}_2\text{O})$ or $\text{Na}_2\text{O}, 4\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 22\text{H}_2\text{O}$.

(Friedheim and Löwy, Z. anorg. 1894, 6. 22.)

$4(5\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 24\text{H}_2\text{O}), 3(5\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 24\text{H}_2\text{O})$. (Friedheim and Löwy.)

$5(5\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 24\text{H}_2\text{O}), 5\text{K}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 24\text{H}_2\text{O}$. (Friedheim and Löwy.)

Silver —, $8\text{Ag}_2\text{O}, 4\text{V}_2\text{O}_5, 16\text{WO}_3, 9\text{H}_2\text{O}$.

Somewhat sol. in cold H_2O , more easily upon addition of little HNO_3 . Decomp. by warm H_2O . (Rosenheim, A. 251. 224.)

$3\text{Ag}_2\text{O}, 2\text{V}_2\text{O}_5, 6\text{WO}_3 + 3\text{H}_2\text{O}$. Nearly insol. in cold H_2O . Decomp. by addition of HNO_3 or upon warming. (Rosenheim.)

Sodium —, $5\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5, 6\text{WO}_3 + 36\text{H}_2\text{O}$.

Sol. in H_2O .

Composition is $3(\text{Na}_2\text{O}, 2\text{WO}_3) + 2(\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5) + 36\text{H}_2\text{O}$. (Friedheim, B. 23. 1527.)

$+ 38\text{H}_2\text{O}$. Sol. in 1.25 pts. H_2O at 13.8° .

(Friedheim and Löwy.) $2\text{Na}_2\text{O}, 2\text{V}_2\text{O}_5, 3\text{WO}_3 + 20\text{H}_2\text{O}$. Very sol. in H_2O .

Composition is $\text{Na}_2\text{O}, 3\text{WO}_3 + \text{Na}_2\text{O}, 2\text{V}_2\text{O}_5 + 20\text{H}_2\text{O}$, double salt of sodium *tritungstate* and *divanadate*. (Friedheim, B. 23. 1523.)

$4\text{Na}_2\text{O}, 3\text{V}_2\text{O}_5, 12\text{WO}_3 + 38\text{H}_2\text{O} = 3(\text{Na}_2\text{O}, 4\text{WO}_3) + \text{Na}_2\text{O}, 3\text{V}_2\text{O}_5 + 38\text{H}_2\text{O}$. Sol. in H_2O . (Rothenbach, B. 23. 3050.)

$8\text{Na}_2\text{O}, 4\text{V}_2\text{O}_5, 16\text{WO}_3, 9\text{H}_2\text{O} + 48\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O . (Rosenheim, A. 251. 210.)

Formula is $6Na_2O, 12WO_3, 3V_2O_5 + 42H_2O$, and is a double salt of sodium *paratungstate*, $5Na_2O, 12WO_3$, and sodium *trivanadate*, $Na_2O, 3V_2O_5$. (Friedheim, B. 23. 1505.)

$7Na_2O, V_2O_5, 12WO_3 + 29H_2O$. Easily sol. in H_2O . (Friedheim and Löwy, Z. anorg. 1894, 6. 15.)

$8Na_2O, V_2O_5, 14WO_3 + 60H_2O$ and $+66H_2O$ (Friedheim and Löwy.)

Strontium vanadiotungstate, $19SrO, 36WO_3, 10V_2O_5 + 122H_2O = 3(5SrO, 12WO_3, 2(2SrO, 5V_2O_5) + 122H_2O$.

Sol. in H_2O . (Rothenbach, B. 23. 3053.)

Vanadious acid.

See Hypovanadic acid.

Vanadiovanadicomolybdic acid.

Ammonium vanadiovanadicomolybdate, $11(NH_4)_2O, 4V_2O_5, VO_3, 28MoO_3 + 48H_2O$.

Sl. sol. in cold, sol. in hot H_2O without decomp. (Gibbs, Am. Ch. J. 5. 391.)

Barium —, $14BaO, 2V_2O_5, 3VO_3, 30MoO_3 + 48H_2O$.

Precipitate. Very sl. sol. in cold, decomp. by hot H_2O . (Gibbs.)

Vanadiovanadicotungstic acid.

Ammonium vanadiovanadicotungstate, $6(NH_4)_2O, 2V_2O_5, 3VO_3, 12WO_3 + 12H_2O$.

Sol. in H_2O . (Gibbs, Am. Ch. J. 5. 391.)

Silver —, $6Ag_2O, 2V_2O_5, 3VO_3, 12WO_3 + 8H_2O$.

Precipitate. Very sl. sol. in cold, sol. in much warm H_2O . (Gibbs.)

Sodium —, $6Na_2O, 2V_2O_5, 3VO_3, 12WO_3$. Very sol. in H_2O . (Gibbs.)

Vanadium, V.

Insol. in H_2O , HCl , dil. $H_2SO_4 + Aq$, and cold conc. H_2SO_4 . Sol. in hot conc. H_2SO_4 . Slowly sol. in $HF + Aq$. Easily sol. in dil. or conc. $HNO_3 + Aq$. Not attacked by hot or cold $NaOH$ or $KOH + Aq$. (Roscoe, A. suppl. 7. 85.)

Does not alter in the air; not acted upon by HCl and very slowly by H_2SO_4 . (Moissan, C. R. 1896, 122. 1299.)

Vanadium boride, VB.

Sol. in HF and HNO_3 . Decomp. by fused *alkalies*. (Wedekind, B. 1913, 46. 1203.)

Vanadium tribromide, VBr_3 .

Very deliquescent; quickly decomposes (Roscoe, A. suppl. 8. 99.)

$+6H_2O$. Sol. in H_2O with decomp. (Locke and Edwards, Am. Ch. J. 1898, 20. 600.)

Sol. in H_2O , alcohol and ether. Insol. in HBr . (Piccini, Z. anorg. 1899, 19. 395.)

Vanadium carbide, VC.

Insol. in HCl and H_2SO_4 .

Sol. in HNO_3 in the cold, and in fused KNO_3 and $KClO_3$. (Moissan, C. R. 1896, 122. 1300.)

Vanadium dichloride, VCl_2 .

Very deliquescent. Sol. in H_2O , alcohol, and ether. (Roscoe, A. suppl. 7. 79.)

Vanadium trichloride, VCl_3 .

Deliquescent. Sol. in H_2O , absolute alcohol, and ether.

$+6H_2O$. Sol. in H_2O ; sl. sol. in conc. $HCl + Aq$.

Sol. in alcohol and ether. (Piccini, Z. anorg. 1899, 19. 395.)

Vanadium tetrachloride, VCl_4 .

Sol. with decomp. in H_2O , alcohol, and ether. (Roscoe.)

Sol. in H_2O with decomp.

Sol. in fuming HCl with decomp.

Sol. in anhydrous $CHCl_3$ or glacial acetic acid apparently without chemical change (Koppel, Z. anorg. 1905, 45. 346.)

Vanadium difluoride, VF_2 .

Sol. in HF with evolution of H_2 and formation of VF_3 . (Manhot, A. 1907, 357. 135.)

Vanadium trifluoride, VF_3 .

Nearly insol. in H_2O and organic solvents. (Ruff, B. 1911, 44. 2544.)

$+3H_2O$. Efflorescent. Easily sol. in cold, extremely sol. in hot H_2O with decomp. Can be recryst. from $HF + Aq$. Insol. in strong alcohol. (Petersen, J. pr. (2) 40. 48.)

Vanadium tetrafluoride, VF_4 .

Very hygroscopic.

Easily sol. in H_2O .

Difficultly sol. in SO_2Cl_2 and $SiCl_4$.

Sol. in $POCl_3$ with evolution of gas.

Sol. in acetone and acetic acid.

Difficultly sol. in alcohol and $CHCl_3$. (Ruff, B. 1911, 44. 2545.)

Vanadium pentafluoride, VF_5 .

Easily sol. in H_2O .

Easily sol. in alcohol, $CHCl_3$, acetone, and ligroin. Insol. in CS_2 . Decomp. toluene and ether. (Ruff, B. 1911, 44. 2549.)

Vanadium trifluoride with MF .

See Fluovanadate, VI.

Vanadium hydride,

Stable. Does not react with boiling H_2O .
Not attacked by boiling HCl , but oxidized by hot H_2SO_4 .

Sol. in boiling HNO_3 . (Muthmann, A. 1907, **355**. 86.)

Vanadium dihydroxide, $\text{VO}, x\text{H}_2\text{O}$.

Insol. in KOH or $\text{NaOH} + \text{Aq}$.

$\text{V}(\text{OH})_2$. Sol. in HCl . (Locke and Edwards, Am. Ch. J. 1898, **20**. 598.)

Vanadium trihydroxide, $\text{V}_2\text{O}_3, x\text{H}_2\text{O}$.

Easily sol. in all acids. (Petersen, J. pr. (2) **40**. 49.)

Vanadium tetrahydroxide (Hypovanadic hydroxide), $\text{V}_2\text{O}_5(\text{OH})_4 + 5\text{H}_2\text{O}$.

Easily sol. in acids or alkalies. (Crow, Chem. Soc. **30**. 453.)

Vanadium triiodide, $\text{VI}_3 + 6\text{H}_2\text{O}$.

Very hygroscopic; sol. in H_2O . Only sl. sol. in cold conc. $\text{HI} + \text{Aq}$.

Sol. in alcohol. (Piccini, Z. anorg. 1899, **19**. 399.)

Vanadium nitride, V_2N .

Insol. in H_2O , HCl , and $\text{KOH} + \text{Aq}$. Sol. in $\text{HNO}_3 + \text{H}_2\text{SO}_4$.

Decomp. by fusing with KOH . (Muthmann, A. 1907, **355**. 93.)

Insol. in liquid NH_3 . (Gore, Am. Ch. J. 1898, **20**. 830.)

VN . (Roscoe, A. suppl. **6**. 114.)

VN_2 . Not attacked by cold $\text{HNO}_3 + \text{Aq}$. (Uhrlaub, Pogg. **103**. 134.)

Vanadium dioxide, VO .

Insol. in H_2O ; easily sol. in dil. acids. (Roscoe, A. suppl. **6**. 95.)

Vanadium trioxide, V_2O_3 .

Oxidized in H_2O in contact with air and then dissolves. Insol. in acids, except HNO_3 , and in alkalies + Aq . (Roscoe, A. suppl. **6**. 99.)

Easily sol. in HF . (Petersen, J. pr. (2) **40**. 48.)

Vanadium tetroxide, VO_2 .

Sol. in acids and alkalies + Aq .

Vanadium pentoxide, V_2O_5 .

Sol. in about 1000 pts. H_2O . (Berzelius.)

Sol. in acids, alkali hydrates, and carbonates + Aq . Insol. in absolute, very sl. sol. in dil. alcohol.

Insol. in glacial $\text{HC}_2\text{H}_3\text{O}_2$.

Sol. in conc. $\text{KF} + \text{Aq}$. (Ditte, C. R. **105**. 1067.)

Sol. in $\text{H}_2\text{C}_2\text{O}_4 + \text{Aq}$ and alkali oxalates + Aq . (Halberstadt, Z. anal. **22**. 1.)

Three modifications.—(a) Forms hydrates

with 2, and $5\text{H}_2\text{O}$. Sol. in H_2O . 1 l. of sat. solution contains 8 g. V_2O_5 .

(β) $\text{V}_2\text{O}_5, 2\text{H}_2\text{O}$. Very sl. sol. in H_2O . 1 l. of sat. solution contains 0.5 g. V_2O_5 .

(γ) $\text{V}_2\text{O}_5, 5\text{H}_2\text{O}$. Less sol. in H_2O than β 1 l. H_2O contains 0.05 g. V_2O_5 when saturated. (Ditte, C. R. **101**. 698.)

See Vanadic acid.

Vanadium oxide, $\text{V}_4\text{O}_7 = 2\text{VO}_2, \text{V}_2\text{O}_5$.

See Vanadate, vanadium.

$\text{V}_2\text{O}_4, \text{V}_2\text{O}_5 + \frac{1}{2}\text{H}_2\text{O}$. (Brierley, Chem. Soc. **49**. 30.)

See also Vanadovanadic acid.

$\text{V}_2\text{O}_4, 2\text{V}_2\text{O}_5 + 8\text{H}_2\text{O}$.

See Vanadate, vanadium.

Vanadium pentoxide with MF.

See Fluoxyvanadate, M.

Vanadium oxy compounds.

See Vanadyl compounds.

Vanadium silicide, V_2Si .

Insol. in H_2O .

Not attacked by HCl , HNO_3 or H_2SO_4 . Readily attacked by HF .

Not attacked by $\text{KOH} + \text{Aq}$, $\text{NaOH} + \text{Aq}$ or NH_4OH . Decomp. by fused KOH .

Insol. in alcohol, ether and benzene. (Moissan, C. R. 1902, **135**. 496.)

VSi_2 . Sol. in HF ; insol. in acids and alkalies.

Decomp. by fused KOH . (Moissan, C. R. 1902, **135**. 78.)

Vanadium disulphide, V_2S_3 .

Insol. in boiling dil. or conc. HCl , dil. $\text{H}_2\text{SO}_4 + \text{Aq}$, or cold conc. H_2SO_4 . Easily sol. in hot dil. or conc. $\text{HNO}_3 + \text{Aq}$, or in boiling conc. H_2SO_4 . Insol. in alkalies + Aq . Sl. sol. in $\text{KSH} + \text{Aq}$; sol. in $\text{NH}_4\text{SH} + \text{Aq}$. (Kay, Chem. Soc. **37**. 728.)

Vanadium trisulphide, V_2S_5 .

Insol. in cold HCl or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. Very sl. sol. in hot HCl or dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. More sol. in $\text{HNO}_3 + \text{Aq}$ or conc. H_2SO_4 . Sl. sol. in NaOH or $\text{NH}_4\text{OH} + \text{Aq}$. Easily sol. in $(\text{NH}_4)_2\text{S}$ or $\text{NH}_4\text{SH} + \text{Aq}$, also in $\text{K}_2\text{S} + \text{Aq}$. (Kay, Chem. Soc. **37**. 728.)

Vanadium pentasulphide, V_2S_8 .

Sl. attacked by hot conc. HCl or hot dil. $\text{H}_2\text{SO}_4 + \text{Aq}$; sol. in hot conc. H_2SO_4 . Sol. in hot dil. $\text{HNO}_3 + \text{Aq}$. Sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$, but easily dissolved by $\text{NaOH} + \text{Aq}$. Sl. sol. in $\text{Na}_2\text{S} + \text{Aq}$. Sol. in $\text{NH}_4\text{SH} + \text{Aq}$. (Kay.)

Vanadium sulphochloride sulphur chloride, $4\text{VSCl}_3, \text{S}_2\text{Cl}_2$.

Decomp. in the air. (Koppel, Z. anorg. 1905, **45**. 357.)

Vanadous acid.**Ammonium vanadite, basic, $2(\text{NH}_4)_2\text{O}, \text{V}_2\text{O}_4$.**

Sl. sol. in cold, easily in hot H_2O . (Ditte, C. R. 102. 1310.)

Ammonium vanadite, $(\text{NH}_4)_2\text{V}_4\text{O}_9 + 3\text{H}_2\text{O}$.

Sol. in H_2O . (Crow, Chem. Soc. 30. 460.)
 $+x\text{H}_2\text{O}$. Sol. in H_2O .

Insol. in alcohol, ether and ammonia. (Koppel, Z. anorg. 1903, 36. 297.)

$+3\text{H}_2\text{O}$. Easily sol. in H_2O . (Mawrow, Z. anorg. 1907, 55. 150.)

Barium vanadite, $\text{BaV}_4\text{O}_9 + 4\text{H}_2\text{O}$.

Ppt. (Koppell, Z. anorg. 1903, 36. 300.)
 $+5\text{H}_2\text{O}$. Precipitate. Easily sol. in HNO_3 ,
 or $\text{HCl} + \text{Aq}$. (Crow, Chem. Soc. 30. 460.)

Lead vanadite, PbV_4O_9 .

Ppt. (Crow.)

Potassium vanadite, $\text{K}_2\text{V}_4\text{O}_9 + 4\text{H}_2\text{O}$.

Sol. in H_2O .

Insol. in alcohol, ether and ammonia (Koppel, Z. anorg. 1903, 36. 300.)

$+7\text{H}_2\text{O}$. Easily sol. in H_2O . Insol. in cold, sol. in hot $\text{KOH} + \text{Aq}$. Insol. in alcohol. (Crow.)

$+ \text{H}_2\text{O}$. (Ditte, C. R. 102. 1310.)

Silver vanadite, $\text{Ag}_2\text{V}_4\text{O}_9$.

Ppt. (Crow.)

Sodium vanadite, $\text{Na}_2\text{V}_4\text{O}_9 + 4\text{H}_2\text{O}$.

Sol. in H_2O .

Insol. in alcohol, ether and ammonia. (Koppel, Z. anorg. 1903, 36. 299.)

$+7\text{H}_2\text{O}$. Easily sol. in H_2O . (Crow, Chem. Soc. 30. 459.)

Vanadosotungstic acid.**Ammonium vanadosotungstate, $5(\text{NH}_4)_2\text{O}, 2\text{V}_2\text{O}_5, 14\text{WO}_3 + 13\text{H}_2\text{O}$.**

Very sol. in H_2O . (E. F. Smith, J. Am. Chem. Soc. 1903, 25. 1228.)

Vanadous acid.

See Hypovanadic acid.

Vanadovanadic acid.

See Vanadicovanadic acid.

Vanadyl bromide, VOBr .

Very sl. sol. in H_2O , acetic anhydride, ethyl acetate, and acetone.

Insol. in alcohol, ether, acetic acid, CHCl_3 , toluene and CCl_4 . (Ruff, B. 1911, 44. 2537.)

Vanadyl dibromide, VOBr_2 .

Very deliquescent, and sol. in H_2O . (Roscoe.)

Vanadyl tribromide, VOBr_3 .

Very deliquescent, and quickly decomposes in moist air. Sol. in H_2O . (Roscoe.)

Vanadyl bromide, $\text{V}_2\text{O}_5\text{Br}_2, 2\text{HBr} + 7\text{H}_2\text{O}$.

Very deliquescent. (Ditte, C. R. 102. 1310.)

Vanadyl semichloride, $\text{V}_2\text{O}_5\text{Cl}$.

Insol. in H_2O . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Roscoe, A. suppl. 6. 114.)

Vanadyl monochloride, VOCl .

Insol. in H_2O . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Roscoe.)

Vanadyl dichloride, VOCl_2 .

Deliquescent. Slowly decomp. by H_2O . Easily sol. in $\text{HNO}_3 + \text{Aq}$. (Roscoe.)

Vanadyl trichloride, VOCl_3 .

Deliquescent. Sol. in H_2O and alcohol with decomp. (Bedson, A. 180. 235.)

Sol. in ether with combination.

Divanadyl chloride, $\text{V}_2\text{O}_5\text{Cl}_2 + 5\text{H}_2\text{O}$.

Deliquescent, and sol. in H_2O , fuming HCl , or alcohol. (Crow, Chem. Soc. 30. 457.)

Vanadyl chloride, $\text{V}_2\text{O}_5\text{Cl}_2 + 4\text{H}_2\text{O}$.

Very deliquescent. (Ditte, C. R. 102. 1310.)

Vanadyl platinum chloride.

See Chloroplatinate, vanadyl.

Vanadyl trichloride ammonia, $\text{VOCl}_3, x\text{NH}_3$.

Decomp. by H_2O . (Roscoe.)

Vanadyl difluoride, VOF_2 .

Insol. in H_2O . Insol. in alcohol, ether, CHCl_3 . Sl. sol. in acetone. (Ruff, B. 1911, 44. 2546.)

Vanadyl trifluoride, VOF_3 .

Very hygroscopic. Easily sol. in H_2O . Sol. in POCl_3 with evolution of gas. Difficultly sol. in PCl_3 and AsCl_3 . Sol. in hot CHCl_3 and acetic anhydride. (Ruff, B. 1911, 44. 2547.)

Vanadyl fluoride with MF.

See Fluoxyvanadate, and Fluoxhypovanadate, M.

Vanadyl iodide, $\text{V}_2\text{O}_5\text{I}_2, 3\text{HI} + 10\text{H}_2\text{O}$.

Very deliquescent, and sol. in H_2O . (Ditte, C. R. 102. 1310.)

$\text{V}_2\text{O}_5\text{I}_2, 2\text{HI} + 8\text{H}_2\text{O}$. As above.

Vanadyl sulphide, VOS (?) .

(a) Insol. in H_2O , alkalies, alkali sulphides $+ \text{Aq}$. Sol. in acids, except nitric acid and aqua regia. (Berzelius.)

(b) Sol. in alkalies, alkali carbonates, and sulphides + Aq. Insol. in acids. (Berzelius.)

Water, H₂O.

Water is the most universal solvent. It absorbs all gases, usually with an increase of volume, seldom, as in the case of NH₃, with a diminution of volume. It dissolves almost all solids in greater or less quantity, and mixes with or dissolves considerable amounts of many liquids.

Miscible with alcohol. Sol. in 36 pts. ether.

Sol. in 30–33 vols. ethyl acetate. (Becker.)

Sol. in 5 vols. iodhydrin.

Sl. sol. in most of the fatty oils.

Solubility in organic solvents at t°.

Solvent	t°	G. H ₂ O in 100 g. of the solution
Benzene	+3	0.030
	23	0.060
	40	0.114
	55	0.184
	66	0.255
	77	0.337
Petroleum bpt. 190–250° at atmos. pressure	–2	0.0012
	+18	0.005
	23	0.007
	30	0.008
	36	0.012
	53	0.026
	59	0.031
	61	0.035
	66	0.043
	79	0.063
	85	0.075
	94	0.097
Paraffin oil bpt. 200–300° at 10 mm. pressur	+16	0.003
	50	0.013
	65	0.022
	73	0.030
	77	0.035
	94	0.055

(Groschuff, Z. Elektrochem, 1911, 17. 350.)

White precipitate, fusible.

See Mercuridiammonium chloride.

White precipitate, infusible.

See Mercuric chloramide.

Xanthochromium bromide,



Sol. in H₂O. Resembles the chloride. (Christensen, J. pr. (2) 24. 74.)

— carbonate, Cr(NO₂)(NH₃)₅CO₃.

Easily sol. in H₂O. (Christensen.)

Xanthochromium chloride,



More sol. in H₂O than the roseo, but less than the purpureo salt.

Solution decomp. by light or boiling. Decomp. by dil. acids. Sol. in NaOH + Aq and in NH₄OH + Aq (sp. gr. 0.91). Insol. in alcohol. (Christensen, J. pr. (2) 24. 74.)

— chloroplatinate, Cr(NO₂)(NH₃)₅PtCl₆.

Insol. in pure H₂O, but sol. when warmed with H₂O containing HCl, with formation of a new double salt. (Christensen.)

— mercuric chloride, Cr(NO₂)(NH₃)₅Cl₂, 2HgCl₂.

Precipitate. Decomp. by long contact with H₂O. (Christensen.)

— chromate, Cr(NO₂)(NH₃)₅CrO₄.

Difficultly sol. in H₂O. (Christensen.)

— dichromate, Cr(NO₂)(NH₃)₅Cr₂O₇.

Difficultly sol. in H₂O. (Christensen.)

— dithionate, Cr(NO₂)(NH₃)₅S₂O₈.

Insol. in cold H₂O. (Christensen.)

— hydroxide, Cr(NO₂)(NH₃)₅(OH)₂.

Known only in solution. (Christensen.)

— iodide, Cr(NO₂)(NH₃)₅I₂.

Quite difficultly sol. in H₂O. (Christensen.)

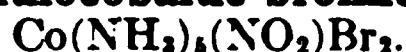
— nitrate, Cr(NO₂)(NH₃)₅(NO₃)₂.

Sol. in about 150 pts. H₂O. (Christensen.)

— sulphate, Cr(NO₂)(NH₃)₅SO₄ + H₂O.

Sol. in H₂O and (NH₄)₂SO₄ + Aq. (Christensen.)

Xanthocobaltic bromide,



Easily sol. in cold H₂O. (Werner and Miolati, Gazz. ch. it. 23, 2. 140.)

— bromonitrate, Co(NO₂)(NH₃)₅(NO₃)Br.

Sl. sol. in cold, more easily in hot H₂O. (Gibbs.)

— chloride, Co(NO₂)(NH₃)₅Cl₂.

Sl. sol. in cold H₂O, and decomp. by boiling therewith. Insol. in HCl + Aq and alkali chlorides + Aq. Easily decomp. by boiling with acids, even dilute. (Gibbs and Genth.)

Sol. in 50 pts. cold H₂O. (Jørgensen, Z. anorg. 5. 172.)

— mercuric chloride, Co(NO₂)(NH₃)₅Cl₂, 2HgCl₂ + H₂O.

Insol. in cold, sl. sol. in hot H₂O without decomp. More sol. in acidified H₂O. (Gibbs and Genth.)

Xanthocobaltic chloraurate,

Can be easily crystallised out of hot H_2O . (Gibbs and Genth, Sill. Am. J. (2) **24**. 90.)

— **chloronitrate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5(\text{NO}_2)\text{Cl}$.
Sl. sol. in cold, more easily in hot H_2O .

— **chloronitrate gold chloride**,
 $\text{Co}(\text{NO}_2)(\text{NH}_3)_5(\text{NO}_2)\text{Cl}$, AuCl_3 .

— **chloronitrate platonic chloride**,
 $2\text{Co}(\text{NO}_2)(\text{NH}_3)_5(\text{NO}_2)\text{Cl}$, PtCl_4 .

— **chloroplatinate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{Cl}_3$,
 $\text{PtCl}_4 + \text{H}_2\text{O}$.

Scarcely sol. in hot or cold H_2O . Can be recryst. from dil. $\text{HNO}_3 + \text{Aq}$. Sol. in hot dil. $\text{HCl} + \text{Aq}$. (Gibbs and Genth, Sill. Am. J. (2) **24**. 91.)

— **chromate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{CrO}_4 + \text{H}_2\text{O}$.
Very sl. sol. in cold, and but slightly sol. in hot H_2O . (Gibbs.)

— **dichromate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{Cr}_2\text{O}_7$.
Easily sol. in hot H_2O .

— **ferrocyanide**, $[\text{Co}(\text{NO}_2)(\text{NH}_3)_5]_2\text{Fe}(\text{CN})_6 + 7\text{H}_2\text{O}$.
Nearly insol. in cold, decomp. by warm H_2O .
 $+ 6\text{H}_2\text{O}$. (Braun, A. **132**. 47.)

— **iodide**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{I}_2$.
Sol. in H_2O . (Gibbs.)

— **iodosulphate**,
 $[\text{Co}(\text{NO}_2)(\text{NH}_3)_5]_2(\text{SO}_4)\text{I}_2$.
Sol. in H_2O .

— **periodosulphate**,
 $[\text{Co}(\text{NO}_2)(\text{NH}_3)_5]_2(\text{SO}_4)\text{I}_4$.
Easily decomp. by hot H_2O .

— **nitrate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5(\text{NO}_3)_2$.
Sl. sol. in cold, moderately sol. in hot H_2O .
Decomp. by boiling. Much less sol. than NH_4Cl or $(\text{NH}_4)_2\text{SO}_4$ in cold H_2O . Insol. in HNO_3 . (Gibbs and Genth.)

— **nitrite**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5(\text{NO}_2)_2 + 2\text{H}_2\text{O}$.
Sol. in H_2O . (Gibbs.)

— **cobaltic nitrite**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5(\text{NO}_2)_2 + 2\text{H}_2\text{O}$.
Sl. sol. in H_2O . (Gibbs, Proc. Am. Acad. **11**. 8.)

Is nitratopurpureocobaltic cobaltic nitrite,

$[(\text{NO}_2)\text{Co}(\text{NH}_3)_5]_2[\text{Co}(\text{NO}_2)_6]_2 + 2\text{H}_2\text{O}$. (Jørgensen, Z. anorg. **5**. 175.)

$[\text{Co}(\text{NO}_2)(\text{NH}_3)_5]_2[\text{Co}(\text{NO}_2)_6]_2$. Not so difficultly sol. as the luteo salt. (Jørgensen.)

Xanthocobaltic tetramine cobaltic nitrite,
 $\text{Co}_2(\text{NO}_2)_2(\text{NH}_3)_{10}[\text{Co}_2(\text{NH}_3)_4(\text{NO}_2)_4]_2$.

Can be recryst. from hot H_2O . (Gibbs, Proc. Am. Acad. **11**. 8.)

$= (\text{NO}_2)\text{Co}(\text{NH}_3)_5[(\text{NO}_2)_2(\text{NH}_3)_3\text{Co}(\text{NO}_2)_2]_2$.
Xanthocobaltic diamine cobaltic nitrite.
Very sl. sol. in cold H_2O . (Jørgensen, Z. anorg. **5**. 180.)

— **oxalate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{C}_2\text{O}_4$.

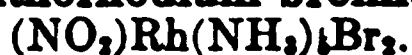
Nearly insol. in cold, sl. sol. in hot H_2O .

— **sulphate**, $\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{SO}_4$.

Moderately sol. in hot, much less in cold H_2O . Sol. without decomp. in $\text{H}_2\text{SO}_4 + \text{Aq}$. (Gibbs and Genth.)

Sol. in 25 pts. hot H_2O acidified with $\text{HC}_2\text{H}_3\text{O}_2$. (Jørgensen, Z. anorg. **5**. 172.)

$4\text{Co}(\text{NO}_2)(\text{NH}_3)_5\text{SO}_4, 3\text{H}_2\text{SO}_4$. Decomp. by H_2O , not by absolute alcohol. (Jørgensen.)

Xanthorhodium bromide,

Moderately sol. in H_2O . (Jørgensen, J. pr. (2) **34**. 394.)

— **chloride**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{Cl}_2$.

Much more sol. in H_2O than the nitrate.

— **chloroplatinate**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{PtCl}_4$.
Ppt. Extremely sl. sol. in cold H_2O .

— **dithionate**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{S}_2\text{O}_8 + \text{H}_2\text{O}$.
Nearly insol. in H_2O .

— **fluosilicate**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{SiF}_6$.
Ppt.

— **hydroxide**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5(\text{OH})_2$.

— **nitrate**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5(\text{NO}_3)_2$.

Moderately sol. in cold, easily in hot H_2O . Insol. in alcohol. Less sol. in conc. $\text{NH}_4\text{OH} + \text{Aq}$ than in H_2O .

Insol. in dil. $\text{HNO}_3 + \text{Aq}$; sol. in $\text{HNO}_3 + \text{Aq}$ of 1.4 sp. gr.

— **oxalate**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{C}_2\text{O}_4$.

Nearly insol. in cold H_2O . Very sl. sol. in warm H_2O . Easily sol. in dil. $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$.

— **sulphate**, $(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{SO}_4$.

Slowly sol. in cold, quite easily in hot H_2O .
 $4(\text{NO}_2)\text{Rh}(\text{NH}_3)_5\text{SO}_4, 3\text{H}_2\text{SO}_4$. Sl. sol. in cold, easily in hot H_2O . Can be recrystallized from dil. $\text{H}_2\text{SO}_4 + \text{Aq}$.

Xenon, Xe.Absorption by H₂O at t°.

t°	Absorption coefficient
0	0.2189
10	0.1500
20	0.1109
30	0.0900
40	0.0812
50	0.0878

(Antropoff, Roy. Soc. Proc. 1910, **83**. A, 480.)**Ytterbium, Yb.****Ytterbium bromide, YbBr₂ + 8H₂O.**Very sol. in H₂O. Hygroscopic. (Cleve, Z. anorg. 1902, **32**. 135.)**Ytterbium chloride, YbCl₃ + 6H₂O.**Very sol. in H₂O. (Cleve, Z. anorg. 1902, **32**. 134.)Mpt. 150–155°. Anhydrous salt is sol. in H₂O and in alcohol. (Matignon, A. ch. 1906, (8) **8**. 442.)**Ytterbium oxide, Yb₂O₃.**

Slowly attacked by cold or warm acids, but easily sol. at 100°.

Ytterbium oxychloride, YbOCl.Ppt. (Cleve, Z. anorg. 1902, **32**. 135.)**Yttrium, Y.**Decomposes H₂O. (Cleve, Bull. Soc. (2) **21**. 344.) Decomp. H₂O slightly at ord. temp., more rapidly by boiling. Easily sol. in dil. acids, even acetic acid. Slightly acted upon by conc. H₂SO₄. Decomposes hot KOH + Aq and cold NH₄Cl + Aq. Not attacked by NH₄OH + Aq. (Popp, A. **131**. 179.)

Popp's yttrium contained erbium.

Yttrium bromide, YBr₃.Sol. in H₂O with evolution of heat. (Duboin, C. R. **107**. 243.)+ 9H₂O. Deliquescent. Easily sol. in H₂O and alcohol. Insol. in ether. (Cleve.)**Yttrium carbide, YC₂.**Decomp. by H₂O and by dil acids; very slowly attacked by conc. acids. (Moissan, C. R. 1896, **122**. 575.)Decomp. by H₂O and dil. acids. (Pettersson, B. 1895, **28**. 2421.)**Yttrium chloride, YCl₃.**Anhydrous. Sol. in H₂O with evolution of heat. (Cleve.)+ 6H₂O. Deliquescent. Very sol. in H₂O. Sl. sol. in alcohol. Insol. in ether. (Cleve.)

Mpt. 156–160°; sol. in alcohol.

60.1 grams anhydrous salt are sol. in 100 grams of abs. alcohol at 15°.

6.5 grams are sol. in 100 grams pyridine. (Matignon, A. ch. 1906, (8) **8**. 437.)**Yttrium fluoride, YF₃ + ½H₂O.**

Nearly insol. in dil. acids. (Cleve.)

Yttrium hydroxide, Y₂O₃, 6H₂O or Y₂O₃.H₂O + 3H₂O.Insol. in H₂O.Insol. in KOH or NaOH + Aq. Easily sol. in acids. Sol. in alkali carbonates + Aq. When freshly pptd., easily sol. in NH₄Cl + Aq.**Yttrium iodide, YI₃.**Very deliquescent. Easily sol. in H₂O and alcohol.

Sl. sol. in ether. (Cleve.)

Yttrium oxide, Y₂O₃.Insol. in H₂O. Sl. sol. in cold HCl, HNO₃, or dil. H₂SO₄ + Aq, but gradually completely sol. on warming. Insol. in NH₄OH and sl. sol. in KOH + Aq. Sol. in HC₂H₃O₂ + Aq. Somewhat sol. in K₂CO₃ + Aq.**Yttrium peroxide, Y₄O₉.**(Cleve, Bull. Soc. (2) **43**. 53.)**Yttrium oxychloride, Y₂O₃.Cl₂.**Insol. in H₂O. (Popp.)**Yttrium sulphide, Y₂S₃.**Not prepared in pure state. Impure is insol. in H₂O, and partially decomp. thereby. Easily sol. in acids with decomp. (Popp.)**Zinc, Zn.**Not attacked by pure cold H₂O. Slowly oxidised by boiling H₂O. Pure H₂O free from O dissolved nothing from 2500 sq. mm. Zn. Presence of air containing CO₂ caused a solution of 3.5 mg. Zn, which maximum was reached in 2 days. Air without CO₂ also caused a slight action. (Snyders, B. **11**. 936.)100 ccm. distilled H₂O dissolved 14 mg. Zn from 11.8 sq. cm. in one week, during which air free from CO₂ was passed through the liquid, and 19 mg. when air containing CO₂ was used. (Wagner, Dingl. **221**. 260.)Filtered rain water was found to contain 20 mg. Zn per l. (Burg, Isis, **1873**. 119.)Very pure H₂O, when conducted through a great length of galvanized iron pipe, contained 1.7 pts. Zn to 100,000 pts. H₂O. (Davies, J. Soc. Chem. Ind. 1899, **18**. 102.)Action of H₂O on Zn in galvanized pipes is caused by electrolysis. (Smetham, C. N. 1879, **39**. 236.)All kinds of H₂O attack Zn, rain water the least.In distilled H₂O exposed to air Zn is abun-

dantly coated with ZnCO_3 , $2\text{ZnO} + 3\text{H}_2\text{O}$. By allowing 32 g. Zn to stand in 270 cc. distilled H_2O in a flask loosely stoppered with filter paper, 1.2 pts. Zn to 100,000 pts. H_2O was found in solution in 1-2 days. (Smith, J. Soc. Chem. Ind. 1904, 23, 475.)

Sol. in all acids. Very slowly sol. in dil. HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$ in glass vessels if Zn is pure. According to Jacquelin, 24 hours were necessary to dissolve 6 g. pure zinc. When fused at the lowest possible temperature, it is much more slowly sol. than when heated to a red heat. In both cases it is much more rapidly dissolved if cooled quickly. (Bolley, A. 95, 294; Rammelsberg.)

Dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ dissolves given % zinc in the same length of time (B=according to Bolley, R=according to Rammelsberg):

	Slowly cooled		Rapidly cooled	
	B	R	B	R
Cast at the melting point	42.5	74.1	13.0	0.9
Cast at a red heat	100.0	69.4	85.5	9.5

50 cc. $\text{H}_2\text{SO}_4 + \text{Aq}$ dissolved in 2 hours the following amts. from 1 sq. cm. Zn at t° :

t°	Strength of acid	Grms. dissolved
20	H_2SO_4	0.000
130	"	0.075
150	"	0.232
20	$\text{H}_2\text{SO}_4 + \text{H}_2\text{O}$	0.002
130	"	0.142
150	"	0.345
20	$\text{H}_2\text{SO}_4 + 2\text{H}_2\text{O}$	0.002
130	"	4.916
150	"	5.450
20	$\text{H}_2\text{SO}_4 + 3\text{H}_2\text{O}$	0.005
130	"	3.080
20	$\text{H}_2\text{SO}_4 + 4\text{H}_2\text{O}$	0.049
130	"	0.456
20	$\text{H}_2\text{SO}_4 + 5\text{H}_2\text{O}$	0.027
130	"	0.337
20	$\text{H}_2\text{SO}_4 + 6\text{H}_2\text{O}$	0.018
100	"	3.16

(Calvert and Johnson, Chem. Soc. 19, 437.)

C. P. zinc is more quickly sol. in dil. acids in vacuo than under normal pressure, the ratio being about 1.65. The rate of solubility increases slowly with rise of temp. from 0° to 98° , when it amounts to about 4 times that at 0° , but from 98° – 100° the increase is thirteenfold. Thus, as an average of 6 experiments, dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ (1.20) dissolves in 30 minutes 2.1 mg. Zn at 0° , 4.9 mg. at 20° ; 7.4 mg. at 60° , 9.3 mg. at 98° ; but 122.1 mg. at 100° . If, however, the acid was prevented from boiling by increasing the pressure, the sudden increase between 98° and 100° does not take place.

The rate of solubility in dil. $\text{H}_2\text{SO}_4 + \text{Aq}$ (1:20) is also increased 175 times by the addition of CrO_3 and 306 times by the addition of H_2O_2 . The above phenomena are explained by assuming the formation of a condensed hydrogen atmosphere around the metal, which prevents the further action of the acid. (Weeren, B. 24, 1785.)

Not attacked by $\text{HNO}_3 + \text{Aq}$ of 1.512 to 1.419 sp. gr. at a temp. of -18° or less, but violently attacked if temp. is raised. $\text{HNO}_3 + \text{Aq}$ of 1.419–1.401 sp. gr. does not attack Zn at temp. of a freezing mixture, but violently at 0° . More dil. $\text{HNO}_3 + \text{Aq}$ attacks Zn even at -20° . (Millon, A. ch. (3) 6, 99.)

Sol. in $\text{H}_2\text{CO}_3 + \text{Aq}$. (Berzelius.)

Solubility of Zn in acids is very much affected by the presence of small quantities of various metallic salts. Small amts. of $\text{PtCl}_4 + \text{Aq}$ accelerated the action of $\text{H}_2\text{SO}_4 + \text{Aq}$ 149 times, and As_2O_3 123 times. HgCl_2 has a strong retarding action owing to pptn. of Hg on the Zn.

The rate of solution of Zn in acids and the effect of changes in concentration and temperature and of the presence of inorganic salts and organic substances on this rate has been studied. 26 Tables are given. (Ericson-Aurén, Z. anorg. 1901, 27, 209–253.)

Speed of solution in H_2SO_4 and in HCl . (Centnerszwer, Z. phys. Ch. 1914, 87, 692.)

Various saline solutions have a strong solvent power in presence of PtCl_4 , i.e. KCl , NaCl , Na_2SO_4 , K_2SO_4 , $\text{MgSO}_4 + \text{Aq}$. PtCl_4 also causes Zn to decompose distilled H_2O . CuSO_4 has a similar but less energetic effect.

In all the above cases the disengagement of hydrogen is slower in the dark than in the light. (Millon, C. R. 21, 37.)

According to Barreswill (C. R. 21, 292) the above reactions are all caused by galvanic action due to pptd. metal, and a piece of Pt in contact with the Zn causes the same action as the PtCl_4 in solution.

Easily sol. in alkalis + Aq , even $\text{NH}_4\text{OH} + \text{Aq}$, especially when the Zn is in contact with Fe. Sol. in $\text{NaCl} + \text{Aq}$ with pptn. of ZnO . (Siersch, J. B. 1867, 257.)

Sol. in sat. alkali and alkali-earth chlorides + Aq . (Post, 1872.)

Sol. in NH_4 salts + Aq . (Lorin, J. B. 1866, 124.)

Sol. in sat. Na_2SO_4 , K_2SO_4 , MgSO_4 , NaNO_3 , KNO_3 , $\text{Ba(NO}_3)_2$, CaCl_2 , MgCl_2 , and $\text{NH}_4\text{NO}_3 + \text{Aq}$. Chlorides and sulphates (especially Na_2SO_4 and MgCl_2) have strongest action, MgSO_4 and nitrates the least. The action was greatly increased by heat. (Seydewitz, B. 11, 936.)

Sol. in boiling $\text{NH}_4\text{Cl} + \text{Aq}$. Sol. in neutral $\text{FeCl}_2 + \text{Aq}$ with pptn. of Fe, especially easily at 100° . (Capitaine, C. R. 9, 737.)

Sol. in $\text{NiSO}_4 + \text{Aq}$ with pptn. of NiO . (Tupputi.)

Sol. in conc. hot $\text{ZnCl}_2 + \text{Aq}$, but Zn oxy-

chloride is pptd. on diluting. (Ordway, Am. J. Sci. (2) **23**. 222.)

Sol. in $\text{GlSO}_4 + \text{Aq}$. (Debray.)

Solubility of Zn in dilute solutions of salts: 100 ccm. of solutions of the given salts were allowed to act one week on 11.8 sq. cm. Zn while a current of air with or without CO_2 was passed through the solution.

Salt	G. salt in 100 ccm. solution	Mg. Zn dissolved without CO_2	Mg. Zn dissolved with CO_2
NaCl or KCl NH ₄ Cl	0.5	7	38
MgCl ₂	1.0	51	36
K ₂ SO ₄	0.83	18	54
KNO ₃	1.0	30	53
Na ₂ CO ₃	1.0	9	37
NaOH	1.0	13	..
CaO, H ₂	0.923	60	..
	Sat	3	..

(Wagner, Dingl. **221**. 260.)

Action of dil. salt solutions (1%) on Zn. The following amts. of Zn in mg. were dissolved from 2500 sq. mm. Zn in 14 days by 400 ccm. 1% solution of the given salts:

Salt	Mg. Zn	Salt	Mg. Zn
NaCl	11.2	NaNO ₃	6
KCl	14.8	Ba(NO ₃) ₂	8
CaCl ₂	15.2	NH ₄ Cl	24.0
MgCl ₂	17.2	(NH ₄) ₂ SO ₄	31.6
BaCl ₂	13.2	NH ₄ NO ₃	26.0
K ₂ SO ₄	12.0	NaHCO ₃	0
MgSO ₄	8.8	K ₂ CO ₃	0
KNO ₃	6.8	Na ₂ CO ₃	0

Sl. attacked by H_2O at 80° , by hot conc. NH_4OH ; attacked by H_3PO_4 or $\text{NaCl} + \text{Aq}$; very sl. attacked by $\text{NaNO}_3 + \text{Aq}$ or $\text{KNO}_3 + \text{Aq}$ at 100° . (Smith, J. Soc. Chem. Ind. 1904, **23**. 476.)

$\frac{1}{2}$ ccm. oleic acid dissolves 0.0240 g. Zn in 6 days. (Gates, J. phys. Chem. 1911, **15**. 143.)

Attacked by cane sugar + Aq at 115° . (Klein and Berg, C. R. **102**. 1170.)

Zinc amide, $\text{Zn}(\text{NH}_2)_2$.

Decomp. by H_2O and alcohol. Insol. in ether. (Frankland, Phil. Mag. (4) **15**. 149.)

Zinc antimonide, ZnSb .

Does not decomp. boiling H_2O except slightly. Not attacked by dil. mineral acids, but decomp. by conc. HCl or $\text{HNO}_3 + \text{Aq}$. (Cooke, Proc. Am. Acad. **5**. 348.)

Zn_2Sb_2 . Decomp. H_2O rapidly at 100° .

Violently decomp. by dil. HCl or $\text{H}_2\text{SO}_4 + \text{Aq}$, also by $\text{HNO}_3 + \text{Aq}$. Completely sol. in $\text{HCl} + \text{Aq}$ mixed with a little HNO_3 . (Cooke.)

Zinc azoimide, basic, $\text{Zn}(\text{OH})\text{N}_3$.

Very sl. sol. in H_2O . Decomp. by hot H_2O . (Curtius, J. pr. 1898, (2) **58**. 293.)

Zinc azoimide ammonia, $\text{ZnN}_3, 2\text{NH}_3$.

Insol. in H_2O , but gradually decomp. thereby. (Dennis, J. Am. Chem. Soc. 1907 **29**. 20.)

Zinc bromide, ZnBr_2 .

Very deliquescent, and sol. in H_2O .

Sat. $\text{ZnBr}_2 + \text{Aq}$ contains at:

-20° $+4^\circ$ 22° 97°
66.3 68.8 77.5 83.6% ZnBr_2 ,

107° 170° 210° 375° (mpt.)
83.8 85.0 89.3 100% ZnBr_2 .

(Étard, A. ch. 1894, (7) **2**. 541.)

Solubility in H_2O .

100 g. of the sat. solution contain at:

35° 40° 60° 80° 100°
85.45 85.53 86.08 86.57 87.05 g. ZnBr_2 .

(Dietz, Z. anorg. 1899, **20**. 250.)

See also below under hydrated salts.

Sp. gr. of $\text{ZnBr}_2 + \text{Aq}$ at 19.5° containing

18.3 31.7 43.2 % ZnBr_2 ,
1.1849 1.3519 1.5276

52.6 59.1 68 % ZnBr_2 ,
1.7082 1.8525 2.1027

(Kremers, Pogg. **108**. 117.)

Sp. gr. of $\text{ZnBr}_2 + \text{Aq}$ at 19.5° .

% ZnBr_2	Sp. gr.	% ZnBr_2	Sp. gr.	% ZnBr_2	Sp. gr.
5	1.045	25	1.265	45	1.560
10	1.093	30	1.330	50	1.650
15	1.196	35	1.400	55	1.755
20	1.204	40	1.475	60	1.875

(Kremers, calculated by Gerlach, Z. anal. **8**. 285.)

Sol. in conc. HCl or $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$, also in $\text{NH}_4\text{OH} + \text{Aq}$.

Sol. in AlBr_3 . (Isbekow, Z. anorg. 1913. **84**. 27.)

Very sl. sol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, **20**. 830.)

Sol. in alcohol and ether. (Berthemot, J. Pharm. **14**. 610.)

Sol. in methyl acetate. (Naumann, B. 1909, **42**. 3790.)

Insol. in ethyl acetate. (Naumann, B. 1910, 43, 314.)

More sol. in anhydrous ether than in abs. alcohol. Insol. in CS_2 . (Hampe, Ch. Z. 1887, 11, 846.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51, 236.)

Mol. weight determined in pyridine. (Werner, Z. anorg. 1897, 15, 22.)

+ H_2O . (Lescœur, A. ch. 1894, (7) 2, 78.)

+ $2\text{H}_2\text{O}$. Very hygroscopic.

Solubility in H_2O .

100 g. of the sat. solution contain at:

—8° 0° 13° 25° 30° 37° (mpt.)
79.06 79.55 80.76 82.46 84.08 86.20 g. ZnBr_2 .

(Diets, Z. anorg. 1899, 20, 250.)

+ $3\text{H}_2\text{O}$. Solubility in H_2O .

100 g. of the sat. solution contain at:

—15° —10° —5° (mpt.)
77.13 78.45 80.64 g. ZnBr_2 .

(Diets, Z. anorg. 1899, 20, 250.)

Zinc bromide ammonia, $\text{ZnBr}_2 \cdot 2\text{NH}_3$.

Decomp. by H_2O . Sl. sol. in cold, more easily in warm $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg, Pogg. 55, 240.)

+ $\frac{1}{2}\text{H}_2\text{O}$. Decomp. by H_2O with separation of ZnO . (André, C. R. 96, 703.)

+ H_2O . Above salt of Rammelsberg's has this composition. (André.)

$3\text{ZnBr}_2 \cdot 8\text{NH}_3 + 2\text{H}_2\text{O}$. Decomp. by H_2O (André.)

$3\text{ZnBr}_2 \cdot 10\text{NH}_3 + \text{H}_2\text{O}$. Decomp. by H_2O (André.)

$2\text{ZnBr}_2 \cdot 10\text{NH}_3$. Efflorescent. Decomp. by H_2O . (André.)

Zinc bromide cupric oxide, $\text{ZnBr}_2 \cdot 3\text{CuO} + 2\text{H}_2\text{O}$.

+ $4\text{H}_2\text{O}$. (Mailhe, C. R. 1901, 133, 227.)

Zinc bromide hydrazine, $\text{ZnBr}_2 \cdot 2\text{N}_2\text{H}_4$.

Decomp. by H_2O

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Fransen, Z. anorg. 1908, 60, 277.)

Zinc chloride, ZnCl_2 .

Very deliquescent, and sol. in H_2O .

Sol. in 0.333 pt. H_2O at 18.75° (Abt.)

$\text{ZnCl}_2 + \text{Aq}$ sat. at 12.5° contains 78.5% ZnCl_2 . (Hassendratz, A. ch. 28, 291.)

Solubility in H_2O .

100 g. of the sat. solution contain at:

15° 20° 41° 60° 100°
79.12 81.19 82.21 83.51 86.01 g. ZnCl_2 .

(Diets, Z. anorg. 1899, 20, 245.)

$\text{ZnCl}_2 + \text{Aq}$ containing 1 pt ZnCl_2 in 1.8205 pts. H_2O at 18° has sp. gr. = 1.3666. (Hittorf, Z. phys. Ch. 1902, 39, 628.)

Solubility in H_2O at t°.
100 g. H_2O dissolve g. ZnCl_2 .

t°	Solid phase	% ZnCl_2	
—5	ice	14	
—10	"	25	
—40	"	83	
—62	ice. 4 aq	104	cryohydrate pt.
—50	$\text{ZnCl}_2 + 4\text{H}_2\text{O}$	113	
—40	"	127	
—30	4 aq. 3 aq	160	transition point
—10	$\text{ZnCl}_2 + 3\text{H}_2\text{O}$	189	
0	"	208	
+5	"	230	
6.5	"	252	mpt.
8	"	282	
0	3 aq. 1 1/2 aq	309	eutectic point
0	$\text{ZnCl}_2 + 2 1/2 \text{H}_2\text{O}$	235	
6.5	2 1/2 aq. 3 aq	252	transition point
10	$\text{ZnCl}_2 + 2 1/2 \text{H}_2\text{O}$	272	
12.5	"	303	mpt.
11.5	2 1/2 aq. 1 1/2 aq	333	eutectic point
9	2 1/2 aq. 1 aq	360	eutectic point
6	$\text{ZnCl}_2 + 2 1/2 \text{H}_2\text{O}$	385	
—6	$\text{ZnCl}_2 + 1 1/2 \text{H}_2\text{O}$	298	
+10	"	330	
20	"	368	
26	1 1/2 aq. 1 aq	423	transition point
26.8	1 1/2 aq. ZnCl_2	433	transition point
0	$\text{ZnCl}_2 + \text{H}_2\text{O}$	342	
10	"	364	
20	"	396	
28	1 aq. ZnCl_2	436	transition point
31	$\text{ZnCl}_2 + \text{H}_2\text{O}$	477	
25	ZnCl_2	432	
40	"	452	
60	"	488	
80	"	543	
100	"	615	
262	"		mpt.

(Mylus and Diets, Z. anorg. 1905, 44, 31)

See also below under hydrated salts.

Sp. gr. of $\text{ZnCl}_2 + \text{Aq}$ at 19.5°.

% ZnCl_2	Sp. gr.	% ZnCl_2	Sp. gr.
13.8	1.1275	37.5	1.2000
25.8	1.2466	49.2	1.3001

(Kremers, Pogg. 106, 360.)

Sp. gr. of $\text{ZnCl}_2 + \text{Aq}$ at 19.5°.

% ZnCl_2	Sp. gr.	% ZnCl_2	Sp. gr.	% ZnCl_2	Sp. gr.
1	1.010	25	1.238	45	1.48
5	1.045	30	1.291	50	1.56
10	1.091	35	1.352	55	1.65
15	1.137	40	1.420	60	1.74
20	1.186				

(Gerlach, Z. anal. 8, 283, calculated from Kremers.)

Sp. gr. of $\text{ZnCl}_2 + \text{Aq}$ at t°.

t°	15°	15°	15°	15°
% ZnCl_2	2.5	4.89	10.0	20.1
Sp. gr.	1.024	1.046	1.094	1.11

t°	15°	15°	15°
% ZnCl_2	29.86	40.0	58.88
Sp. gr.	1.297	1.423	1.728

(Long, W. Ann. 1890, 11, 88.)

Sp. gr. of $\text{ZnCl}_2 + \text{Aq}$ at room temp. containing:

15.334 23.487 33.752% ZnCl_2 .
1.1459 1.2288 1.3431

(Wagner, W. Ann. 1883, 18. 267.)

Sp. gr. of $\text{ZnCl}_2 + \text{Aq}$ at 25°.

Concentration of $\text{ZnCl}_2 + \text{Aq}$	Sp. gr.
1—normal	1.0590
$\frac{1}{2}$ —	1.0302
$\frac{1}{4}$ —	1.0152
$\frac{1}{8}$ —	1.0077

(Wagner, Z. phys. Ch. 1890, 5. 40.)

Sp. gr. of $\text{ZnCl}_2 + \text{Aq}$.

$\frac{1}{2}$ ZnCl_2 g. in 1000 g. of solution	Sp. gr. 16°/16°
0	1.000000
0.5994	1.000560
2.3163	1.002163
5.0406	1.004708
9.8988	1.009243
19.4914	1.018228

(Dijken, Z. phys. Ch. 1897, 24. 108.)

Insol. in SbCl_3 . (Klemensiewicz, C. C. 1908, II. 1850.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 830.)

Easily sol. in hot absolute alcohol, and ether. Sol. in 1 pt. strong alcohol at 12.5°. (Wenzel.)

Sol. in 0.35 pt. absolute alcohol. (Graham.)

Sol. in butyl (Wurtz), and hexyl (Bouis) alcohol at ord. temp., but decomp. on heating.

Very sol. in acetic ether with evolution of heat. (Cann, C. R. 102. 363.)

Easily sol. in acetone. (Krug and M'Elroy, J. Anal. Ch. 6. 184.)

1 g. ZnCl_2 is sol. in 2.3 g. acetone at 18°. Sp. gr. of sat. solution 18°/4° = 1.14. (Naumann, B. 1904, 37. 4338.)

Sol. in acetone and in methylal. (Eidmann, C. C. 1899, II. 1014.)

Sol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Sol. in hot benzonitrile, also in other aromatic nitriles.

Sol. in methyl sulphide and in ethyl sulphide. Very sol. in piperidine. (Werner, Z. anorg. 1897, 15. 7.)

Sol. in benzyl alcohol, furfurol, methylpropylketone, acetophenone, ethyl monochloracetate, ethyl cyanacetate, ethyl acetoacetate, ethyl benzoate, ethyl oxalate, amyl nitrite, pyridine, piperidine, and quinoline.

Insol. in salicylic aldehyde, ethyl nitrate, and nitrobenzene. (Lincoln, J. phys. Chem. 1899, 3. 460.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, 51. 236.)

Sol. in 2 pts. glycerine at ord. temp. (Clever, Bull. Soc. 1872, (2) 18. 372.)

100 g. glycerol dissolve 50 g. ZnCl_2 at 15.5°. (Ossendowski, Pharm. J. 1907, 79. 575.)

Insol. in CS_2 . (Arctowski, Z. anorg. 1894, 6. 257.)

Sol. in urethane. (Castoro, Z. anorg. 1899, 20. 61.)

Mol. weight determined in piperidine; pyridine and methyl sulphide. (Werner, Z. anorg. 1897, 15. 18.)

+ H_2O . Very deliquescent. Contains $1\frac{1}{2}$ H_2O . (Engel, C. R. 102. 1111.)

Solubility in H_2O .

100 g. of the sat. solution contain at:

0° 11° 27° (mpt.)
74.33 78.25 84.61 g. ZnCl_2 .

(Dietz, Z. anorg. 1899, 20. 245.)

+ $1\frac{1}{2}\text{H}_2\text{O}$.

Solubility in H_2O .

100 g. of the sat. solution contain at:

0° 10° 20° 26° (mpt.)
67.45 73.65 80.08 83.43 g. ZnCl_2 .

(Dietz, Z. anorg. 1899, 20. 245.)

+ $2\text{H}_2\text{O}$. Sat. aq. solution contains at:

—20° —14° —10° —4° —1°
54.7 55.4 56.5 57.4 57.9% salt,

+5° 9° 15° 33° 42°
59.1 60.2 62.0 66.8 68.3% salt.

(Étard, A. ch. 1894, (7) 2. 536.)

Solubility in H_2O .

100 g. of the sat. solution contain at:

0° 10° 19° (mpt.)
67.56 73.70 79.07 g. ZnCl_2 .

(Dietz, Z. anorg. 1899, 20. 245.)

+ $2\frac{1}{2}\text{H}_2\text{O}$. Solubility in H_2O .

100 g. of the sat. solution contain at:

0° 8° 13° (mpt.)
67.42 71.96 75.14 g. ZnCl_2 .

(Dietz, Z. anorg. 1899, 20. 245.)

+ $3\text{H}_2\text{O}$. Sol. in 12.5 pts. H_2O at 0°. (Engel.)

Solubility in H_2O .

100 g. of the sat. solution contain at:

—5° 0° +7° (mpt.)
64.5 67.58 71.57 g. ZnCl_2 .

(Dietz, Z. anorg. 1899, 20. 245.)

Zinc hydrogen chloride, $2\text{ZnCl}_2, \text{HCl} + 2\text{H}_2\text{O}$.

Deliquescent. (Engel, C. R. 102. 1068.)

$\text{ZnCl}_2, \text{HCl} + 2\text{H}_2\text{O}$. (Engel.)

Zinc hydrazine chloride, $\text{ZnCl}_2, \text{N}_2\text{H}_4, \text{HCl}$.

Very hygroscopic.

Sol. in H_2O . (Curtius, J. pr. 1894, (2) 338.)

ZnCl₂, 2(N₂H₄, HCl.) Hygroscopic; very sol. in H₂O.

Sol. in hot alcohol and NH₄OH + Aq. (Curtius, J. pr. 1894, (2) 50. 338.)

Zinc chloride ammonia, ZnCl₂, 5NH₃ + H₂O.

Easily sol. in little, but decomp. by much H₂O. Still more sol. in ZnCl₂ + Aq with decomp. (Divers, C. N. 18. 13.)

ZnCl₂, 4NH₃ + H₂O. (Kane.)

ZnCl₂, 2NH₃. Not completely sol. in H₂O; can be recryst. from hot NH₄Cl + Aq. (Ritt-hausen, J. pr. 60. 473.)

Insol. in H₂O. Sol. in NH₄Cl or NH₄OH + Aq. (Thomas, B. 20. 743.)

+1/4 H₂O.

+2/5 H₂O.

+1/2 H₂O.

+H₂O.

} Decomp. by H₂O.

(André, C. R. 1882, 94. 964.)

ZnCl₂, NH₃. Decomp. by H₂O. (Kane, A. ch. 72. 290.)

Zinc chloride cupric oxide, ZnCl₂, 3CuO + 4H₂O.

(Mailhe, C. R. 1901, 134. 226.)

Zinc chloride hydrazine, ZnCl₂, 2N₂H₄.

Ppt. (Franzen, Z. anorg. 1908, 60. 275.)

ZnCl₂, 2N₂H₄. Insol. in H₂O.

Easily sol. in NH₄OH + Aq. (Curtius, J. pr. 1894, (2) 50. 345.)

Zinc chloride hydroxylamine, ZnCl₂, 2NH₂OH.

Sl. sol. in cold, somewhat more in warm H₂O. Very sol. in NH₂OH + Aq. Very sl. sol. in alcohol and other organic solvents. (Crismer, Bull. Soc. (3) 3. 116.)

1 pt. is dissolved in 100 pts aq. solution sat. at 20°. (Antonoff, C. C. 1905, II. 810.)

Zinc fluoride, ZnF₂.

Sl. sol. in cold, more easily in hot H₂O. Insol. in 95% alcohol. Sol. in boiling HNO₃, HCl, or H₂SO₄. (Poulenc, C. R. 116. 581.)

Contrary to older statements, ZnF₂ is quite sol. in H₂O. (Kohlrausch, Z. phys. Ch. 1903, 44. 213.)

Insol. in liquid NH₃. (Gore, Am. Ch. J. 1898, 20. 830.)

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

+4H₂O. Difficultly sol. in H₂O. Somewhat more sol. in H₂O containing HF, HCl, or HNO₃. Easily sol. in NH₄OH + Aq. (Berzelius, Pogg. 1. 26.)

1 l. H₂O dissolves 16 g. at 18°. (Dietz)

Zinc hydrogen fluoride.

Known only in solution.

Zinc zirconium fluoride.

See Fluozirconate, zinc.

Zinc hydrophosphide, Zn₂H₂P₂.

Decomp. by cold H₂O and by dil. HCl + (Drechsel and Finkelstein, B. 1871, 4. 3.)

Zinc hydroxide, ZnO₂H₂.

Insol. in H₂O. Sol. in acids. Sol. in K⁺ NaOH, NH₄OH, or (NH₄)₂CO₃ + Aq.

1 l. H₂O dissolve 0.01 g. ZnO₂H₂ at (Bodländer, Z. phys. Ch. 1898, 27. 66.)

Solubility in H₂O is calculated to be 2 10⁻⁵ g. mols. per l. (Herz, Z. anorg. 1900, 227.)

1 l. H₂O dissolves 0.0042 g. ZnO₂H₂ at (Dupre and Bialas, Z. angew. Ch. 1903, 55.)

See also Zinc oxide.

Solubility in NH₄OH + Aq at 25°.

ZnO ₂ H ₂ used	NH ₃ norm.	G. ZnO pr.
prepared from ZnSO ₄	1.287	7.28
	0.825	3.84
	0.311	0.55
prepared from Zn(NO ₃) ₂	0.321	0.34
	0.643	0.845
	1.215	2.70
	1.928	5.07
	2.570	7.01
	3.213	10.16

(Bonsdorff, Z. anorg. 1904, 41. 189.)

Solubility of ZnO₂H₂ in NH₄OH and ammonium bases + Aq at 17°-19°.

Normality of the base	G. ZnO in 20 cc. of the solution
0.0942 NH ₃	0.00185
0.236 NH ₃	0.01795
0.707 NH ₃	0.0959
0.0944 NH ₂ CH ₃	0.0008
0.472 NH ₂ CH ₃	0.01325
0.944 NH ₂ CH ₃	0.0484
0.068 NH ₂ C ₂ H ₅	0.0005
0.51 NH ₂ C ₂ H ₅	0.0074
0.68 NH ₂ C ₂ H ₅	0.01605
NH(C ₂ H ₅) ₂	insol.
NH(CH ₃) ₂	"

(Herz, Z. anorg. 1902, 30. 280.)

Solubility in NH₄OH + Aq increases with increasing concentration of NH₄OH. (Eul B. 1903, 36. 3401.)

2 pts. ZnO₂H₂ dissolve in 5 pts. KOH + A (sp. gr. = 1.3.) (Bonnet.)

Solubility of ZnO₂H₂ in NaOH+Aq.

G. Na in 20 ccm.	G. Zn in 20 ccm.
0.1012	0.0040
0.1978	0.0150
0.4278	0.0442
0.6670	0.1771
0.9660	0.9630
1.4951	0.2481
2.9901	0.3700

When zinc hydroxide is treated with alkali, more dissolves at first than corresponds with the true equilibrium under the prevailing conditions, for such solutions spontaneously deposit more or less zinc hydroxide according to the concentration. (Rubenbauer, Z. anorg. 1902, **30**. 333.)

Solubility of ZnO₂H₂ in NaOH+Aq at 25°.

G. mol. per l.	
Na	Zn
0.2636	0.00311
0.3871	0.0057
0.5414	0.0129
0.9280	0.0425

(Wood, Chem. Soc. 1910, **97**. 884.)

Freshly pptd. ZnO₂H₂ is easily sol. in KOH+Aq, but it gradually goes over into a stable form which is difficultly sol. in KOH+Aq. (Herz, Z. anorg. 1901, **28**. 474.)

Freshly pptd. ZnO₂H₂ is sol. in dil. salt solutions (1 %) as follows. The given amts. in mg. (calculated as Zn) were dissolved per l. at t°.

Salt	Mg. Zn	t°
NaCl	51	18
KCl	43	20
CaCl ₂	57.5	16
MgCl ₂	65	16
BaCl ₂	38	18
K ₂ SO ₄	37.5	20
MgSO ₄	27	21
KNO ₃	17.5	15
NaNO ₃	22	15
Ba(NO ₃) ₂	25	21
K ₂ CO ₃	0	15
NH ₄ Cl	95	20
NH ₄ NO ₃	77	20
(NH ₄) ₂ SO ₄	88	20

(Snyders, B. **11**. 936.)

+H₂O.
See also Zinc oxide.

Zinc hydrosulphide, Zn(SH)₂.

Very unstable. Decomp. by H₂O. (Zotta, M. **10**. 807.)

Sol. in NaSH+Aq. (Thomsen, B. **11**. 2044.)
Zn₂H₂S₄. (Zotta.)

Zinc iodide, ZnI₂.

Deliquescent. Easily sol. in H₂O.
Sat. ZnI₂+Aq contains at:

—18°	—5°	+17°	47°	62°	73°
70.9	74.0	80.4	80.3	81.3	81.2% ZnI ₂ ,
97°	100°	107°	138°	140°	
82.1	83.0	82.6	83.8%		ZnI ₂ .

(Étard, A. ch. 1894, (7) **2**. 544.)

Solubility in H₂O.
100 g. of the sat. solution contain at:

0°	18°	40°	60°	80°	100°
81.11	81.20	81.66	82.37	83.05	83.62 g. ZnI ₂ .

(Dietz, Z. anorg. 1899, **20**. 251.)

See also under +2H₂O.

Sp. gr. of ZnI₂+Aq at 19.5° containing:

23.1	42.6	56.3	63.5	76.0%
1.2340	1.5121	1.7871	1.9746	2.3976

(Kremers, Pogg. **111**. 61.)

Sp. gr. of ZnI₂+Aq at 19.5° containing:

5	10	15	20	25	% ZnI ₂ ,
1.045	1.091	1.140	1.196	1.255	
30	35	40	45	50	% ZnI ₂ ,
1.368	1.390	1.420	1.560	1.650	
55	60	65	70	75	% ZnI ₂ .
1.754	1.875	2.020	2.180	2.360	

(Kremers, calculated by Gerlach, Z. anal. **8**. 285.)

Sol. in (NH₄)₂CO₃+Aq.
Moderately sol. in liquid NH₃. (Franklin, Am. Ch. J. 1898, **20**. 830.)
Sol. in alcohol.

100 pts. glycerine dissolve 40 pts. at ord. temp. (Klever, Bull. Soc. 1872, (2) **18**. 372.)
100 g. glycerol dissolve 40 g. ZnI₂ at 15.5°. (Ossendowski, Pharm. J. 1907, **79**. 575.)

More sol. in anhydrous ether than in abs. alcohol. Insol. in CS₂. (Hampe, Ch. Z. 1887, **11**. 846.)

Sol. in methyl acetate (Naumann, B. 1909, **42**. 3790); acetone. (Eidmann, C. C. **1899**, II. 1014; Naumann, B. 1904, **37**. 4328.)

Sol. in quinoline. (Beckmann and Gabel, Z. anorg. 1906, **51**. 236.)

Mol. weight determined in methyl sulphate. (Werner, Z. anorg. 1897, **15**. 25.)
+2H₂O. Solubility in H₂O.

100 g. of the sat. solution contain at:
—10° —5° 0° +10° 22° 27°(mpt.)
80.50 80.77 81.16 82.06 83.12 89.52 g. ZnI ₂ .

(Dietz, Z. anorg. 1899, **20**. 251.)

+4H₂O. (Lubarski, Z. anorg. 1898, **18**. 387.)

Zinc tetraiodide, ZnI₄.

Known only in aqueous solution. (Baup, Repert. **14**. 412.)
Sol. in fenchone. (Rimini and Oliveri, C. C. **1907**, II. 241.)

Zinc iodide ammonia, $\text{ZnI}_2, 4\text{NH}_3$.

Decomp. by cold H_2O . Easily sol. in acids and $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg, Pogg. 48. 152.)

$\text{ZnI}_2, 5\text{NH}_3$. Decomp. by cold H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Rammelsberg.)

$3\text{ZnI}_2, 5\text{NH}_3 + 3\text{H}_2\text{O}$. (Tassily, C. R. 1896, 122. 324.)

Zinc iodide hydrazine, $\text{ZnI}_2, 2\text{N}_2\text{H}_4$.

Decomp. by H_2O .

Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Franzen, Z. anorg. 1908, 60. 277.)

Zinc nitride, Zn_3N_2 .

Decomp. by H_2O with the greatest violence. (Frankland, Phil. Mag. (4) 15. 149.)

Easily decomp. by H_2O when finely powdered. (Rossel, C. R. 1895, 121. 942.)

Sol. in HCl . (Fischer, B. 1910, 43. 1468.)

Zinc oxide, ZnO .

Insol. in H_2O . Some preparations of ZnO are sl. sol. in H_2O , never, however, in less than 1 million pts. H_2O . (Bineau, C. R. 41. 510.)

Calculated from electrical conductivity of $\text{ZnO} + \text{Aq}$. 1 pt. ZnO is sol. in 236,000 pts. H_2O at 18° . (Dupre and Bialas, Zeit. angew. Ch. 1903, 16. 55.)

See also **Zinc hydroxide**.

Easily sol. in acids, even after ignition. Easily sol. in acids, even H_2SO_3 , or $\text{H}_2\text{CO}_3 + \text{Aq}$.

Solubility of ZnO in $\text{CrO}_3 + \text{Aq}$ at 25° .

1 l. of the solution contains:

G. CrO_3	G. ZnO	G. CrO_3	G. ZnO
0.010	0.013	101	44.9
0.010	0.013	151	66.1
0.010	0.013	192	83.8
0.604	0.409	192	83.6
2.14	1.16	285	123
4.19	2.24	392	168
11.4	5.84	450	193
11.5	5.89	461	196
22.2	10.7	463	197
31.4	14.9	475	202
43.1	20.1	574	240
57.5	26.7	660	274
66.5	30.3	769	318
66.7	30.4	879	354
70.6	32.2	970	389
93.3	41.5

(Gröger, Z. anorg. 1911, 70. 136.)

When moist is easily sol. in KOH , NaOH , and $\text{NH}_4\text{OH} + \text{Aq}$, but only sl. sol. therein after ignition. Partially repptd. from solution in $\text{NH}_4\text{OH} + \text{Aq}$ by dilution with H_2O .

Anhydrous ZnO is insol. in dil., but sol. in conc. alkali hydrates + Aq , but the hydroxide is easily sol. even in dil. alkalies + Aq . (Fremy, A. ch. (3) 23. 390.)

Very sl. sol. in $\text{NH}_4\text{OH} + \text{Aq}$. After igni-

tion its solubility is greatly increased by traces of K and NH_4 salts. Phosphates have the strongest action, then, in the following order: arsenates, chlorides, sulphites, nitrates, acetates, carbonates, tartrates, citrates, and sulphates. Succinates and benzoates increase the solubility in $\text{NH}_4\text{OH} + \text{Aq}$, only when it is very dil.; borates, iodides, chlorates, arsenites, gallates, and oxalates do not increase the solubility. (Schindler.)

ZnO is sol. in $\text{NH}_4\text{OH} + \text{Aq}$. only in presence of NH_4 salts. (Brandhorst, Zeit. angew. Ch. 1904, 17. 513.)

Solubility in KOH , NaOH , and $\text{NH}_4\text{OH} + \text{Aq}$.

An excess over 4 mols. KOH to 1 mol. ZnO is necessary for solution, but that excess may be neutralised after solution, until only 4 mols. are left, without pptn. of ZnO . Solution is pptd. by addition of 12 vols. H_2O . $\text{KOH} + \text{Aq}$ containing 16.5 g. KOH to a litre H_2O is the weakest solution which will dissolve ZnO . Three times as much alkali are necessary for solution at 50° as at $16-17^\circ$. Less excess of NaOH than of KOH is necessary. 3 mols. NH_4OH will dissolve 1 mol. ZnO , and the temp. and dilution are in this case of little influence. (Prescott.)

100 cc. of 20% $\text{NaOH} + \text{Aq}$ dissolve in many hours at most 2.97 g. ignited ZnO . Pptd. ZnO is more quickly dissolved but the action becomes very slow after 100 cc. of the solution contain 3.87 g. of Zn . (Förster and Günther, Z. Elektrochem. 1900, 6. 301.)

Solubility of ignited ZnO in $\text{NaOH} + \text{Aq}$ gradually decreases. (Kunschert, Z. anorg. 1904, 41. 343.)

Sol. in hot $\text{NH}_4\text{Cl} + \text{Aq}$, either when moist or dry.

Somewhat less sol. in $\text{NH}_4\text{NO}_3 + \text{Aq}$.

Somewhat sol. in water glass + Aq . (Ordway.)

Slowly sol. in cold, easily in hot $\text{NaCl} + \text{Aq}$. (Siersch, J. B. 1867. 255.)

Solubility of ZnO in $\text{ZnCl}_2 + \text{Aq}$ at room temp.

G. ZnCl_2 per 100 g. H_2O	G. ZnO per 100 g. H_2O
8.22	0.0137
23.24	0.138
45.95	0.497
51.50	0.604
56.90	0.723
62.85	0.884
96.00	1.792
124.70	3.213
144.80	2.640
203.00	1.590

The solubility curve has a maximum at a point corresponding to about 125 g. ZnCl_2 per 100 g. H_2O . On the first branch of the curve the solid phase in equilibrium with the solution is $\text{ZnCl}_2, 4\text{ZnO}, 6\text{H}_2\text{O}$; on the second branch it is $\text{ZnCl}_2, \text{ZnO}, 1.5\text{H}_2\text{O}$.

(Driot, C. R. 1910, 150. 1427.)

Sol. in boiling $\text{Fe}(\text{NO}_3)_3$, and $\text{Pb}(\text{NO}_3)_2$ + Aq with pptn. of oxides. Not attacked by $\text{Co}(\text{NO}_3)_2$, $\text{Ni}(\text{NO}_3)_2$, and $\text{Ce}(\text{NO}_3)_3$ + Aq. (Persoz.)

Sol. in boiling KCN + Aq.

Insol. in boiling K tartrate + Aq. (Kahlenberg and Hillyer, Am. Ch. J. 1894, 16. 101.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 830.)

Tartaric acid somewhat hinders the pptn. of ZnO_2H_2 .

Insol. in methyl acetate. (Naumann, B. 1909, 42. 3790.)

Insol. in acetone. (Naumann, B. 1904, 37. 4329; Eidmann, C. C. 1899, II. 1014.)

Sol. in methyl amine, but insol. in amyl amine + Aq. (Wurtz.)

1 l. solution containing 174.4 g. sugar and 14.1 g. CaO dissolves 0.24 g. ZnO . (Bodenbender, J. B. 1865, 600.)

Min. *Zincite*. Sol. in acids.

Zinc peroxide.

1 pt. sol. in 45,000 pts. H_2O .

Very sol. in acids. (Foregger and Philipp, J. Soc. Chem. Ind. 1906, 25. 300.)

ZnO_2 (?). Ppt. Decomp. by acids with evolution of H_2O_2 . (Haass, B. 17. 2249.)

ZnO , H_2O , H_2O_2 . (de Forcrand, A. ch. 1902, (7) 27. 58.)

3ZnO , $2\text{H}_2\text{O}_2$. (de Forcrand.)

3ZnO , H_2O , $2\text{H}_2\text{O}_2$. (de Forcrand.)

4ZnO , H_2O , $3\text{H}_2\text{O}_2$. (de Forcrand.)

ZnO_2 , ZnO_2H_2 . Insol. in NH_4OH + Aq. (Kouriloff, A. ch. (6) 23. 431.)

3ZnO_2 , $\text{Zn}(\text{OH})_2$. Sol. in NaOH + Aq with evolution of O . (Eijkman, C. C. 1905, I. 1628.)

Zn_4O_7 , 3ZnO + $4\text{H}_2\text{O}$. Completely sol. in dil. H_2SO_4 . (de Forcrand.)

10ZnO_2 , 4ZnO + $5\text{H}_2\text{O}$. Ppt. (Teletow, C. C. 1911, I. 1799.)

Zinc oxybromide, ZnBr_2 , ZnO + $13\text{H}_2\text{O}$.

ZnBr_2 , 4ZnO + 10, 13, and $19\text{H}_2\text{O}$. Decomp. by H_2O into—

ZnBr_2 , 6ZnO + $35\text{H}_2\text{O}$. (André.)

ZnBr_2 , 5ZnO + $6\text{H}_2\text{O}$. (André.)

All oxybromides are sol. in KOH and NH_4OH + Aq. (André, C. R. 96. 703.)

Zinc oxybromide ammonia, ZnBr_2 , 3ZnO , 2NH_3 + $5\text{H}_2\text{O}$.

Decomp. by H_2O . (André, C. R. 96. 703.)

Zinc oxychloride, ZnO , 3ZnCl_2 + H_2O .

Decomp. by H_2O .

Very sol. in dil. acids. (Ephraim, Z. anorg. 1908, 59. 67.)

+ $4\text{H}_2\text{O}$. Sl. sol. in H_2O ; more sol. in ZnCl_2 + Aq.

Easily sol. in acids, or NH_4OH , or KOH + Aq. (Schindler, Mag. Pharm. 36. 45.)

+ $5\text{H}_2\text{O}$ and $8\text{H}_2\text{O}$. (André, A. ch. (6) 3. 94.)

ZnO , ZnCl_2 + $1\frac{1}{2}\text{H}_2\text{O}$. (Driot, C. R. 1910, 150. 1427.)

3ZnO , ZnCl_2 + $2\text{H}_2\text{O}$. Sl. sol. in H_2O , more easily sol. in ZnCl_2 + Aq. Easily sol. in acids and in NH_4OH or KOH + Aq. (Schindler, Mag. Pharm. 36. 45.)

+ $3\text{H}_2\text{O}$. (Werner, B. 1907, 40. 4443.)

+ $5\text{H}_2\text{O}$. (Mailhe, A. ch. 1902, (7) 27. 367.)

3ZnO , 2ZnCl_2 + $11\text{H}_2\text{O}$. (André, C. R. 1888, 106. 854.)

4ZnO , ZnCl_2 + $6\text{H}_2\text{O}$. (André, C. R. 1888, 106. 854.)

+ $11\text{H}_2\text{O}$. (André, A. ch. (6) 3. 94.)

5ZnO , ZnCl_2 + $6\text{H}_2\text{O}$. (Perrot, Bull. Soc. 1895, (3) 13. 976.)

+ $8\text{H}_2\text{O}$. (André, C. R. 1882, 94. 1524.)

5ZnO , 2ZnCl_2 + $26\text{H}_2\text{O}$. Sol. in KOH or NH_4OH + Aq. Decomp. by H_2O into—

5ZnO , ZnCl_2 + $26\text{H}_2\text{O}$. Sol. in KOH or NH_4OH + Aq. Decomp. by H_2O into—

6ZnO , ZnCl_2 + $6\text{H}_2\text{O}$. Insol. in H_2O . (Kane, A. ch. 72. 296.)

8ZnO , ZnCl_2 + $10\text{H}_2\text{O}$. (Mailhe, A. ch. 1902, (7) 27. 367.)

9ZnO , ZnCl_2 + $3\text{H}_2\text{O}$. Insol. in H_2O . Less sol. in NH_4OH + Aq than ZnCl_2 , 3ZnO + $2\text{H}_2\text{O}$, but easily sol. in + $14\text{H}_2\text{O}$.

9ZnO , 2ZnCl_2 + $12\text{H}_2\text{O}$. Insol. in hot or cold H_2O . (Habermann, M. 5. 432.)

Zinc oxychloride ammonia, 6ZnCl_2 , ZnO , 12NH_3 + $4\text{H}_2\text{O}$.

Decomp. by H_2O and boiling alcohol. (André, A. ch. (6) 3. 90.)

ZnCl_2 , 3ZnO , 2NH_3 + $5\text{H}_2\text{O}$. Decomp. by H_2O . (André.)

ZnCl_2 , 2ZnO , 2NH_3 + $3\text{H}_2\text{O}$. (André.)

6ZnCl_2 , 3ZnO , 10NH_3 + $13\text{H}_2\text{O}$. (André.)

4ZnCl_2 , ZnO , 8NH_3 + $2\text{H}_2\text{O}$. (André.)

Zinc oxyiodide, ZnI_2 , 3ZnO + $2\text{H}_2\text{O}$.

Insol. in cold, sl. sol. in boiling H_2O . (Müller, J. pr. 26. 441.)

ZnI_2 , 9ZnO + $24\text{H}_2\text{O}$. Insol. in cold H_2O .

ZnI_2 , 5ZnO + $11\text{H}_2\text{O}$. Decomp. by H_2O . (Tassilly, C. R. 1896, 122. 324.)

Zinc oxyphosphide, ZnP_2O .

(Renault, A. ch. (4) 9. 162.)

Probably is a mixture of zinc phosphate and phosphorus. (Vigier, Bull. Soc. 1861. 5.)

Zinc oxysulphide, ZnO , ZnS .

Sol. in HCl + Aq. (Arfvedson, Pogg. 1. 59.)

4ZnS , ZnO . Not decomp. by boiling $\text{HC}_2\text{H}_3\text{O}_2$ + Aq. (Kersten, Schw. J. 57. 186.)

Min. *Volzite*. Sol. in HCl + Aq.

Zinc phosphide, ZnP .

Less easily attacked by HCl + Aq than Zn_3P_2 .

ZnP_2 . Not attacked by hot HCl + Aq. (Hvoslef, A. 100. 99.)

ZnP_4 . Insol. in dil. HCl + Aq. (Renault.)

Zn_3P_2 . Insol. in H_2O . Sol. in dil. HCl , H_2SO_4 , or $\text{HNO}_3 + \text{Aq}$, with evolution of PH_3 . (Renault, A. ch. (4) 9. 162.)

Zn_3P_4 . Insol. in $\text{HCl} + \text{Aq}$. (Renault.)

Zinc selenide, ZnSe .

Cold dil. $\text{HNO}_3 + \text{Aq}$ dissolves out Zn , and Se separates out, which dissolves on warming as H_2SeO_3 . (Berzelius.)

$+x\text{H}_2\text{O}$. Insol. in H_2O . (Berzelius.)

Zinc sulphide, ZnS .

Anhydrous. Insol. in H_2O . Sol. in $\text{HCl} + \text{Aq}$; insol. in $\text{HC}_2\text{H}_3\text{O}_2 + \text{Aq}$. (Ebelmen, A. ch. (3) 25. 97.)

Sol. in $\text{H}_2\text{S} + \text{Aq}$ under pressure in a sealed tube. (Senarmont, A. ch. (3) 32. 168.)

Min. *Blende*, *Sphalerite*. Sl. attacked by acids, expecting aqua regia.

1 l. H_2O dissolves 6.65×10^{-6} mols. zinc blende at 18° .

1 l. H_2O dissolves 6.63×10^{-6} mols. artificial cryst. ZnS at 18° . (Wiegel, Z. phys. Ch. 1907, 58. 294.)

Sol. in an alkaline solution of NaClO . (Sadler, Trans. Am. Electrochem. Soc. 1902, 1. 142.)

Insol. in liquid NH_3 . (Franklin, Am. Ch. J. 1898, 20. 830.)

$+1/2$, $2/3$, or $1\text{H}_2\text{O}$.

Pptd. ZnS .

1 l. H_2O dissolves 70.60×10^{-6} mols. pptd. ZnS at 18° . (Wiegel, Z. phys. Ch. 1907, 58. 294.)

Insol. in alkali hydrates, carbonates, and sulphides $+ \text{Aq}$. Insol. in NH_4OH , HCl , or $(\text{NH}_4)_2\text{CO}_3 + \text{Aq}$. Easily sol. in very dil. HCl and $\text{HNO}_3 + \text{Aq}$, but H_2S ppts. ZnS in presence of very dil. $\text{HCl} + \text{Aq}$, or $\text{H}_2\text{SO}_4 + \text{Aq}$. (Eliot and Storer.)

More easily sol. in $\text{HNO}_3 + \text{Aq}$ than in $\text{HCl} + \text{Aq}$. (Fresenius.)

Only sl. sol. in acetic acid. (Wackenroder.)

When still moist is sol. in $\text{H}_2\text{SO}_3 + \text{Aq}$.

Insol. in NH_4Cl or $\text{NH}_4\text{NO}_3 + \text{Aq}$.

$\text{K}_2\text{S} + \text{Aq}$ when added to $\text{ZnSO}_4 + \text{Aq}$ produces a ppt. in presence of 10,000 pts. H_2O , and a slight opalescence with 20,000 pts. (Lassaigne.)

Slowly sol. in conc. $\text{KCN} + \text{Aq}$. (Halm, J. B. 1870. 1008.)

Sl. sol. in $\text{Na}_2\text{S} + \text{Aq}$; sol. in $\text{NaSH} + \text{Aq}$. (Becker, Sill. Am. J. (3) 33. 199.)

Zinc pentasulphide, ZnS_5 .

Sol. in acids, with separation of S . (Schiff, A. 115. 74.)

Zinc sulphosilicide, ZnSiS .

Decomp. by acids and by alkalies. (Fraenkel, Metall, 1909, 6. 683.)

Zinc telluride, ZnTe .

Decomp. by acids. Sol. in $\text{Br}_2 + \text{Aq}$. (Fabre, C. R. 105. 277.)

Zincic acid.

Zinc hydroxide shows weak acid properties and forms the following salts.

Ammonium zincate, $3\text{ZnO}, 4\text{NH}_3 + 12\text{H}_2\text{O} = 3\text{ZnO}, 2(\text{NH}_4)_2\text{O} + 10\text{H}_2\text{O}$.

Decomp. by much H_2O .

Barium zincate, $\text{BaH}_2\text{Zn}_2\text{O}_4 + 7\text{H}_2\text{O}$.

Decomp. by H_2O . (Bertrand, C. R. 111. 939.)

Calcium zincate, $\text{CaH}_2\text{Zn}_2\text{O}_4 + 4\text{H}_2\text{O}$.

Decomp. by H_2O . Sol. in $\text{NH}_4\text{OH} + \text{Aq}$. (Bertrand, C. R. 115. 939.)

Cobaltous zincate, $x\text{CoO}, y\text{ZnO}$.

Rinman's green. Sol. in acids. $\text{H}_2\text{CO}_3 + \text{Aq}$ dissolves out ZnO . (Corney.)

Potassium zincate, $\text{ZnO}, \text{K}_2\text{O}$.

Easily sol. in H_2O , but decomp. by boiling. (Laux, A. 9. 183.)

$2\text{ZnO}, \text{K}_2\text{O}$. Decomp. immediately by cold H_2O . (Fremy, C. R. 15. 1106.)

Sodic zincate, $\text{Na}_2\text{O}, 2\text{ZnO} + 8\text{H}_2\text{O}$, or $2\text{NaHZnO}_2 + 7\text{H}_2\text{O}$.

Decomp. by H_2O or alcohol. (Corney and Jackson, Am. Ch. J. 11. 145.)

$+7\text{H}_2\text{O}$. (Förster and Günther, Z. Elektrochem, 1899, 6. 301.)

$2\text{Na}_2\text{O}, 3\text{ZnO} + 18\text{H}_2\text{O}$ or $\text{Zn}_2\text{O}_4\text{Na}_4\text{H}_4 + 17\text{H}_2\text{O}$. Decomp. by H_2O or alcohol. Insol. in ether. (Corney and Jackson.)

Strontium zincate, $\text{SrH}_2\text{Zn}_2\text{O}_4 + 7\text{H}_2\text{O}$.

Decomp. by H_2O . (Bertrand.)

Zirconic acid.

See Zirconium hydroxide.

Barium zirconate, BaZrO_3 .

Insol. in acids. (Ouvrard, C. R. 113. 80.)

Calcium zirconate, CaZrO_3 .

Insol. in acids. (Ouvrard, C. R. 113. 80.)

Calcium zirconate, acid.

Insol. in H_2O or $\text{HCl} + \text{Aq}$. (Hiordthal, A. 137. 237.)

Calcium potassium zirconate, $(\text{Ca}, \text{K})\text{LaO}_3$ (small quantity of CaO substituted by K_2O).

Sol. in HCl . (Venable, J. Am. Chem. Soc. 1896, 18. 444.)

Cupric zirconate.

(Berthier, A. ch. 59. 195.)

Lithium zirconate, Li_2ZrO_3 .

Easily attacked by acids. (Ouvrard, C. R. **112**. 1444.)

Magnesium zirconate.

Insol. in H_2O or $\text{HCl} + \text{Aq.}$ (Hiordthal, C. R. **61**. 215.)

Potassium zirconate.

Decomp. by $\text{HCl} + \text{Aq.}$ (Knop, A. **159**. 44.)

Sodium zirconate, Na_2ZrO_3 .

Decomp. by H_2O .

Na_4ZrO_4 . Decomp. by $\text{HCl} + \text{Aq.}$ and is dissolved by subsequent addition of H_2O .

Na_2O , $8\text{ZrO}_2 + 12\text{H}_2\text{O}$. (Hiordthal.)

Strontium zirconate, SrZrO_3 .

As CaZrO_3 . (Ouvrard.)

Zirconium, Zr.

Crystallized. Attacked by conc. $\text{HCl} + \text{Aq.}$ above 50° , but very slowly even at 100° ; rapidly by hot aqua regia. Sol. in cold conc. $\text{HF} + \text{Aq.}$ (Troost, C. R. **61**. 109.)

Very violently attacked by a mixture of HNO_3 and HF . (Berzelius, Pogg. **4**. 117.)

Amorphous. Slowly attacked by boiling aqua regia, H_2SO_4 , or conc. $\text{HCl} + \text{Aq.}$ (Berzelius.)

Easily sol. in HF or $\text{HNO}_3 + \text{HF}$.

Zirconium bromide, ZrBr_4 .

Very hygroscopic. Violently decomp. by H_2O to form oxybromide. (Melliss, Zeit. Ch. (2) **6**. 296.)

Sl. sol. in organic solvents. (Matthews, J. Am. Chem. Soc. 1898, **20**. 840.)

Zirconium bromide ammonia, $\text{ZrBr}_4 \cdot 4\text{NH}_3$.

Ppt. Insol. in organic solvents. (Matthews, J. Am. Chem. Soc. 1898, **20**. 840.)

$\text{ZrBr}_4 \cdot 10\text{NH}_3$. Very hygroscopic.

Decomp. by H_2O . (Stähler, B. 1905, **38**. 2612.)

Zirconium carbide, ZrC .

Insol. in H_2O and $\text{NH}_4\text{OH} + \text{Aq.}$ and HCl Aq even when heated. Sol. in HNO_3 , H_2SO_4 and fused alkali nitrates, chlorates, or hydroxides. (Moissan, C. R. 1896, **122**. 653.)

Zirconium chloride, ZrCl_4 .

Sol. in H_2O with evolution of much heat to form ZrOCl_2 . Sol. in alcohol. (Hinsberg, A. **239**. 253.)

Very unstable.

Probably substances so described in the literature by Nylander and others were oxychlorides. (Venable, J. Am. Chem. Soc. 1894, **16**. 471.)

Sol. in ether. (Matthews, J. Am. Chem. Soc. 1898, **20**. 821.)

Zirconium chloride ammonia, $\text{ZrCl}_4 \cdot 2\text{NH}_3$.

Fumes in the air. Decomp. by H_2O . (Matthews, J. Am. Chem. Soc. 1898, **20**. 821.)

$\text{ZrCl}_4 \cdot 3\text{NH}_3$. (Stähler, B. 1905, **38**. 2611.)

$\text{ZrCl}_4 \cdot 4\text{NH}_3$. Decomp. by H_2O . (Paykull.)

Unstable. Decomp. by H_2O . (Matthews, J. Am. Chem. Soc. 1898, **20**. 821.)

$\text{ZrCl}_4 \cdot 8\text{NH}_3$. Stable in the air. Decomp. by H_2O . Insol. in ether. (Matthews, J. Am. Chem. Soc. 1898, **20**. 821.)

Very hygroscopic. Decomp. by H_2O . (Stähler, B. 1905, **38**. 2611.)

Zirconium fluoride, ZrF_4 .

Anhydrous. Insol. in H_2O and acids. (Deville, A. ch. (3) **49**. 84.)

Only sl. sol. in H_2O .

1.388 g. dissolve in 100 cc. H_2O without hydrolysis. On warming the solution, zirconium hydrate begins to ppt. out at about 50° . (Wolter, Ch. Z. 1908, **32**. 606.)

+ $3\text{H}_2\text{O}$. Sol. in H_2O , but solution decomposes on diluting, with pptn. of an insol. basic salt. Sol. in dil. $\text{HF} + \text{Aq.}$ (Berzelius.)

Zirconium fluoride ammonia, $5\text{ZrF}_4 \cdot 2\text{NH}_3$.

(Wolter, Ch. Z. 1908, **32**. 607.)

Zirconium hydride, ZrH_2 .

Not attacked by acids. (Winkler, B. **24**. 873.)

Zirconium hydroxide, $\text{Zr}(\text{OH})_4$.

Insol. in H_2O or alcohol. Sol. in 5000 pts. H_2O . (Melliss.)

Sol. in acids, even oxalic or tartaric acid, when precipitated cold. If precipitated hot, it is slowly dissolved upon heating with conc. acids.

Sol. in dil. or conc. min. acids except HI . Readily sol. in oxalic, only sl. sol. in acetic acid.

Much less sol. when pptd. from hot solution than when pptd. from cold solution. (Venable, J. Am. Chem. Soc. 1898, **20**. 274.)

Sl. sol. in $(\text{NH}_4)_2\text{CO}_3 + \text{Aq.}$ Insol. in K_2CO_3 and $\text{Na}_2\text{CO}_3 + \text{Aq.}$ Insol. in NaOH , KOH , and $\text{NH}_4\text{OH} + \text{Aq.}$

Sol. in $(\text{NH}_4)_2\text{C}_4\text{H}_4\text{O}_6 + \text{NH}_4\text{OH} + \text{Aq.}$ Insol. in NH_4 salts + Aq.

Zirconium iodide, ZrI_4 .

Fumes in the air.

Sol. in H_2O and acids with violent reaction.

Decomp. by alcohol. Sol. in abs. ether.

Sl. sol. in benzene and CS_2 . (Stähler, B. 1904, **37**. 1137.)

Insol. in H_2O , HNO_3 , HCl , aqua regia, and CS_2 .

Sol. in H_2SO_4 with decomp; unchanged by boiling H_2O . (Dennis, J. Am. Chem. Soc. 1896, **18**. 678.)

Zirconium iodide ammonia, $\text{ZrI}_4 \cdot 6\text{NH}_3$;
 $\text{ZrI}_4 \cdot 7\text{NH}_3$; $\text{ZrI}_4 \cdot 8\text{NH}_3$; $\text{ZrI}_4 \cdot 10\text{NH}_3$.

All above comps. are hygroscopic and lose NH_3 in the air. (Stähler, B. 1905, **38**. 2615.)

Zirconium nitride.

Scarcely attacked by acids, aqua regia, and caustic alkalies. Slowly decomp. by long contact with H_2O . (Mallet, Sill. Am. J. (2) **28**. 346.)

Zr_3N_2 . Decomp. when heated in the air; sol. in HF ; insol. in other min. acids. (Matthews, J. Am. Chem. Soc. 1898, **20**. 844.)

Zr_2N_3 . Decomp. when heated in the air. Sol. in HF ; insol. in other min. acids. (Matthews, J. Am. Chem. Soc. 1898, **20**. 844.)

Zirconium oxide, ZrO_2 .

When ignited, is insol. in all acids except HF and H_2SO_4 . Sl. sol. in HF ; sol. in H_2SO_4 only when very finely powdered and heated with a mixture of 2 pts. H_2SO_4 and 1 pt. H_2O until the H_2SO_4 volatilises. (Berzelius.)

Zirconium peroxide, ZrO_3 .

(Cleve, Bull. Soc. (2) **43**. 53), or Zr_2O_5 according to Bailey (Chem. Soc. **49**. 150).

Not attacked by cold dil. $\text{H}_2\text{SO}_4 + \text{Aq}$. (Bailey.)

Zirconium silicon oxide.

Min. *Zircon*. See Silicate, zirconium.

Zirconium oxy-compounds.

See Zirconyl compounds.

Zirconium phosphide, ZrP_2 .

Insol. in dil. or conc. acids and alkalies.

Sl. sol. in aqua regia. (Gewecke, A. 1908, **361**. 85.)

Zirconium silicide, ZrSi_2 .

Sol. in HF , insol. in other min. acids.

Not acted upon by 10% $\text{KOH} + \text{Aq}$ or $\text{NaOH} + \text{Aq}$.

Decomp. by fusion with KOH . (Hönigschmid, C. R. 1906, **143**. 225.)

Zirconium sulphide.

Insol. in H_2O . Sol. in HF ; slowly sol. in aqua regia. Insol. in HNO_3 , HCl , H_2SO_4 , or $\text{KOH} + \text{Aq}$. (Berzelius.)

Insol. in dil. acids. Sol. in conc. $\text{HNO}_3 + \text{Aq}$ (perhaps an oxysulphide). (Fremy.)

Zirconomolybdic acid.

Ammonium zirconomolybdate,

$2(\text{NH}_4)_2\text{O}, \text{ZrO}_2, 12\text{MoO}_3 + 10\text{H}_2\text{O}$.

Sol. in H_2O . (Pechard, C. R. 1893, **117**. 790.)

Potassium zirconomolybdate,
 $2\text{K}_2\text{O}, \text{ZrO}_2, 12\text{MoO}_3 + 18\text{H}_2\text{O}$.

Sol. in H_2O . (Pechard.)

Zirconotungstic acid.

Ammonium zirconodecatungstate,

$3(\text{NH}_4)_2\text{O}, \text{ZrO}_2, 10\text{WO}_3 + 13\text{H}_2\text{O}$.

Very sol. in H_2O .

Efflorescent. (Hallopeau, Bull. Soc. (3) **15**. 921.)

$3(\text{NH}_4)_2\text{O}, \text{H}_2\text{O}, \text{ZrO}_2, 10\text{WO}_3 + 13\text{H}_2\text{O}$. Sol. in H_2O ; unstable; effloresces in the air. (Hallopeau.)

Potassium zirconotungstate, $4\text{K}_2\text{O}, \text{ZrO}_2, 10\text{WO}_3 + 20\text{H}_2\text{O}$.

Sol. in hot H_2O . (Hallopeau.)

$4\text{K}_2\text{O}, \text{ZrO}_2, 10\text{WO}_3 + 15\text{H}_2\text{O}$. More sol. in hot than in cold H_2O .

Sol. in fused alkali carbonates. (Hallopeau.)

Potassium dizirconodecatungstate,

$4\text{K}_2\text{O}, 2\text{ZrO}_2, 10\text{WO}_3 + 20\text{H}_2\text{O}$.

More sol. in hot than in cold H_2O .

Sol. in fused alkali carbonates. (Hallopeau.)

Zirconyl bromide, $\text{ZrOBr}_2 + 3\text{H}_2\text{O}$.

Deliquescent. Decomp. in moist air. Very sol. in H_2O . (Venable, J. Am. Chem. Soc. 1898, **20**. 324.)

$+7\text{H}_2\text{O}$. Sol. in H_2O . (Melliss.)

$+8\text{H}_2\text{O}$. Deliquescent. Decomp. in air. Very sol. in H_2O . (Venable, J. Am. Chem. Soc. 1898, **20**. 324.)

$+13\text{H}_2\text{O}$. Deliquescent. Decomp. in moist air.

Very sol. in H_2O . (Venable.)

$+14\text{H}_2\text{O}$. Deliquescent. Decomp. in moist air. (Venable.)

Very sol. in H_2O . (Venable.)

$\text{ZrBr}(\text{OH})_2 + \text{H}_2\text{O}$, and $+2\text{H}_2\text{O}$. Decent, and decomp. in moist air. (Venable.)

Zirconyl chloride, ZrOCl_2 .

Sol. in H_2O ; insol. in HCl . (Venable, J. Am. Chem. Soc. 1894, **16**. 475.)

$+2\text{H}_2\text{O}$. (Chauvenet, C. R. 1912, **822**.)

$+3\text{H}_2\text{O}$. Sol. in H_2O . (Venable.)

$+3.5\text{H}_2\text{O}$. (Chauvenet, C. R. 1912, **822**.)

$+6\text{H}_2\text{O}$. (Chauvenet.)

Sol. in H_2O .

Insol. in HCl . (Venable.)

$+4\frac{1}{2}\text{H}_2\text{O}$, $6\frac{1}{2}\text{H}_2\text{O}$, and $8\text{H}_2\text{O}$.

Efflorescent. Easily sol. in H_2O and in conc. $\text{HCl} + \text{Aq}$. (Venable, J. Am. Chem. Soc. 1898, **20**. 321.)

$+8\text{H}_2\text{O}$. Sol. in H_2O . (Venable, J. Am. Chem. Soc. 1898, **20**. 321.)

Effloresces in the air.

l. in H_2O . Less sol. in $HCl + Aq.$ and
 y insol. in conc. HCl . (Chauvenet,
 . 1912, **154**. 822.)
 $ZrOCl_2$. Sol. in H_2O and alcohol. (Ende-
 i, J. pr. (2) **11**. 219.)
 t decomp. by H_2O .
 l. in dil. HCl . (Chauvenet, C. R. 1912,
 1236.)
 H_2O . (Chauvenet, C. R. 1912, **154**.
)
 $3H_2O$. Decomp. by H_2O . Sol. in dil
 (Chauvenet.)
 $rO_2, 7HCl$. Sol. in H_2O . (E.)
 OCl_6 . (Troost and Hautefeuille, C. R.
 163.)

$Zr_2OCl_4 = ZrCl_4, 2ZrO_2$. Insol. in H_2O .
 (Hermann.)

Zirconyl iodide, $ZrOI_2 + 8H_2O$.

Very sol. in H_2O and alcohol.

Very hygroscopic. ((Stähler, B. 1904,
37. 1138.)

$ZrI(OH)_3 + 3H_2O$. Easily sol. in H_2O .
 (Hinsberg, A. **239**. 253.)

Ppt. Sl. sol. in $HI + Aq.$ (Venable, J.
 Am. Chem. Soc. 1898, **20**. 328.)

Zirconyl sulphide (?)

Decomp. by HNO_3 with separation of S.
 (Fremy, A. ch. (3) **38**. 326.)



APPENDIX

FORMULÆ FOR CONVERTING AREOMETER DEGREES INTO SPECIFIC GRAVITY.

n = no. of degrees on the areometer scale; sp. gr. = specific gravity.

Areometer	Temp.	Liquids heavier than H ₂ O	Liquids lighter than H ₂ O
<p>1. Baumé. (a) <i>According to Baumé's original directions.</i> For liquids heavier than H₂O. Sp. gr. of a solution of 15 pts. NaCl dissolved in 85 pts. H₂O at 12.5° ($d_{12.5^\circ}^{12.5^\circ} = 1.1118988$) = 15°; H₂O = 0°. For liquids lighter than H₂O. Sp. gr. of 10% NaCl + Aq at 12.5° ($d_{12.5^\circ}^{12.5^\circ} = 1.0737665$) = 0°; H₂O = 10°.</p> <p>(b) <i>Old Form.</i> Liquids heavier than H₂O, 10% NaCl + Aq at 15° ($d_{15^\circ}^{15^\circ} = 1.073350$) = 10°; H₂O = 0°. Liquids lighter than H₂O, 10% NaCl + Aq = 0°; H₂O = 10°.</p> <p>(c) <i>New Form.</i> So-called "Rational Scale." Liquids heavier than H₂O, H₂SO₄ + Aq $\frac{15^\circ}{15^\circ} = 1.842 = 66^\circ$; H₂O = 0°.</p>	<p>15°</p> <p>12.5°</p> <p>15°</p> <p>17.5°</p> <p>15°</p>	<p>Sp. gr. = $\frac{149.05}{149.05 - n}$</p> <p>Sp. gr. = $\frac{145.88}{145.88 - n}$</p> <p>Sp. gr. = $\frac{146.3}{146.3 - n}$</p> <p>Sp. gr. = $\frac{146.78}{146.78 - n}$</p> <p>Sp. gr. = $\frac{144.3}{144.3 - n}$</p>	<p>Sp. gr. = $\frac{145.56}{135.56 + n}$</p> <p>Sp. gr. = $\frac{145.88}{135.88 + n}$</p> <p>Sp. gr. = $\frac{146.3}{136.3 + n}$</p> <p>Sp. gr. = $\frac{146.78}{136.78 + n}$</p> <p></p>
<p>2. Beck. H₂O = 0°; liquid of 0.850 sp. gr. ($\frac{12.5^\circ}{12.5^\circ}$) = 30°. Scale continued above and below.</p>	12.5°	Sp. gr. = $\frac{170}{170 - n}$	Sp. gr. = $\frac{170}{170 + n}$
<p>3. Twaddle. H₂O = 0°. Each degree corresponds to an increase of 0.005 in the sp. gr.</p>	Given on the instrument	Sp. gr. = 1.000 + 0.005n	

TABLES FOR THE CONVERSION OF BAUMÉ DEGREES
INTO SP. GR.

Since the original directions of Baumé there have been many slight modifications suggested, so that there are several varieties of Baumé hydrometers with somewhat varying readings, tables for the two principal ones of which are here-given.

1. According to Baumé's original directions.

For liquids heavier than H₂O. Sp. gr. of 15 % NaCl+
 $Aq \left(\frac{12.5^{\circ}}{12.5^{\circ}} \right) = 1.1118988 = 15^{\circ}; H_2O = 0^{\circ}.$

Calculated according to the formula, sp. gr. = $\frac{149.05}{149.05 - n}.$

Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.
0	1.00000	20	1.15497	39	1.35438	58	1.63701
1	1.00675	21	1.16399	40	1.36680	59	1.65519
2	1.01360	22	1.17316	41	1.37945	60	1.67378
3	1.02054	23	1.18246	42	1.39234	61	1.69279
4	1.02757	24	1.19192	43	1.40547	62	1.71223
5	1.03471	25	1.20153	44	1.41885	63	1.73213
6	1.04194	26	1.21129	45	1.43248	64	1.75250
7	1.04927	27	1.22122	46	1.44638	65	1.77335
8	1.05671	28	1.23131	47	1.46056	66	1.79470
9	1.06426	29	1.24156	48	1.47501	67	1.81657
10	1.07191	30	1.25199	49	1.48971	68	1.83899
11	1.07968	31	1.26260	50	1.50479	69	1.86196
12	1.08755	32	1.27338	51	1.52014	70	1.88551
13	1.09555	33	1.28436	52	1.53580	71	1.90967
14	1.10366	34	1.29522	53	1.55179	72	1.93446
15	1.11189	35	1.30688	54	1.56812	73	1.95989
16	1.12025	36	1.31844	55	1.58471	74	1.98601
17	1.12873	37	1.33621	56	1.60182	75	2.01283
18	1.13735	38	1.34218	57	1.61923	76	2.04038
19	1.14609

For liquids lighter than H₂O. Sp. gr. of 10 % NaCl

$Aq \left(\frac{12.5^{\circ}}{12.5^{\circ}} \right) = 1.0737665 = 0^{\circ}; H_2O = 10^{\circ}.$

Calculated according to the formula, $sp. \text{ gr.} = \frac{145.56}{135.56 + n}.$

Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.
10	1.00000	30	0.87919	50	0.78443	65	0.72577
15	0.96679	35	0.85342	55	0.76385	70	0.70811
20	0.93571	40	0.82912	60	0.74432	75	0.69130
25	0.90657	45	0.80616

2. According to the so-called Rational Scale.

$Sp. \text{ gr. of } H_2SO_4 + Aq \left(\frac{15^{\circ}}{15^{\circ}} \right) = 1.842 = 66^{\circ} ; H_2O = 0^{\circ}.$

Calculated according to the formula, $sp. \text{ gr.} = \frac{144.3}{144.3 - n}.$

Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.	Deg. Baumé	Sp. gr.
1	1.007	18	1.142	35	1.320	51	1.547
2	1.014	19	1.152	36	1.332	52	1.563
3	1.021	20	1.161	37	1.345	53	1.580
4	1.029	21	1.170	38	1.357	54	1.598
5	1.036	22	1.180	39	1.370	55	1.616
6	1.043	23	1.190	40	1.384	56	1.634
7	1.051	24	1.200	41	1.397	57	1.653
8	1.059	25	1.210	42	1.411	58	1.672
9	1.066	26	1.220	43	1.424	59	1.692
10	1.074	27	1.230	44	1.439	60	1.712
11	1.082	28	1.241	45	1.453	61	1.732
12	1.091	29	1.251	46	1.468	62	1.753
13	1.099	30	1.262	47	1.483	63	1.775
14	1.107	31	1.274	48	1.498	64	1.797
15	1.116	32	1.285	49	1.514	65	1.820
16	1.125	33	1.296	50	1.530	66	1.842
17	1.133	34	1.308

SYNCHRONISTIC TABLE OF CHEMICAL

Year	A.	A. ch.	Am. J. Sci.	Ann. Min.	Ans. Phil.	Arch. Pharm.	Ch. Gas.	C. R.	Dugl.
1800	.	(1) 32-34
1801	.	35-39
1802	...	40-43
1803	.	44-47
1804	.	48-51
1805	..	52-55
1806	.	56-60
1807	.	61-64
1808	..	65-68
1809	.	69-72
1810	...	73-76
1811	..	77-80
1812	..	81-84
1813	...	85-88	..	.	(1) 1, 2
1814	...	89-92	.	.	3, 4
1815	..	93-96	.	.	5, 6
1816	.	(2) 1-3	.	.	7, 8
1817	.	4-6	.	1, 2	9, 10
1818	.	7-9	.	3	11, 12
1819	.	10-12	(1) 1	4	13, 14
1820	.	13-15	2	5	15, 16	.	.	.	1-3
1821	.	16-18	3	6	(2) 1, 2	.	.	.	4-6
1822	..	19-21	4, 5	7	3, 4	1, 2	.	.	7-9
1823	.	22-24	6	8	5, 6	3-6	.	.	10-12
1824	.	25-27	7, 8	9	7, 8	7-10	.	.	13-15
1825	.	28-30	9	10, 11	9, 10	11-14	.	.	16-18
1826	.	31-33	10, 11	12, 13	11, 12	16-19	.	.	19-22
1827	.	34-36	12	(2) 1, 2	.	20-23	.	.	24-26
1828	.	37-39	13, 14	3, 4	.	24-26	.	.	27-30
1829	.	40-42	15, 16	5, 6	.	27-30	.	.	31-34
1830	.	43-45	17, 18	7, 8	.	31-34	.	.	35-38
1831	.	46-48	19, 20	.	.	35-39	.	.	39-42
1832	1-4	49-51	21, 22	(3) 1, 2	.	40-43	.	.	43-47
1833	5-8	52-55	23, 24	3, 4	.	44-47	.	.	48-50
1834	9-12	56-57	25-27	5, 6	.	48-50	.	.	51-54
1835	13-16	58-60	28, 29	7, 8	.	(2) 1-4	.	1	55-58
1836	17-20	61-63	30, 31	9, 10	.	5-8	.	2, 3	59-62
1837	21-24	64-66	32, 33	11, 12	.	9-12	.	4, 5	63-66
1838	25-28	67-69	35, 35	13, 14	.	13-16	.	6, 7	67-70
1839	29-32	70-72	36, 37	15, 16	.	17-20	.	8, 9	71-74
1840	33-36	73-75	38, 39	17, 18	.	21-24	.	10, 11	75-78
1841	37-40	(3) 1-3	40, 41	19, 20	.	25-28	.	12, 13	79-82
1842	41-44	4-6	42, 43	(4) 1, 2	.	29-32	.	14, 15	83-86
1843	45-48	7-9	44, 45	3, 4	.	33-36	1	16, 17	87-90
1844	49-52	10-12	46, 47	5, 6	.	37-40	2	18, 19	91-94
1845	53-56	13-15	48-50	7, 8	.	41-44	.	20, 21	95-98
1846	57-60	16-18	(2) 1, 2	9, 10	.	45-48	3	22, 23	99-102
1847	61-64	19-21	3, 4	11, 12	.	49-52	4	24, 25	103-106

AND OTHER SCIENTIFIC PERIODICALS—Part I.

Glab. Ann.	J. Chim med.	J. Pharm	J. pr.	Phil. Mag.	Pogg.	Proc. Am Acad.	Proc Roy. Soc.	Q. J. Sci.	Scher J	Schw. J.
4-6	6-8		.	.	.	3, 4	..
7-9	9-11		.	.	.	5, 6	.
10-12	12-14	7, 8	...
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